Low-level analysis of microarray data

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LOW-LEVEL ANALYSIS OF MICROARRAY DATA

HENRIK BENGTSSON

LUND INSTITUTE OF TECHNOLOGY
Lund University
Centre for Mathematical Sciences
Mathematical Statistics
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Identifying differentially expressed genes in cDNA microarray experiments: making aging visible

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Lund (Mondo), August 26, 2004.

Henrik Bengtsson
x
Low-level analysis of microarray data

That’s one small step for mankind; one giant leap for a man.

1 Introduction

In today’s life sciences much attention is on systems biology and functional genomics where researchers try to identify and study important genes and complex genetic regulatory networks in order to understand the fundamental properties of living organisms and especially the dynamics of the cells. This thesis is all about cleaning up measurements obtained by one very promising and less than a decade old technology, a technology that makes it possible for scientists to study the cells in ways that have never been done before.

The introductory part of this seven-paper PhD thesis in Mathematical Statistics is outlined as follows. In Section 2, we give a brief introduction to the molecular biology of the cell. In the following section we introduce the marvelous technology, explain how it works, and give some initial motivations for the area of our own work. In Section 4, in order to further motivate and give more background to the papers presented, we give our formalized view of gene-expression analysis in general. In Section 5, we describe the software developed making it possible for us to quickly communicate new statistical methods with the research community. In the final section, based on these introductory sections, we conclude by providing a summary for each of the papers.

2 On DNA, proteins, and gene activities

There is no exact definition of what a gene is as it is used slightly differently depending on the context. Historically, its definition has evolved from the time pre-dating the discovery of DNA (deoxyribonucleic acid) when a gene was just “the atom” of an inheritable trait (now a genotype) and which later was pinned
down to be a region on a chromosome. Now a gene is said to be a specific region on the DNA that produces a functional product, that is a protein (or a special type of RNA (ribonucleic acid) molecule). Proteins are complex macro molecules that are essential for the function, structure, and dynamics of a cell. Each protein has a unique function. A cell interacts and responds to its environment by means of proteins and other molecules through a complex regulatory network. By studying these regulatory networks, scientists aim to understand and develop better treatments for complex diseases such as cancer, brain disorders, diabetes, but also viral diseases such as Acquired Immune Deficiency Syndrome (AIDS) and Severe Acute Respiratory Syndrome (SARS).

A gene is said to be activated or expressed when it, or more precisely, its coded information, is converted into proteins. Note, there is no single definition for this term. Somewhat simplified, gene expression takes place in two steps. In the first step, the gene’s so called template strand of the double-helix DNA is transcribed into messenger RNA (mRNA), which is a complementary copy of the sequence of nucleotides (bases) on the DNA. In the second step, the mRNA moves out of the nucleus into the cell’s cytoplasm and gets translated into a protein. This is done such that continuous triplets of mRNA nucleotides (codons) are converted into a sequence of amino acids that folds into a protein. Because the same amino-acid sequence can be subsequently modified in different ways, but also for other reasons, there is not a one-to-one relationship between the DNA and the final functional proteins; there are many more types of proteins than genes. Moreover, several protein copies of the same gene can be produced, which is done by the transcription of multiple mRNA copies or by multiple translations of the same mRNA copy. The more product, the more the gene is expressed. In biology, the aforementioned process is referred to as the Central Dogma of Biology. For further details see [1, 9].

When a cell responds to its environment, undergoes division, is attacked by viruses etc., different genes are activated and expressed in various amounts. By studying the gene-expression levels of cells under various (controlled or uncontrolled) conditions, researchers wish to infer the properties and regulatory networks of the cells. For various reasons not discussed here, it is easier to study the gene expressions by measuring the amount of mRNA than the amount of proteins. Standard protocols have been developed making it possible for any lab-
3. The revolutionizing technology

The microarray technology has revolutionized the research area on gene-expression analysis by providing a relatively cheap and high-throughput platform for quantifying tens of thousands of gene expressions simultaneously. This makes it possible to collect huge sets of data that can be analyzed in order to infer functions of the cells. These data sets of high dimensions raise many new and interesting questions on statistical methodology. Some of them may be answered using classical theory of multivariate and cluster analysis, Bayesian statistics, robust and nonparametric statistics, variance components techniques and multiple testing, but others require new tools from these (and other) fields; see [17]. This calls for more statisticians and mathematicians to get involved.

The most general way to think of a microarray-based gene-expression experiment is as a black box. In one end the cell sample of interest is inserted and in the other end its gene-expression levels are output. However, as this thesis shows, in order to get correct and high quality measurements this box has to be opened and its components need to be investigated.

3.1 Brief about the technology

A microarray has well defined regions (probes) that each consists of immobilized sequences of DNA that are unique to a specific gene. When fluorophore labeled complementary DNA (cDNA) sequences (targets), obtained by reverse transcription of mRNA extracted from the samples of interest are hybridized to the probes, each region on the microarray will specifically bind a certain amount of DNA unique to the corresponding gene. Depending on if a two-channel or single-channel microarray platform is used, either several and differentially labeled targets are hybridized to the same array, or different targets are each hybridized to an exclusive array using identical labels. Fluorescent and radioactive molecules are standard types of labels. Next, the array is scanned at different wavelengths to excite the fluorescent molecules using a light source, typically a laser. Shortly after
being excited, the fluorophores emit photons, which are registered and quantified in each position resulting in one high-resolution digitized image for each channel. Using image analysis methods, the pixels that belong to the regions that contain the probes are identified and their intensities are averaged giving estimates of the transcript abundances for all genes involved. For visualization purposes, the red and the green images are often combined into one image where the spots (probes) are falsely colored on a scale from red to yellow to green corresponding to the amount of relative gene expression. A schematic overview of this process is given in Figure 1. Spotted DNA microarrays [15] is a multi-channel technique that

![Diagram](image)

**Figure 1:** A schematic overview of a two-channel microarray technology. The gene-expression level, that is, the amount of mRNA, for gene \(i\) in the red and the green channel is \(x_{R,i}\) and \(x_{G,i}\), respectively. For each sample, the mRNA is extracted, purified, and reversely transcribed into cDNA which is labeled with a dye that is unique for each channel. The labeled cDNA is mixed in equal amounts and hybridized to the probes on the microarray glass slide. The amount of hybridized cDNA of gene \(i\) is denoted by \(z_{R,i}\) and \(z_{G,i}\), respectively. To be explained in details below, we formally denote these steps by the submeasurement functions \(g_{R}\) and \(g_{G}\). Next, with help of a scanner and image-analysis procedures, we try to estimate the amount of hybridized cDNA. The estimates are called \(y_{R}\) and \(y_{G}\) and the latter steps are denoted by the functions \(h_{R}\) and \(h_{G}\), respectively.
3. The revolutionizing technology

measures the (relative) abundances of mRNA of 5,000-50,000 genes from two (or more) samples by competitive co-hybridization of aliquot labeled cDNA targets. To date, two-channel microarrays are by far the most commonly used types of arrays. Contrary, microarray techniques such as nylon membrane microarrays and high-density (100,000-500,000 probes) oligonucleotide in-situ microarrays [8] measure the abundances of gene expressions in one single sample. For a further introduction to the microarray technology, see Papers B, E, and D in this thesis. For a thorough description and history of the microarray technology, see [14].

3.2 Other usage

The DNA microarray technology is not limited to measure gene-expression levels and to pinpoint genes that are characteristic to a certain cell type, although most frequently used so. To give a few examples, the technique is also used to sequence DNA and detect genomic abnormalities such as amplifications and deletions of DNA. Low-resolution classical comparative genomic hybridization (CGH) methods, which are widely used in cancer research, are being replaced by more high-throughput microarray-based alternatives. Furthermore, with the advent of microarrays it has become practically and economically feasible to provide diagnostic tests on an individual basis as a complement to traditional clinical tests. More targeted treatments with less suffering for the patients can be developed.

3.3 Calls for improvements

There is a continuous and exponentially growing stream of publications with biological results of great value that are obtained by means of microarrays. Although the technology is highly praised and many advances have been made since the first proof-of-concept experiments performed a decade ago [15], there is also an increasing number of calls for improvements of protocols and statistical methods [7]. One crucial problem is that data generated with equal biological setups but by different laboratories and/or by different platforms may not easily be combined and integrated into the same analysis. This is even the case when experiments generated over time by the same facility are compared. To give a real-world example, a local collaborator is facing the tremendous task to pre-process and analyze data from almost 1000 hybridizations of scarce RNA extracts. These experiments are produced over a period of more than one year including many different cell cultures, different reagents, different people, various print batches (including dif-
ferent array types), climate conditions and so on. The first major task is to identify and remove systematic variation that obscures the biological signals by a process referred to as normalization and calibration or, more general, as low-level analysis. This thesis provides substantially improved methods for calibration and normalization of microarray data.

4 On microarray and gene-expression analysis

To give a more detailed motivation, but also a better understanding for the studies presented in this thesis, we will next give our view of microarray experiments from the perspective of gene-expression analysis. We will allow ourselves to generalize as much as possible in order to keep the discussion simple and we ask the reader to keep in mind that exceptions may exist.

4.1 Gene expression

In the context of microarray analysis, we refer to number of gene transcripts, that is, the number of mRNA transcripts for a specific gene available at a certain time, as the expression level of that gene. Note that this is a refinement of the more general “definition” introduced in Section 2. Although the sensitivity and specificity of the microarray techniques improve continuously, gene-expression levels are still not measured on a single-cell basis. In reality, the average expression level in a large number of cells is measured. Thus, when talking about the expression level of a gene in the context of microarray experiments it is typically meant the average number of mRNA transcripts in a set of cells. To formalize this, assume we are interested in a set of genes $I = \{1, \ldots, I\}$ in the cell samples $C = \{1, \ldots, C\}$ where $I$ is the number of genes and $C$ is the number of cell samples. Let $N_c$, which is typically large, be the total number of cells in sample $c \in C$, and let $n_{c,i}$ be the total number of gene transcripts for gene $i \in I$ in all those cells. Then, the average number of gene transcripts per cell for gene $i$ is $x_{c,i} = n_{c,i}/N_c$. We refer to $x_{c,i}$ as the gene-expression level for gene $i$ in sample $c$.

4.2 Expressed and differentially expressed genes

The simplest microarray experiment has only one cell sample, that is $C = 1$. Ideally, it allows us to classify genes to either be non-expressed or expressed. For
4. On microarray and gene-expression analysis

A gene to be non-expressed no gene transcripts are allowed, otherwise it is expressed. Formally, for sample \( c \), genes \( I_{x_c=0} = \{ i \in I : x_{c,i} = 0 \} \) are labeled "non-expressed", and all other genes, \( I_{x_c \neq 0} = I \setminus I_{x_c=0} \), are labeled "expressed". However, rather than being either on or off in this sense, in reality a gene can be expressed with any number of gene transcripts. Moreover, it does not make much sense to describe the gene activity as the (absolute or average) number of gene transcripts. For instance, for a certain gene it might be enough to be expressed in one mRNA copy in order to affect a cell's state whereas for another gene there might be a threshold of, say, 2000 transcripts in order to make a difference. It can also be the opposite that a gene is normally expressed in 2000 transcripts per cell and makes a difference first when its expression level drops below 500 transcripts, and so on. Instead, it is better to quantify the amount of activity as the expression levels compared to a reference. The reference should not be the level of another gene, because it makes little sense to compare \( x_{c,i} \) with \( x_{c,j} \) for \( i \neq j \), but instead the expression level of the same gene under other circumstances, that is \( x_{c,i} \) compared to \( x_{d,i} \) for \( c \neq d \). As an example, it is more informative to compare the expression level of a promoter oncogene in a tumor tissue to its expression level in a healthy tissue, than to compare the expression level of a promoter oncogene with the expression level of an inhibitor oncogene. Thus, in addition to classifying genes as non-expressed or expressed, an experimental setup with two or more samples allows us to classify genes as differentially or non-differentially expressed. Formally the set of genes labeled "non-differentially expressed" is \( I_{x_c=x_d} = \{ i \in I : x_{c,i} = x_{d,i} \} = \{ i \in I : N_{d,n_{c,i}} = N_{c,n_{d,i}} \} \) for some \( c \neq d \) with \( c, d \in C \). All other genes, \( I_{x_c \neq x_d} = I \setminus I_{x_c=x_d} \), are labeled "differentially expressed".

Given the above foundation and binary classification schemes, it is possible to test rather complicated biological hypotheses of different flavors involving the cell samples of interest. By far, the most common application of microarray experiments to date has been to scan for genes that are differentially expressed, that is, to test the null hypothesis \( H_0 \) against \( H_1 \):

\[
H_0 : x_{c,i} = x_{d,i} \\
H_1 : x_{c,i} \neq x_{d,i}
\]

(1)

In some analyses such as cell-cycle studies, it may indeed be of interest to look at the expression level of a single gene in relation to all genes, that is, \( x_{c,i} / \sum_{j \in I} x_{c,j} \).
for all genes $i \in I$ and in samples $c \neq d; c, d \in C$.

### 4.3 Relative gene expression and ordering of expression levels

In addition to the above, many experiments try to order the gene-expression levels relative to each other. For instance, the question asked may be if the expression of gene $i$ is increasing between time $t_1$ and $t_0$ or not. This can be done by testing the null hypothesis $H_0$ against $H_1$:

$$H_0 : x_{c,i} > x_{d,i}$$
$$H_1 : x_{c,i} \leq x_{d,i}$$

(2)

where $c$ and $d$ represent the samples obtained at time $t_1$ and $t_0$, respectively. Hence, this test provides a way to order gene-expression measurements of the same gene. We say that gene $i$ is up-regulated in sample $c$ compared to sample $d$ if $H_0$ is true, and vice versa if it is down-regulated.

Continuing, when comparing gene-expression levels of two samples, typically the relative gene-expression in one sample compared to another\(^2\) is considered, that is $x_{c,i} / x_{d,i}$ for $c \neq d$ where $c, d \in C$ and $i \in I$. Since $x_{c,i} / x_{d,i}$ is not symmetric in $x_{c,i}$ and $x_{d,i}$, it is more common to take the logarithm because its absolute value is the same regardless of whether gene $i$ is up- or down-regulated by a certain fraction. Taking the logarithm is so common that the transform has its own name and its own symbols; the log-ratio and the (average) log-intensity for gene $i$ [20] is

$$M_i = \log_2 \frac{x_{c,i}}{x_{d,i}} = \log_2 x_{c,i} - \log_2 x_{d,i}$$
$$A_i = \frac{1}{2} \log_2 (x_{c,i} x_{d,i}) = \frac{\log_2 x_{c,i} + \log_2 x_{d,i}}{2},$$

(3)

where, due to the binary nature of the image-analysis quantified signals, base two has become a de facto standard, although any base would do. $M$ and $A$ are mnemonics for “minus” and “add”, respectively [16]. Estimates of the gene-expression levels may be non-positive. For this reason, the second step of equalities are included to make it explicit that in such cases the log-ratios and the

\(^2\)We here allow ourselves to think of the channels in two- or multi-channel microarray experiments conceptually as separate measurements performed in parallel.
log-intensities are not defined. Moreover, as long as the signals are positive the transform is bijective. It follows that, for up-regulated genes \( M_i > 0 \) and for down-regulated \( M_i < 0 \). Non-differentially expressed genes have \( M_i = 0 \). That is, under transform (3) the null hypothesis \( H_0 \) and the alternative hypothesis \( H_1 \) can be written as

\[
H_0 : M_i = 0 \\
H_1 : M_i \neq 0
\]  

for all genes \( i \in I \) where it is implicit that samples \( c \neq d; c, d \in C \) are considered, and analogously for hypothesis (2).

To interpret the value of \( x_{c,i}/x_{d,i} \) beyond the ordering of expression levels, additional biological assumptions have to be added. For instance, it is reasonable to assume that when the gene expression changes, the gene effect (phenotype) changes too, and that an additional small change in the same direction \emph{ceteris paribus} will further change the gene effect in the same direction. We may also assume a linear functional relationship between gene-expression level and gene effect locally. On the other hand, it is much harder to assume a linear relationship on the global scale. This is also unlikely to be true. Even worse, the biological effect of one gene is likely to be affected by the biological output of other genes and so on. Considering the huge number of genes, this illustrates the essence of the enormous task researchers in the field of functional genomics are facing trying to infer genetic regulatory networks of the cells based on gene-expression data.

4.4 Fundamental assumptions and approximations

A gene-expression experiment based on, say, microarrays, ideally gives estimates that are proportional to \( \{n_{c,i}\}_{c,i} \), but they include no estimates of \( \{N_c\}_c \) per se. However, in order to classify genes as differentially or non-differentially expressed according to (1) or (4), we have to know the relative number of cells in the two samples compared, that is, \( N_c/N_d \) for \( c, d \in C \). In practice, this is done by assuming that \( N_c \) is approximately proportional to the sum \( \sum_{i \in I} n_{c,i} \), which implies that it is assumed that all mRNA transcripts have equal (physical) weight and that at any time there is a large set of genes that are expressed. More precisely, this is typically assumed implicitly in the step of the microarray protocol where it is said that aliquot extracted and purified total amount of mRNA (or labeled
cDNA) should be used. Any microarray-based comparative gene-expression analysis relies on these types of fundamental assumptions. Without them there would be no reference available. In this thesis we will not discuss these assumptions and approximations further but assume them to be correct or at least reasonable enough.

4.5 Measurement functions

In reality, microarray experiments are much more complicated than outlined in Section 3.1, which has immediate implications on the data analysis. To summarize the model notation used in Papers D and E, let the complete microarray process from gene-expression levels \( \{x_{c,i}\}_{i \in I} \) to the corresponding set of quantified signals \( \{y_{c,i}\}_{i \in I} \) in channel \( c \) be represented by the unknown measurement function \( f_c \) such that in the error-free case we have

\[
y_{c,i} = f_c(x_{c,i}); \quad \forall c, i.
\]

(5)

To avoid complicating the model too much, we assume that \( f_c \) is the same for all genes. However, it is straightforward to extend the definition such that the measurement function is specific to disjoint subgroups of genes such as print-tip groups, plate groups, tissue-specific gene groups etc. A plate group is a set of spots that originate from the same microtiter plate, and analogously, a print-tip group is a set of spots that were printed by the same print tip. More complicated, but nevertheless possible, is to define spatially dependent measurement functions.

Note, we do assume that for any two channels \( c \neq d \), the observed signals \( y_{c,i} \) and \( y_{d,i} \) are independent given the gene-expression levels \( x_{c,i} \) and \( x_{d,i} \). A generalized definition of a measurement function that does not assume separate components is

\[
y_{1,i}, \ldots, y_{C,i} = f_C(x_{1,i}, \ldots, x_{C,i}); \quad \forall i.
\]

Note that (5) would not be true for two-color microarrays if there were spectral cross-talk between channels, which may occur if for instance excitation and emission spectra of the two fluorophores overlap. Any cross-talk is assumed to be negligible or already corrected for. We

\footnote{Alternatively one could define the gene expression to be a gene’s relative weight of mRNA transcripts compared to the total weight of all genes and in all cells, that is \( x_{c,i} = w_{c,i} / \sum_{j \in I} w_{c,j} \). where \( w_{c,i} \) is the weight of all mRNA transcripts of type \( i \) in sample \( c \). The weight \( w_{c,i} \) of all sequences for gene \( i \) depends on the number of sequences \( n_{c,i} \) and their lengths and compositions. This definition would relax some of the assumptions and approximations. However, it would at the same time move the definition of gene expression away from what we normally would mean by a gene being expressed.}
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will return to this assumption when introducing the concept of submeasurement functions below.

Avoiding quenching, scanner saturation, and other non-linear phenomena, we assume that $f_c$ is strictly increasing. In such case, a unique inverse, $f^{-1}_c$, exists and $x_{c,i}$ can be calculated by the back transform (still assuming no errors)

$$x_{c,i} = f^{-1}_c(y_{c,i}).$$

(6)

4.5.1 Testing for differentially expressed genes

Returning to our test for differentially expressed genes and assuming zero noise, we see that when the measurement functions are identical, that is, when

$$f = f_c = f_d; \ c, d \in C,$$

(7)

the set of non-differentially expressed genes can be expressed as $I_{x_c=x_d} = \{i \in I : x_{c,i} = x_{d,i}\} = \{i \in I : f^{-1}_c(y_{c,i}) = f^{-1}_d(y_{d,i})\} = \{i \in I : y_{c,i} = y_{d,i}\}$. If $f_c \neq f_d$, the last equality will not hold anymore.

4.5.2 Linear and affine measurement functions

A common assumption, implicit or explicit, is that the measurement function is linear (with zero intercept);

$$f_c(x_{c,i}) = b_c x_{c,i} \iff f^{-1}_c(y_{c,i}) = \frac{y_{c,i}}{b_c}$$

(8)

for channel $c$ and gene $i$ where $b_c \neq 0$. This measurement function simplifies downstream statistical analysis and allows direct interpretation of the observed signals, that is, the signals are assumed to be proportional to the corresponding gene-expression levels. However, it does not capture systematic effects that express themselves as channel biases, which for instance is the case with imperfect background correction, scanner biases, etc. The observed log-ratios and log-intensities under a linear transform are

$$M_i = \log_2 \frac{f_c(x_{c,i})}{f_d(x_{d,i})} = \log_2 \beta + \log_2 \frac{x_{c,i}}{x_{d,i}}$$

$$A_i = \frac{1}{2} \log_2 (f_c(x_{c,i}) f_d(x_{d,i})) = \frac{1}{2} \log_2 (b_c b_d) + \frac{1}{2} \log_2 (x_{c,i} x_{d,i})$$

(9)
where $\beta = b_c/b_d$ and $b_c, b_d \neq 0$. Thus, the linear transform introduces a bias $\log_2(\beta)$ in the log-ratios, which is seen as a (constant) vertical shift in the log-ratio versus log-intensity plots. See Figure 2 for an example. There is also a (constant) shift in the log-intensities, which is of less importance since the log-ratios are the quantities of interest.

Figure 2: Example of log-ratio versus log-intensity scatter plots between channels $\{R, G\}$ where a linear transform introduces a constant bias in the observed log-ratios. The same data set is used in all cases. **Left:** There is a balanced scale in the channels, that is, $\beta = b_R/b_G = 1$. **Middle:** Relative scale is $\beta = 4/1$, which introduces a $\log_2\beta = +2$ shift in the log-ratios. **Right:** A relative scale $\beta = 1/4$ introduces a $\log_2\beta = -2$ shift. The data points are colored on a red-to-yellow-to-green scale corresponding to the observed log-ratios and a dark-to-bright scale corresponding to the observed log-intensities to imitate the (false) colors of the spots as seen in typically two-color microarray images.

The second simplest measurement function to consider, which is also more powerful, is the affine measurement function

$$f_c(x_{c,i}) = a_c + b_c x_{c,i} \iff f_c^{-1}(y_{c,i}) = \frac{y_{c,i} - a_c}{b_c} \quad (10)$$

where $b_c \neq 0$. The affine transformation can be motivated physically, biologically, and mathematically; see Papers D and E. To continue, the observed log-ratios
under an affine transform are
\[ M_i = \log_2 \frac{f_c(x_{c,i})}{f_d(x_{d,i})} \]
\[ = \log_2 \beta + \log_2 \frac{\alpha^2 + \frac{x_{c,i}}{x_{d,i}} \beta^2 A_i}{\alpha^2 + \frac{x_{c,i}}{x_{d,i}} \beta^2 A_i} + \log_2 \frac{x_{c,i}}{x_{d,i}} \] (11)

where \( \alpha_i = a_c - \beta a_d (x_{c,i}/x_{d,i}) \) and \( A_i \) is the observed log-intensity for gene \( i \).

In addition to the constant bias seen in the linear transform there is a bias in the log-ratios that depends on the size of true relative gene-expression ratio \( x_{c,i}/x_{d,i} \) and the (observed) log-intensities \( A_i \), cf. Figure 3. See Paper D for details. The existence of this non-linear intensity-dependent bias complicates test (4) for non-differential expression. Of less importance, there is also an intensity and relative-gene-expression dependent bias in the observed log-intensities.

Figure 3: Example where an affine transform introduces an intensity-dependent bias in the observed log-ratios. The same data set with relative scale \( \beta = 1/1 \) is used in all cases. Left: No bias in either channel. Middle: A bias of size 20 in channel \( R \), that is \((a_G, a_R) = (0, 20)\). Right: An additional bias of size 200 in the other channel so that \((a_G, a_R) = (200, 20)\).

4.5.3 Submeasurement functions

The concept of a measurement function can be applied to any part of the microarray process, not only the complete process from gene expression to the quantified
signals in the computer. For instance, as done in Paper E, we can choose to focus on the submeasurement function describing the scanner and the image-analysis steps. This is also illustrated in Figure 1 where the complete measurement function $f_c$ is split up into two subsequent submeasurement functions $g_c$ and $h_c$, that is, $f_c = h_c \circ g_c$ assuming the two channels $C = \{R, G\}$ act separately. Submeasurement functions can be defined and studied at any resolution. Furthermore, when investigating single submeasurement functions alone, the assumption (5) that the measurement functions are separate between channels may be less questionable or at least has less impact on the final conclusions. Notice that linear and affine submeasurement functions $h_c$ and $g_c$ are closed under composition, which implies that $f_c$ belongs to the same class. This is very convenient, not only in theory but also in practice, because it allows some normalization methods to be applied in any order, which makes it possible to postpone some required decisions to a later step in the analysis process.

4.6 Calibration and normalization

Systematic effects or artifacts in the observed signals are observed when the measurement functions are non-linear. As exemplified in Section 4.5.2 and illustrated in Figure 3, calibration and normalization is about identifying and removing such unwanted bias in the observed signals in order to reveal the true relative gene-expression levels.

4.6.1 Calibration

When a parametric model for $f_c$ is used and data is of the form $\{(x_{c,i}, y_{c,i})\}_i$, the measurement function $f_c$ can be computed in the noise-free case. The true expression levels are then recovered through

$$\hat{x}_{c,i} = \hat{f}_c^{-1}(y_{c,i}); \forall c, i,$$

(12)

where $\hat{f}_c$ is the computed version of $f_c$. We refer to a method that calculates the calibrated signal $\hat{x}_{c,i}$ by (12) as a calibration method. An example is when known amounts of certain RNA, also known as spike-ins, for a set of genes $I_{\text{known}}$ are added to the original set of RNA transcripts, then measured and observed in

\footnote{Without noise, we often have $\hat{f}_c = f_c$. However, we also regard the case when $f_c$ is known only up to a multiplicative constant ($f_c \propto f_c$) as a calibration method. In that case $\hat{x}_{c,i} \propto f_c^{-1}(y_{c,i}) = x_{c,i}$. This is because the absolute scale of $x_{c,i}$ is of subordinary interest.}

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order to obtain \( \{(x_{c,i}, y_{c,i})\} \) for \( i \in \mathcal{I}_{\text{known}} \). With classical regression techniques, the measurement function \( \hat{f}_c \) for the spike-ins can be estimated by \( \hat{\hat{f}}_c \). With the assumption that the other genes have the same measurement function, that is \( f_c = \hat{f}_c \), we have \( \hat{\hat{f}}_c = \hat{f}_c \) and all \( x_{c,i} \) can be estimated from (12). For example, with known expression levels and observations of the spike-ins, and assuming an affine transformation, the bias and scale parameters \( a_c \) and \( b_c \) in channel \( c \) can be estimated, respectively. Assuming the same parameters for the measurement function of all other genes, estimates of the gene-expression levels can be obtained.

In Paper E, another type of calibration method is discussed. It does not estimate the complete measurement function, but instead a submeasurement function specific to the scan and image analysis steps of the microarray process. For the analysis of that paper, there are no known data points \( \{(x_{c,i})\} \) available, but the data points are known to be fixed between repeated measurements at different sensitivity levels. We impose an additional assumption about the measurement function stating that no matter what sensitivity level \( k = 1, \ldots, K (K \geq 2) \) is used the bias is the same;

\[
y_c^{(k)} = f_c^{(k)}(x_{c,i}) = a_c^{(k)} + b_c^{(k)} x_{c,i} = a_c + b_c^{(k)} x_{c,i}, \quad \forall c, i, k.
\]  

Here superscript \((k)\) denotes sensitivity level with index \( k \). The sensitivity is adjusted by modifying the photomultiplier tube (PMT) gain. With this additional constraint, which can be motivated physically and is strongly supported by data, it is possible to infer the submeasurement function based solely on the observations \( y_c^{(k)} \) (up to a scale factor). Thus, in the absence of noise, the signals \( x_{c,i} \) of interest can be estimated up to a multiplicative constant, which is the reason why this method is a calibration method.

4.6.2 Normalization

Even if no calibration data points such as spike-ins or fixed data points are available, it is still possible to infer something about the measurement functions. Consider a two-sample experimental setup for which it is possible to identify a subset of non-differentially expressed genes, \( \mathcal{I}_{\text{norm}} = \{i \in \mathcal{I} : x_{c,i} = x_{d,i}\} \subset \mathcal{I}_{x_c = x_d} \) for \( c \neq d \). This is typically done by assuming that most genes are non-differentially expressed. With this set we can find a smooth normalization function such that \( g_c^{-1}(y_{c,i}) = g_d^{-1}(y_{c,i}) \) for \( \mathcal{I}_{\text{norm}} \). This is analogue to the above where we used
separate calibration functions for each channel. Next, assume that this is true for all non-differentially-expressed genes, and otherwise not, that is,

\[ x_{c,i} = x_{d,i} \iff g_c^{-1}(y_{c,i}) = g_d^{-1}(y_{d,i}) \]  \hspace{1cm} (14)

for \( i \in I \). A normalization method is then a method that transforms the signals as

\[ \bar{y}_{c,i} \leftarrow g_c^{-1}(y_{c,i}); \forall c, i. \]  \hspace{1cm} (15)

When the channels are normalized separately, orthogonality between channels will be preserved, otherwise artificial cross-talk will be introduced. We immediately note that \( g_c^{-1} \) can be chosen as the inverse of the measurement functions \( f_c^{-1} \), if known. In other words, calibration is a special case of normalization. Moreover, note that it is always possible to choose, say, \( g_c \) as the identity function such that \( x_{c,i} = x_{d,i} \iff y_{c,i} = g_d^{-1}(y_{d,i}) \) for \( i \in I \). As an example, consider the affine model (10) with two channels \( c \) and \( d \), and suppose there are no fixed or spike-in data points available. Then \( a_c, b_c, a_d \) and \( b_d \) cannot be estimated. However, without noise we have

\[ y_{c,i} = \alpha_i + \beta x_{c,i} x_{d,i} \]  \hspace{1cm} (16)

with \( \alpha_i \) and \( \beta \) as in (11). Hence, if there is a known set \( I_{\text{norm}} \) of non-differentially expressed genes, the corresponding points \( \{(y_{c,i}, y_{d,i})\} \) are located along a line with slope \( \beta \) and intercept \( \alpha = a_c - \beta a_d \). Therefore, the parameters \( \alpha \) and \( \beta \) are known (or can be estimated when noise is present). Hence, the normalized signals

\[ \bar{y}_{c,i} = y_{c,i} = a_c + b_c x_{c,i} \]
\[ \bar{y}_{d,i} = \alpha + \beta y_{d,i} = a_c + b_c x_{d,i} \]  \hspace{1cm} (17)

are computable and satisfy (14). Moreover, it follows from (11) that the intensity-dependent bias is eliminated (\( \alpha_i = 0 \)) for the log-ratios \( M_i = \log_2(\bar{y}_{c,i}/\bar{y}_{d,i}) \) of all non-differentially expressed genes.

Another example of normalization is given in Paper E (and D). There we show that for some scanners there exist PMT-independent (scale-independent) biases \( a_c^{(k)} = a_c \) as in (13). Thus, by adjusting the gain of the two PMT and hence the scale parameters \( b_c \) and \( b_d \), it is possible to find a configuration such that
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Figure 4: Example where an adjustment of the gain (scale parameters) can normalize affinely-transformed signals, that is, remove the curvature. The same data set is used in all cases. Left: No bias and equal scale in both channels. Middle: The same data with channel biases \((a_G, a_R) = (200, 20)\) and equal scale \((\beta = 1)\). Right: The same biases now compensated by the relative scale \(\beta = 1/10\). A constant bias in the log-ratios remains, but there is no curvature for the non-differentially expressed genes.

\[ \alpha_i = 0 \text{ for } i \in I_{x_c=x_d} \text{ in Equation (11) resulting in no curvature for the non-differentially genes.} \]  
Most likely this is done in quite a few laboratories when the arrays are pre-scanned and the PMT settings are adjusted such that the two channels are "balanced". This rather mechanical normalization procedure is illustrated in Figure 4. Note, if the PMT-independent bias would be zero, adjusting the PMT gains would just shift the log-ratios the same amount up or down for all intensities as in Figure 2. Having said this, a quick and simple test for linearity of a scanner's submeasurement function is then also given.

It follows immediately from (14) that normalized signals can be used to test for non-differential expression; testing \(H_0\) against \(H_1\) in (1) is the same as testing \(H'_0 : \tilde{y}_{c,i} = \tilde{y}_{d,i} \text{ against } H'_1 : y_{c,i} \neq y_{d,i} \text{ for } c \neq d\). However, testing \(H_0\) against \(H_1\) in (2) is not necessarily the same as testing \(H'_0 : \tilde{y}_{c,i} > \tilde{y}_{d,i} \text{ against } H'_1 : y_{c,i} \leq y_{d,i} \text{ for } c \neq d\). Although the latter is less likely to be a problem in reality, more problematic is the ordering of relative gene-expression levels. Observing \(\tilde{y}_{c,i_1} / \tilde{y}_{d,i_1} > \tilde{y}_{c,i_2} / \tilde{y}_{d,i_2}\) for \(i_1, i_2 \in I\) does not imply \(x_{c,i_1} / x_{d,i_1} > x_{c,i_2} / x_{d,i_2}\). To illustrate this with a real-world example, consider a two-channel microarray experiment with replicated spots \(i_1\) and \(i_2\) of the same gene \(i\). Although the underlying relative expression levels for the two spots are identical by definition,
due to a non-linear measurement function and differences in spot concentrations, the normalized log-ratio can be greater for spot $i_1$ than for spot $i_2$, which becomes a problem when (if) averaging log-ratios. This is why we differentiate between calibration and normalization.

Other normalization methods such as quantile normalization methods and curve-fit normalization methods can be seen in the light of (14) and (15). For instance, a curve-fit (intensity-dependent) normalization method normalizes the signals by assuming $M_i = h(A_i)$ for all non-differentially expressed genes. The normalization method in this case differs slightly from (15) and acts jointly on the two channels $c$ and $d$ through the log-ratio. The normalized log-ratios are $\tilde{M}_i = M_i - h(A_i)$, where $h$ is estimated using a robustified version of some nonparametric smoothing method such as local polynomial regression (for instance loess or smoothing splines). Putting $\tilde{y}_{c,i} = 2^{-0.5h(A_i)}y_{c,i}$ and $\tilde{y}_{d,i} = 2^{0.5h(A_i)}y_{d,i}$, we find that $x_{c,i} = x_{d,i}$ is equivalent to $\tilde{y}_{c,i} = \tilde{y}_{d,i}$ for $i \in I_{\text{norm}}$.

### 4.7 Error models and error propagation

Error terms are still to be discussed. Any measurement contains noise

$$y_{c,i} = f_c(x_{c,i}) + \varepsilon_{c,i}$$

(18)

where $\{\varepsilon_{c,i}\}_{c,i}$ are independent and unobserved zero-mean error terms for $i \in I$ and $c \in C$. With noise, the estimate (12) of $x_{c,i}$ contains two sources of error: the possible estimation error of $\hat{f}_c$ (as an estimate of $f_c$) and the presence of $\varepsilon_{c,i}$ in (18). For this reason, $\hat{x}_{c,i}$ will usually depart from $x_{c,i}$ even when $f_c$ is estimated with good accuracy. For normalization methods we may still use (15) provided $g_c^{-1}$ is replaced by an estimate $\hat{g}_c^{-1}$. However, formula (14) is only valid in the noise-free case although it still serves as a guiding principle for defining the normalization method.

With error terms, the test of $H_0$ against $H_1$ becomes a statistical test; the region were $H_0$ is rejected is a region defined by the distribution of the error terms.

Various error models have been suggested. The most naive one assumes independent and identical distributed (i.i.d.) normal error terms for all genes, say, $\varepsilon_{c,i} \in N(0, \sigma^2_c)$; $\forall c$. The exact error model depends on the which (sub)measurement function is studied, but typically a heteroscedastic model is more useful, that is,
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A model where the standard deviation of the noise is a function of the signal. An example is

$$\sigma_{c,i}(x_{c,i}) = \delta_c + \lambda_c x_{c,i}$$

(19)

where $\delta_c \geq 0$ and $\lambda_c \geq 0$. Another error model with additive and multiplicative noise was suggested by [11, 12] and utilized in [4, 5]. The latter two suggest variance stabilizing models that also contain an affine component. Depending on measurement function and noise structure, the errors propagate differently resulting in a more or less complicated noise structure of, say, the log-ratios, which in turn affects the region of rejection in a log-ratio-based hypothesis test. This thesis does not explicitly study error propagation further, but Paper G suggests a heteroscedastic affine error model for microarray data with dilution series that allows us to identify a calibration function. See also Section 6.7, which gives a brief overview of this paper. More research on error models for microarrays is needed.

4.8 The search for an omnibus way to compare calibration and normalization methods

Quite a few normalization methods and some calibration methods for single-channel and two-channel microarray data have been published to date. For a description of some of them, see Paper D. Depending on the error structure, a test of $H_0$ against $H_1$ in (1) based on normalized signals may be as simple (or as hard) to conduct as a test based on calibrated signals. However, it is our belief that in general a correct calibration method is to prefer over a normalization method, because the former tries to estimate the true gene-expression levels (up to a scale factor) contrary to the latter. Not to be forgotten, calibration allows us to test $\tilde{H}_0$ against $\tilde{H}_1$ in (2). More important, though, is how to know which calibration or normalization method to use. This calls for quantitative methods in order to compare any two normalization or calibration procedures. Unfortunately, no generic method has been presented to date. It is still an open problem which calibration or normalization method is optimal for a given data set.

A reasonable criterion for a normalization method to be good is that normalized signals from repeated measurements of identical samples should be “as similar as possible”. Statistically we say that the variability (or variance) of the normalized
signals should be small (or equivalent, that the precision should be high). In addition to this, small bias (high accuracy; "closeness to the truth") of the estimated signals is preferred, although not required by all tests. For example, consider a single two-channel microarray where each gene has been replicated \( K \) times for which we obtain log-ratios \( M_i = (M_{i,1}, \ldots, M_{i,K}) \) for each gene \( i \). Let \( M = \{M_i\}_i \) be the set of all log-ratios. The average sample variance of all genes is

\[
\hat{\sigma}^2(M) = \frac{1}{I} \sum_{i=1}^{I} \hat{\sigma}^2_i(M_i) \tag{20}
\]

where \( \hat{\sigma}^2_i(M_i) \) is the sample variance of the log-ratios for gene \( i \). Let this be our precision measure. Next, assume that we apply two different normalization methods, called A and B, to the same data set. This will result in two sets of normalized log-ratios, \( \tilde{M}^A = \{\tilde{M}^A_i\}_i \) and \( \tilde{M}^B = \{\tilde{M}^B_i\}_i \). By comparing \( \hat{\sigma}^2(\tilde{M}^A) \) and \( \hat{\sigma}^2(\tilde{M}^B) \) we then wish to say which of the two methods A and B that performs best on the given data set. However, we cannot use (20) alone for this purpose. One reason may be heteroscedasticity of the log-ratios. Another is that (20) is not invariant under affine transformations. To illustrate this, consider the case where method A is a "perfect" calibration method in the sense that there are no systematic effects remaining, and where method B is such that is first does what method A does, but then adds a large constant to both channels. The absolute value of the log-ratios for the latter method will decrease as the added channel bias gets larger resulting in \( \hat{\sigma}^2(\tilde{M}^B) < \hat{\sigma}^2(\tilde{M}^A) \), although method A is the best by definition. A statistician would of course immediately object and remind that we do not control for bias. The accuracy of \( \tilde{M}^B \) will of course decrease as it departs from the true values. However, the problem in microarray analysis is that the truth is almost never known and therefore we have to rely on other methods. We have chosen the average standard deviation as an example because we know it is indeed used for the purpose of comparing normalization and calibration methods. In Paper B, we used a similar measure ourselves. Similar concerns occur when trying to define a distance space for clustering and classification.

A possible solution is to use a variance stabilizing transform such as [4, 5] before calculating (20). However, as these transforms are normalization methods themselves, it is not clear how to use them for the purpose of comparing signals normalized by other means. This is an open problem and more research is needed.
5. Making methods available to the research community

This thesis neither discuss nor provide answers to the above, but we hope that by better understanding measurement functions for microarrays we are one step closer to an answer. In the meantime, we have to rely on subjective judgements based on visual comparisons of log-ratio versus log-intensity plots, density plots, summary statistics, comparisons of “top 100 gene lists”, comparisons to measurements from other techniques, and most importantly, biological validation of the results.

5 Making methods available to the research community

Since the end of year 2000, we have developed an open-source microarray analysis package in R [6] called aroma (formerly known as com.braju.sma) for the purpose of making existing and new statistical tools easily accessible to life science researchers. It is aimed to be user-friendly and is used by international research groups [19, 2] and by academia who in turn give us most valuable practical (tests and bug reports) and scientific feedback. In addition to this, aroma provides us with a structured and expandable platform to implement new statistical algorithms and compare them with published ones. The maintenance of this project has been and is still tightly coupled to computationally intensive projects of our own. Aroma pre-dates the promising Bioconductor.org project [18], which is one reason for why it is currently not available via that forum. We hope to be able to conform toward their standards and conventions in a way that is backward compatible with our standards [3] (and Paper C) and with users’ existing aroma scripts. We plan to broaden the access to our statistical tools further by making the package available as a plug-in for the BioArray Software Environment (BASE) [13]. Part of this work has already been initiated by the group responsible for BASE. With the upcoming release of BASE v2 the opposite will also be true in which all of BASE’s database and analysis tools can be accessed from within R and aroma via a Java application programming interface (API). The aroma package is presented in Paper F.

6 Summary of the papers

A brief presentation of the papers presented in this thesis follows. Naturally, our knowledge on microarray analysis has evolved in the course of our research, which
is also reflected in the papers. The papers are listed in chronological order.

6.1 Paper A

Paper A was written as a complementary article to an online discussion at Science’s Science Aging Knowledge Environment (SAGE KE). It was intended to give a short introduction of basic normalization methods and statistical tests for microarray data to an audience consisting of experimentalists. The one-sample t-test and the one-sample empirical Bayes’ test (“B-test”) [10] are explained and a simple empirical comparison of the two tests is performed illustrating how different methods and different number of arrays give different results. A simple comparison of the sets of identified genes from applying two different image analysis methods was also conducted. Data was provided by Matt Callow and Eric Rubin. Saira Mian contributed with the introduction and Terry Speed contributed with valuable scientific feedback. Brent Calder did the Axon GenePix image analysis and replicated the analysis done for the Spot image-analysis data, which was done by Henrik Bengtsson who also summarized the results and wrote the remaining text.

6.2 Paper B

Paper B is about print-order and plate effects in spotted microarray data. It introduces a novel way of displaying microarray data called print-order plots. When producing spotted microarrays the spots are not printed simultaneously, but subsequently in groups of size equal to the number print tips used by arrayer. The arrayer does a so called source visit, fills the print tips with the DNA clones in the microtiter-plate wells, and moves to the area where the glass arrays are mounted. After that, it will print a set of spots at once on the array, then move over to the second array and so on. The amount of DNA picked up at a single source visit will often last 50-100 arrays before a new source visit is needed. When the first round is completed another set of clones are picked up and printed on array one, two and so on until the arrays are filled. For this reason, it will take many hours to print the arrays, and, if the printing environment is not controlled well enough, the properties of the spots may vary over time. Thus, by displaying, say, the observed log-ratios in the order that the corresponding spots were printed, such abnormalities may be detected. The print-order plots illustrated in the paper show strong apparent print-order effects.
6. Summary of the papers

Different sequences of normalization methods applied to all genes or subset of genes are applied and the results are visually compared. In order to compare normalization strategies objectively, the paper suggest a measure of reproducibility (MOR), which is a precision measure of how dissimilarly replicated log-ratios are across arrays similar to how (20) is defined. Using this measure, one of the optimal strategies found was to apply a print-tip lowess normalization followed by a plate-group lowess normalization. Comparing the MOR before and after normalization, and comparing with print-tip lowess normalization alone, most of the systematic effects were removed by print-tip normalization (the MOR decreased from 100% to 46%) and succeeding plate-group lowess normalization would remove a little bit more (MOR decrease to 41%).

It turns out that the majority of the print-order effects seen in the data set studied are actually “fake” visual artifacts. They are due to the fact that different types of clones originating from different sets of microtiter plates were printed at different times. As shown in one of the figures in the paper, the brain clones were spotted during one day and liver clones during another day with the effect that the data points from the two groups are located in separate parts of print-order plots. Moreover, since the experiment was done with RNA extracted from liver, the genes corresponding to the liver clones were strongly expressed whereas the brain clones were hardly expressed at all. Together with the fact that there was a strong intensity-dependent effect in the log-ratios, the brain-clone signals became highly up-regulated at the same time as the strongly expressed liver genes were less affected by the intensity effect giving only little bias. The results also “indicate” that doing background subtraction is worse because it gives a higher MOR compared to when it is not applied. In reality this may or may not be true, but the indications are probably false due to reasons discussed above. This is an example where it is important to have an intensity-invariant precision measure. However, we still believe it is fair to compare MOR values for background and non-background corrected signals separately. We have not revisited this analysis with affine normalization procedures.

6.3 Paper C

Paper C presents the R package R.oo, which provides the object-oriented core used in the microarray aroma package, which all of our applied microarray re-
search is based on. R.oo extends the S3 method dispatching mechanism in R with an object-oriented layer that provides references. By implementing support for reference variables, it allows a more user-friendly and memory efficient, but also more developer-friendly design. Reference variables are controversial, because they break the boundaries of the otherwise functional nature that the S language was intended for\(^5\). In spite of this, it provides us with the necessary language foundation for developing and maintaining the aroma package. Many other packages, not published here, rely on R.oo. It has proven to be very useful.

### 6.4 Paper D

Paper D conducts a detailed study of affine transformations of two-channel microarray data and their effects on the log-ratio transform. Many common normalization methods are investigated with respect to this transform. We conclude the study by suggesting a robust nonparametric normalization method for affinely transformed microarray data. With channels \( C = \{ R, G \} \), \( \alpha = a_R - \beta a_G \) and \( \beta = b_R/b_G \) as in (11), we have for non-differentially expressed genes (without noise) that

\[
y_{R,i} = \alpha + \beta y_{G,i},
\]

which is a special case of (16). Define the observation vector \( y_i = (y_{G,i}, y_{R,i}) \). With noise (18) included, the points \( \{ y_i \}_{i=1}^I \) are scattered around a line \( L(\alpha, \beta) \) in \( \mathbb{R}^2 \) with intercept \( \alpha \) and slope \( \beta \). In order to estimate \( \alpha \) and \( \beta \), we define the objective function

\[
Q(\alpha, \beta; y) = \sum_{i=1}^I w_i r_i(\alpha, \beta; y_i)^2
\]

where \( r_i(\alpha, \beta; y_i) > 0 \) is the orthogonal Euclidean distance between \( y_i \) and the line \( L(\alpha, \beta) \). The estimate of \( \alpha \) and \( \beta \) is then

\[
(\hat{\alpha}, \hat{\beta}) = \arg\min_{(\alpha, \beta)} Q(\alpha, \beta; y).
\]

\(^5\)As a comment, we would like to note that the same functionalities that the R.oo core is based on is added to the data type environment in R v1.9.0, which in some sense should make R.oo less controversial.
Different weight functions minimize the orthogonal distances in different norms. For \( w_i = 1 \) minimization is done in \( L_2 \), just like principal component analysis (PCA), using \( w_i = 1/(r_i(\hat{\alpha}, \hat{\beta}; y_i) + \delta) \) minimization is done in \( L_1 \) (if we let \( \delta \to 0^+ \)), and so on. The above minimization problem was implemented using iteratively reweighted PCA (IWPCA) and the method is very fast. Since the expression levels \( x_{c,i} \) and \( x_{d,i} \) are unknown, \( a_R \) and \( a_G \) are not individually identifiable. As described in Section 4.6.2, we obtain a normalization method by plugging the estimated parameters \( \hat{\alpha} \) and \( \hat{\beta} \) into (17) and then computing the estimated log-ratio \( \tilde{M}_i = \log_2(\tilde{y}_{c,i}/\tilde{y}_{d,i}) \). In order to make the bias and scale parameters \( a_e \) and \( b_e \) identifiable, we suggest additional constraints in the paper. The method generalizes directly to multiple channels and even multiple arrays, which therefore makes so called between-array (scale) normalization unnecessary.

6.5 Paper E

Paper E gives evidence for existence of affine submeasurement functions in real microarray data. We investigate two different brands of scanners together with two different image-analysis applications and we find that the scanners introduce a significant bias in both channels. The results show that the bias is most likely introduced by the PMT and/or the following electronics such as the analog-to-digital converter. The image-analysis methods seem to introduce a small bias too, but of subsidiary order. As described in Section 4.6 above, by scanning the arrays at different PMT settings, using model (13) the channel biases can be uniquely identified. The estimation of bias and scale parameters is done by a modified version of the IWPCA method presented in Paper D. Model (13) is strongly supported by data. We refer to this method as the multiscan calibration method. Because the same array is scanned multiple times and the calibrated signals are averaged, the effective measurement noise is decreased and the effective dynamical range of the scanner is increased.

6.6 Paper F

Paper F describes the aroma (An R Object-oriented Microarray-Analysis environment) package discussed in Section 5. The package alone is written in more than 15000 lines of R code and has a 280-page manual with many examples. It is under continuous development.
6.7 Paper G

Paper G suggests a calibration method for spotted microarray data containing dilution series (replicated probes of various concentrations). The existence of dilution series makes it possible to identify a calibration function, instead of just a normalization function. The method is based on an affine heteroscedastic model. Let \( y_{c,i,j} \) be the observed expression level for replicate \( j \) of gene \( i \) in channel \( c \). We assume that

\[
y_{c,i,j} = a_c + b_c b_{i,j} x_{c,i} + \varepsilon_{c,i,j}; \quad \forall c, i, j
\]  

(24)

where \( a_c \) is the channel bias, \( b_c \) is the channel gain with \( b_1 = 1 \), \( b_{i,j} \) is the probe concentration for replicate \( j \) of gene \( i \), and \( \varepsilon_{c,i,j} \) is independent noise with \( E[\varepsilon_{c,i,j}] = 0 \) and \( V[\varepsilon_{c,i,j}] = \sigma^2_{c,i,j} \). A variance function that is quite flexible and at the same time generates feasible parameter estimates is

\[
\sigma_{c,i,j} = k_c \sigma_i; \quad \forall c, i
\]  

(25)

where \( k_c \) is the relative standard deviation in channel \( c \) with \( k_1 = 1 \) for identifiability. Recalling Section 4.7, contrary to a model with noise structure as in (19) and [11], this model assumes that the standard deviation is the same for all replicates regardless of spot concentration at the same time as it allows it to vary between genes. The model parameters are estimated with maximum likelihood (ML) techniques utilizing profile-likelihood based grid search and PCA. In the paper, we study properties of bias parameter estimates \( \hat{a}_c \) for microarray data simulated from model (24)-(25). We compute confidence intervals and standard errors for estimates of \( a_c \) using both bootstrap and analytical methods. The latter are based on normal approximations of the parameter estimates and computation of Fisher information of the structural parameters \( \{a_c\} \), treating all other parameters as nuisance parameters. The estimated \( a_c \) is used for calibration of the probe signals. As a direct implication of the model design, estimates of the relative expression levels of the dilution series (genes) follow. For two channels, this calibration can either be defined through the replicate-specific log-ratios

\[
\hat{M}_{i,j} = \log_2 \frac{y_{2,i,j} - \hat{a}_2}{y_{1,i,j} - \hat{a}_1}
\]  

(26)

\(^6\)Note the differences in notation here and in the paper.
or the gene-specific version

\[ \hat{M}_i = \log_2 \frac{\sum_j w_{i,j}(y_{2,i,j} - \hat{a}_2)}{\sum_j w_{i,j}(y_{1,i,j} - \hat{a}_1)}. \] (27)

Here \( w_{i,j} \) are weights chosen by the user, for instance \( w_{i,j} = 1 \) or \( w_{i,j} = w(b_{i,j}) \), in case the latter are known. We also estimate relative expression levels \( \beta_{c,i} = (b_{c,b_{i,j}x_{c,i}})/(b_{1,b_{i,j}x_{1,i}}) = b_{c}(x_{c,i}/x_{1,i}) \). For two channels, this defines another calibration method

\[ \hat{M}_i = \log_2(\hat{\beta}_{2,i}). \] (28)

The log-ratios (26)-(28) are estimates of the true log-ratios up to an additive constant \( \log_2 b_2 \). Compared to (27), (28) has the advantage that standard errors and confidence intervals for \( \hat{M}_i \) can be provided. These may be computed using either parametric bootstrap or analytical methods based on Fisher information and normal approximation. We also compare different types of dilution series \( b_{i,j} \). Finally, we investigate robustness toward model misspecification in that we simulate data from (24), but with a variance function different from (25) and close to (19).
References


Paper A

Identifying differentially expressed genes in cDNA microarray experiments: making aging visible

H. Bengtsson1, B. Calder2, I.S. Mian3, M. Callow4, E. Rubin4, T.P. Speed5,6

1 Mathematical Statistics at the Centre for Mathematical Sciences, Lund University, Lund
2 Health Science Center at San Antonio, The University of Texas, Texas
3 Genome Sciences Department, Lawrence Berkeley National Laboratory, Berkeley
4 Radiation Biology and Environmental Toxicology, Life Sciences Division, Lawrence Berkeley National Laboratory, Berkeley
5 Department of Statistics, University of California, Berkeley
6 Division of Genetics and Bioinformatics, Walter and Eliza Hall Institute of Medical Research, Melbourne

This is a version of an online report accompanying the Science’s SAGE KE discussion forum with the same title.

1 Introduction

Transcription profiling monitors simultaneously the abundance of tens of thousands of transcripts in a biological sample. In biogerontology, such profiling is but one arrow in the quiver required to ascertain how the physiological vigor of an organism declines over time. Currently, expression profiling technology comes in a variety of flavors including cDNA microarrays, high-density nylon membrane arrays, serial analysis of gene expression, short or long oligonucleotide arrays, fiber optic arrays, and so on [17]. The exponential rise in the number of profiling-related MEDLINE articles is a testament to its popularity. If this experimental tool is to deliver on its promises, however, a growing list of statistical, computational, technological and biological issues will need to be addressed (for incomplete lists, see [18, 15]).

This discussion forum focuses on the elemental, but critical issue of statistical approaches to identifying differentially expressed genes given cDNA microarray data.
(it should be noted that many of the same issues arise with the other technologies). The nature and type of data to be analyzed as well as a need to distinguish between biologically and statistically significant differential expression highlights the multiplicity of potential answers to such a deceptively simple question.

The starting premise for the forum is that a suitable scientific question has been formulated, the appropriate study designed, standard operating procedures employed to extract mRNA samples from relevant specimens, the transcription profiling experiments performed, and images of hybridized microarrays acquired. Accounting for technological and/or experimental confounding factors can reduce observation noise, errors associated with the measurement process or technology itself, but has little influence on model noise, variation due to the stochastic nature of gene expression and the underlying biology. It is assumed that strategies for minimizing observation noise and data analysis are independent of the scientific question. That is, aging is sufficiently similar to cancer, development and other biological processes so that no “new” statistical methods need to be invented to analyze transcription profiling data from biogerontology studies. Thus, employing a study of mouse lipid metabolism in mice ([7]) as the framework within which to structure the discussion forum and to illustrate pertinent issues seems appropriate.

The background corrected fluorescent intensities as well as the eight original scanned images processed using the Spot software [1] for the ApoAI experiments can be downloaded from [8]. The statistical analyses described on this page were performed using the object-oriented add-on com.braju.sma [4] to the sma (Statistics for Microarray Analysis) package [6] written using the R language [13]. How it can be done in R is described in the R program [5].

The outline of this article is as follows. In Section 2, we describe the setup of the experiments of the study and in Section 3 we shortly discuss the problem of extracting signal from scanned images. In Section 4, we will identify systematic variations in data that do not have a biological source, but are believed to stem from different error sources along the cDNA microarray process. We will show how different normalization methods can correct for these artifacts. In Section 5, we discuss the problems and methods on how to identify differentially expressed genes. Throughout the article we use the ApoAI data set to exemplify the different ideas. In Section 6, we give a list of 40 genes that we identified to be differentially
expressed. At the end, we also do a comparison between the genes that are found when Spot is used and those that are found when GenePix is used.

2 Experimental setup

The goal of the study of mouse lipid metabolism in mice was to identify genes with altered expression in two mouse models with very low HDL cholesterol levels (treatment groups) compared to inbred control mice. The first mouse model compared eight mice with the apolipoprotein AI (ApoAI) knocked-out with a control group consisting of eight “normal” C57B1/6 mice. The second mouse model compares in a similar way eight scavenger receptor BI transgenic mice (SRBI) with the same control group. To identify the altered gene expressions a cDNA microarray technique was used. For the first model, target cDNA was obtained from mRNA for all sixteen mice by reverse transcription and labeled using the red-fluorescent dye Cy5. The reference sample was prepared by pooling cDNA from the eight control mice and was labeled with the green-fluorescent dye Cy3. An analog setup was used for the second mouse model. The mix of target and reference cDNA was hybridized to microarrays containing \( N = 5548 \) DNA probes. The slides contained 6384 spots, but some of them were empty spots. Here we will focus on the comparison between the eight ApoAI mice \((K = 8)\) and the pool of control mice without making use of the fact that a comparison between each of the eight control mice with a pool of the same mice was also done. To include the eight slides from the control group in the analysis both two-sample \( t \)-tests and adjusted \( p \)-values are required, an approach which is beyond the scope of this article. For an analysis of the ApoAI data set using two-sample \( t \)-tests, see [11].

3 Image analysis

Raw data for one cDNA microarray slide typically consists of two scanned 16-bit images. While scanning, a green and a red laser excite the Cy3 and Cy5 dyes, and the emitted light is registered in every location. From the two images acquired, at least four different measures for each spot are extracted. The foreground signals, \( R_{fg} \) and \( G_{fg} \), are often measured as the mean or the median of the intensities of the pixels that are identified to belong to the spot. There are several ways to define what the background of a spot is and different definitions often result in different
background estimates, $R_{bg}$ and $G_{bg}$. All background identification methods strive to get a representative estimate of the background noise in the area of the spot, noise that is also expected to be added to the foreground signals. With a good estimate of the foreground and the background signals it is reasonable to believe that the background subtracted signals, $R = R_{fg} - R_{bg}$ and $G = G_{fg} - G_{bg}$, reflect the gene-expression levels. It is important to remember that the spots are not always easily identifiable regions in the images, but can often be weak, or smeared out as comets (as an effect from the wash-off of the poly-L-lysine). It also happens that two or more spots are so large that they overlap or that the microarray slides are contaminated with dust and scratches. For reasons like these it is important to have good methods for estimating the foreground and the background signals.

Yang et al. [20] compare the different foreground background identification and estimation methods that are used by the most commonly used image analysis software, such as GenePix [2], QuantArray [14], ScanAlyze [12], and Spot [1]. The authors show that the latter gives better foreground and background estimates than the others.

4 Normalization

Before any method for identifying differentially expressed genes can be applied, data has to be normalized. Normalization is a process of removing systematic variation in microarray experiments, which affects the measured gene-expression ratios. The sources of these artifacts can be many and they often reveal themselves in different ways. For instance, a too large imbalance in the PMT (photomultiplier tube) voltage setting for the red and the green laser in the scanner will give a bias in the fluorescence intensities toward one of channels. Another reason for having such a bias could be due to different physical properties of the two dyes (green Cy3 and red Cy5 dye), such as differences in half-life, size and weight, and heat and light sensitivity. Other systematic variations could be due to variations in the quality of the cDNA clones printed on the cDNA microarray slides. Varying printing conditions, such as temperature and humidity, during the printing process might further contribute to the variability in the measured gene-expression levels, or, as shown below, there may be problems with the print-tips.

The normalization process can be divided into two main steps. The first step identifies and normalizes for artifacts within each slide. The within-slide normal-
4. Normalization

Normalization is applied to each slide independently of the others. To be able to compare the measured gene-expression levels between replicated slides the spread of expression levels must be the same for all slides. This is taken care of by the second normalization step called between-slide normalization.

4.1 Transformation of variables

Since the quantity of interest is the relative gene-expression level and \( R \) and \( G \) typically range from 0 to 65535, we transform data by taking the logarithm of the ratio between the red and the green signals, \( M = \log_2(R/G) \). Another interesting quantity is the average log-intensity of the two signals, \( A = 1/2 \cdot \log_2(RG) \). The rational for the factor \( 1/2 \) is that \( A \) will have the same natural range as \( \log_2 R \) and \( \log_2 G \) and log-base two is used because data is originally binary. It is easy to show that this transform is reversible (and therefore the information contained in \( R \) and \( G \) is preserved). We have that

\[
M = \log_2(R/G), \quad A = 1/2 \cdot \log_2(RG) \quad \iff \quad R = (2^{2A+M})^{1/2}, \quad G = (2^{2A-M})^{1/2}.
\]

(1)

Many cDNA microarray experiments are set up in such a way that most of the genes are expected to be non-differentially expressed. In these cases an \( M \) versus \( A \) scatter plot will be more revealing than an \( R \) versus \( G \) scatter plot, because most of the data points in an \( R \) versus \( G \) plot will be distributed around the diagonal line \( R = G \) whereas the same data points in an \( M \) versus \( A \) plot will be distributed around the horizontal line \( M = 0 \). In Figure 1 an \( R \) versus \( G \) and an \( M \) versus \( A \) scatter plot for the same set of data are shown. The normalization methods discussed below are operating in the \((A, M)\) space. Also, if the error in the two channels is multiplicative, that is, \( R = c \cdot R_{\text{true}} \) and \( G = c \cdot G_{\text{true}} \), the transform to \( M \) will remove this noise.

It is important to have in mind that all the normalization methods used below are based on the assumption that there is only a small fraction of outliers \((1 - 2\%)\). We believe that this assumption is true for the ApoAI data set.
4.2 Within-slide normalization

4.2.1 Global lowess normalization

Taking the mean (or the median) of the log-ratios for each slide we should, since we assume that most genes are non-differentially expressed, get values close to zero. A box plot of the $M$ values for each of the eight ApoAI slides shows that this is not the case (cf. left plot in Figure 2). For slides 2, 3, and 5-8 the relative expression levels are biased toward the green channel (down-regulated genes) and for slides 1 and 4 there is a small bias toward the red channel (up-regulated genes). A first approach to normalize data would simply be to subtract the bias. However, the $M$ versus $A$ plot (for the third ApoAI slide), shows that the shift is not constant, but intensity-dependent (cf. right plot in Figure 2). The solid line in the plot is the lowess-fitted curve. The lowess function uses robust (not much affected by outliers) local linear regression to smooth scatter plots [9, 10]. All slides show this intensity-dependent pattern. This suggests that an intensity-dependent
4. Normalization

Figure 2: *Left:* Box-and-whisker plot showing the sample distribution of the log-ratios for each of the eight ApoAI slides. Since we assume most of the genes to be non-differentially expressed, ideally this plot should show a median around zero for all slides. *Right:* $M$ versus $A$ scatter plot for the third ApoAI slide with a lowess-fitted curve revealing that the log-ratios are intensity dependent.

Normalization [8] is needed:

$$M_{\text{norm}} = M - c(A)$$  \hspace{1cm} (2)

where $c(A)$ is the lowess fit to the $M$ versus $A$ plot.

4.2.2 Print-tip normalization

The printing step in the cDNA microarray process can also introduce systematic differences. In the ApoAI data set it is believed that there is an artifact present due to variation in the quality of the print tips [11, 21]. This could be due to different lengths of the print-tips making some of them leaving more cDNA on the slides than others, which in turn, due to dilution, could result in different concentration of cDNA in the wells on clone plates. Another reason could be deformation of some of the print-tip slits after hours of printing. For the ApoAI microarrays a print head of four-by-four print-tips were used ($J = 16$). Each print-tip printed a group of 399 spots, called a *print-tip group* (or grid). The print-tip groups are laid out on the slide from left-to-right and from top-to-bottom counting from the
upper left corner to the lower right corner (cf. legend in Figure 4). Within each print-tip the spots are printed from left-to-right and from top-to-bottom. In other words, the sixteen spots in the upper left corner in the sixteen print-tip groups were printed at the same time, then the spots right next to them were printed and so forth. Even if the slides were printed with a well-organized clone library, this mapping of spots assures that any interesting genes are located “all over” the slide. For this reason we can assume that the “true” log-ratios have the same zero-mean distribution across the print-tip groups. A box plot of the log-ratios for the sixteen groups (cf. Figure 3) shows that there are systematic differences between the groups and especially that print-tip groups 13-16 are different from the others. A plot of the spatial distribution of the top 5% absolute log-ratios (cf. Figure 3) confirms this. This suggests that some kind of normalization within each print-tip group is needed. The (global) lowess normalization method above

![Figure 3: Before normalization. Left: Box-and-whisker plot of the data points in each of the sixteen print-tip groups. The lower four groups, 13-16, shows a different behavior than the rest. A perfect slide would have the same average and variance across the print-tip groups. Right: The spatial position of the top 5% absolute log-ratios (|M|) on the slide. Each small rectangle represents the log-ratio of a spot and the large four-by-four rectangles represent the print-tip groups. It is clear that there is, especially in the lower four groups, some kind of artifact.](image)

can be applied to each print-tip group individually. The sixteen lowess curves in Figure 4 confirm that the groups 13-16 are different from the others and that the
differences are intensity dependent. The print-tip normalization [21] is

\[ M_{j,\text{norm}} = M_j - c_j(A_j); \quad j = 1, 2, \ldots, J, \] 

where \( J \) is the number of print-tip groups and \( c_j(A_j) \) is the lowess fit of the log-ratios \( M_j \) and the log-intensities \( A_j \) of the spots belonging to print-tip group \( j \). When doing a within-print-tip normalization a slide-intensity-dependent normalization as discussed in the previous section is unnecessary.

Figure 4: The \( M \) versus \( A \) plot with sixteen lowess-estimated curves, one for each of print-tip groups. The legend in the lower right corner shows how the print-tip groups are numbered, starting with one in the upper left group and counting left-to-right to sixteen in the lower right group. The signals from the four lowest print-tip groups, 13-16, have different properties than the others.
4.2.3 Scaled print-tip normalization

Doing a box plot and a spatial plot (not shown) similar to the ones in Figure 3 one gets that the print-tip normalization method not only removes the bias and the intensity dependency, but it also removes some of the spatial effects and makes the differences in variance between the groups smaller. However, there may still be differences in variance, which can be normalized by scaling the log-ratios within each print-tip group. The scale method used in [21] assumes that the log-ratios in each group, $j$, follow a distribution with zero mean and a variance $\sigma_j^2$. Not showing the details, it is straightforward to scale the print-tip groups so they get the same variance $\sigma^2$. To assure that the scaling is not affected by outliers the robust statistic MAD (median absolute deviation) is used to estimate the individual variances. The print-tip normalization method followed by a scaling step is called the scaled print-tip normalization method. The result of scaled print-tip normalization is shown in Figure 5.

Figure 5: After within-slide normalization. Left: Box-and-whisker plot of normalized data for each of the sixteen print-tip groups. The scaled print-tip normalization method first adjusts data in each print-tip group so that their lowess lines become zero for all intensities. Second, it scales data within each group so that their variances become equal. Right: The spatial distribution on the slide of the top 5% absolute log-ratios, $|M|$, from ApoAI slide 8.
5. Identifying differentially expressed genes

4.3 Between-slide normalization

Scaled print-tip normalization applied to all slides separately removes the intensity dependency of the log-ratios within the print-tip groups and this also removes the intensity dependency and the bias within the slides. However, the spread of the log-ratios might still vary from slide to slide and the same scaling technique that was applied to the print-tip groups may be used to scale variances between slides. Between-slide normalization makes it possible to compare the gene-expression levels measured by the different slides. The box plot in Figure 6 shows that scaled print-tip normalization followed by between-slide normalization makes the slides comparable (cf. Figure 2). The “average” slide is shown as an $M$ versus $A$ scatter plot in Figure 6. The bars over the axis labels denote the mean of $M$ and the mean of $A$, respectively.

![Figure 6: Left: Box-and-whisker plot showing the sample distribution of the log-ratios of the eight ApoAI slides after scaled print-tip normalization and between-slide normalization. Right: The “average” slide where the mean of the log-ratios and the mean of the intensities have been calculated for all spots across the slides.](image)

5 Identifying differentially expressed genes

5.1 Absolute log-ratios and the t-statistic

The most simple way to select genes being differentially expressed is to stratify on the absolute average log-ratios, e.g. by choosing genes that have $|M| > M_{\text{min}}$. 

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The main argument against using the average $M$ value to decide if a gene is differentially expressed or not, is that a large mean might be driven by an outlier, something which is not rare for cDNA microarray measurements. Thus, a better statistic to be used is the traditional $t$-statistic:

$$t_n = \frac{\bar{M}_n}{SE_n}; \quad n = 1,2,\ldots,N$$

where $\bar{M}_n$ is the (sample) mean of the log-ratios for gene $n$; $(M_{n,1}, \ldots, M_{n,K})$. $SE_n = s_n/K^{1/2}$ is the standard error of the mean, where $s_n$ is the (sample) standard deviation of the log-ratios for gene $n$. Since we do not care if the genes are up or down regulated, we look at the absolute $t$ value, $|t|$. For two genes with the same mean of log-ratios, the gene with the smaller standard error will get the larger $|t|$ value. The rational for this is that if repeated measurements of a gene-expression ratio give very similar results, i.e. the standard error is small, we should trust these measurements more than in the case where they differ a lot.

There are problems with the $t$-statistics. A gene with a small absolute mean might get a large $|t|$ value only because of a small SE, which might not be the true reflection of the gene. Remember that we are not only comparing $t$ values from two genes but from several thousands of genes. For this reason, by chance there will be genes that have larger $|t|$ values because they have small SE, not because they are differentially expressed. An ad hoc way to solve this problem is to discount genes with a small absolute mean whose standard errors are small. A small SE could be defined as SEs that are among the bottom 1%. Tusher et al. [19] presented another solution to the problem where they add a constant to each SE. Their statistic is referred to as the $S$-statistic.

### 5.2 The B-statistic

A better and more correct method for dealing with the above problem and for identifying differentially expressed genes is an empirical Bayes method developed by Lönnstedt et al. [16]. The authors introduce a new statistic called the $B$-statistic. The $B$ value calculated for gene $n$ is a log-posterior odds ratio for the gene to be differentially expressed, against to be non-differentially expressed, given the observed log-ratios for all genes and all experiments:

$$B_n = \log \frac{P(\text{gene } n \text{ is differentially expressed}|\{M_{n,k}\})}{P(\text{gene } n \text{ is non-differentially expressed}|\{M_{n,k}\})}$$
5. Identifying differentially expressed genes

where \( n = 1, 2, \ldots, N \). The probabilistic model behind this B-statistic assumes that the \( K \) replicated log-ratio measurements for gene \( n \), \( M_{n,k}; k = 1, 2, \ldots, K \), are normally distributed with mean \( \mu_n \) and variance \( \sigma^2_n \). However, the mean and the variance are not treated as fixed parameters, but as random variables themselves, and therefore we look at the distribution of \( M_{n,k} \) as a conditional distribution \( M_{n,k}|\mu_n, \sigma^2_n \sim N(\mu_n, \sigma^2_n) \). The assumption is still that most genes are non-differentially expressed, that is, they have \( \mu_n = 0 \), but a small proportion \( p \) of genes have \( \mu_n \neq 0 \). The expression above can now be written \( B_n = \log \left[ P(\mu_n \neq 0|\{M_{n,k}\})/P(\mu_n = 0|\{M_{n,k}\}) \right] \). Assuming special conjugate-prior distributions for the mean and the variances (for details see [16]), the authors get the full expression for \( B_n \) to be:

\[
B_n = \log \left( \frac{p}{1-p} \cdot \frac{1}{(1 + Nc)^{-1/2}} \cdot \left\{ \frac{a + s^2_n + \bar{M}^2_n}{a + s^2_n + M^2_n/(1 + Nc)} \right\}^{\nu + N/2} \right)
\]

where \( \bar{M}_n \) is the sample mean and \( s^2_n \) is sample variance of the \( K \) values for gene \( n \) with \( N \) as the number of genes. Leaving out the details, \( a, \nu \) and \( c \) are hyperparameters that are estimated from the log-ratios from all genes and all slides. Finally, \( p \) is the proportion of differentially expressed genes and \( p \) is normally the only parameter to be tuned. The B-statistic works in a similar fashion as the S-statistic, but the former gives a slightly smaller number of false positives and false negatives [16].

5.3 Number of replicates needed

Before presenting the results, a relevant and important question to be asked is how many replicates are that needed to get trustful results. We did all data analysis discussed above for the cases where \( n = 2, 4, 6, 8 \) slides were used. Since the number of slides does not affect within-slide normalization we did not have to repeat that step. However, we had to do the between-slide normalization for each \( n \). The top 40 genes when two, four, six and eight slides were used are listed in Table 1. The genes are ranked according to their \( B \) values. For \( n = 2 \) there were no genes with a positive log-posterior odds ratio among the top 40 genes. For \( n = 4 \) there were 12 with positive \( B \) values, and for \( n = 6 \) and \( n = 8 \) all top 40 genes had positive \( B \) values. More interestingly, when data from two slides are used, we can only identify 10 out of the top 40 genes found when all slides were used. When four slides were used, 17 out of 40 were found, and when
Table 1: Genes found as a function of the number of replicates. The table is listing the top 40 (0.6%) ApoAI differentiated genes found when the number of slides used in the analysis is $n = 2, 4, 6, 8$, respectively. In each column, the genes are ranked by their B-statistic. The number listed are the genes’ spot numbers as they occur on the microarray slides. The genes marked with a star (*) are the eight genes listed in [11]. The color and style coding is only used to help locate the same gene across the columns. All genes found when $n = 8$ slides were used are **bold-faced**. To simplify the identification of these genes in the $n = 2$, $n = 4$ and $n = 6$ columns, they are also color coded and some of them are in italic. Genes not found in the $n = 8$ column are non-bold and colored in silver.

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6. Results

6.1 Top forty differentially expressed genes in the ApoAI data set

Using the B-statistic on the ApoAI data set we identify a set of genes that we believe to be differentially expressed. The genes are listed Table 2, but can also be downloaded in a computer-readable format from [3]. Almost all the identified genes are down regulated, which makes sense since we are looking at a knock-out model. A more detailed discussion of the biological relevance of these genes are left to the SAGE KE discussion forum. The B values were calculated assuming 1% of the genes to be differentially expressed ($p = 0.01$). If we instead assume 0.5% of the genes to be differentially expressed ($p = 0.005$) we get that the 40 genes with largest log-posterior odds ratio are still ranked the same as in the $p = 0.01$ case. A $B$ versus average $M$ plot (cf. left plot in Figure 7) of the ApoAI data set shows that selecting genes based solely on the average log-ratios would not give the same set of genes as if the log-posterior odds ratio would be used. In the plot, the 40 (0.6%) genes with highest ranked $B$ values are highlighted red. To the right in Figure 7, the same genes are highlighted, but now in an average $M$ versus average $A$ plot.

6.2 Comparison between GenePix and Spot

In addition to using the Spot software for extracting the signals in the cDNA microarray images, we used the GenePix software. We analyzed data from GenePix in exactly the same way as we did with data from Spot and we compared the genes identified. The genes with the top 40 largest $B$ values found by GenePix and the genes with the top 40 largest $B$ values found by Spot are listed in Table 3. Both Spot and GenePix ranked the gene of spot 1496 to be the most differentially expressed gene. Spot ranked 2149 second whereas GenePix ranked spot 4941 second and so on. In GenePix’s top 40 list there are 15 genes that are found in Spot’s top 40 list, or equivalent in GenePix’s top 40 list, there are 25 genes that are not found in Spot’s top 40 list. Descriptions of these 25 genes are listed in the last column in Table 3.

Finally, looking at the $B$ values, Spot seems to produce larger $B$ values than GenePix. This can be interpreted as suggesting that the quality of data (signal to noise) from Spot is slightly higher than that from GenePix. However, a preferred comparison would be one based on their respective abilities to correctly identify
Figure 7: Left: The log-posterior odds ratio versus the average log-ratios. The B-statistic is the log-posterior odds ratio for each gene, which here is the same as each spot, to be differentially expressed. It is clear from this plot, that identification of differentially expressed genes based only on the average log-ratios would not give the same genes as if it was based on the log-posterior odds ratio. Small $|M|$ values tend to correspond to small $B$ values, but the reverse is not true. The genes with the top 0.6% largest B-statistics are highlighted red. We used the parameter value $p = 0.01$, i.e. we expect about 1% of the genes to be differentially expressed. Right: A scatter plot of the average log-ratios versus the average intensities with the same identified genes highlighted.

Genuinely differentially expressed genes, and on their false positive rates. Such a comparison is not possible at this point.
6. Results

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Table 2: Differentially expressed genes. The columns are 1) the rank (from B-values), 2) the spot number, 3) the log-ratio, 4) the intensity, 5) the log-posterior odds ratio, 6) the Cy5 intensity, 7) the Cy3 intensity, 9) the gene id, and 9) a short description of the gene/est.

The list of genes can be downloaded in a computer-readable format from [18].
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Table 3: Genes ranked highly by Spot versus genes ranked highly by GenePix. In column 2 are the spot number for the genes found using Spot for the image analysis. In column 6 are the spot number for the genes found by using GenePix for the image analysis. Except for the image analysis the rest of the data analysis was identical. In the last column are the descriptions of the genes that were highly ranked by GenePix but not by Spot listed. For the rest of the description, both for the GenePix and the Spot genes, see Table 2. The color and style coding is done in the same fashion as in Table 1.
References


Identification and normalization of plate effects in cDNA microarray data

Henrik Bengtsson

Mathematical Statistics at the Centre for Mathematical Sciences, Lund University, Box 118, SE-221 00 Lund, Sweden. Email: hb@maths.lth.se.

ABSTRACT

By introducing a novel way of visualizing cDNA microarray data we have identified a new type of systematic variation, which we refer to as plate effects. We believe that plate effects are due to non-biological differences in the DNA clones spotted onto the microarray slides. By comparing the variability of replicated spots (both within and between slides) after performing 42 different normalization strategies, we claim that the plate effects are non-negligible and should be corrected for in addition to the already known intensity-dependent effects. For comparison between normalization strategies we introduce a robust genewise variability measure, which we call measure of reproducibility. In addition, for the data set studied, we find that not doing background correction improves the reproducibility significantly.

Keywords: cDNA microarrays; print-order plot; systematic variation; artifacts; plate effects; clone effects; normalization.

1 Introduction

The cDNA microarray technology is a relatively new technique for measuring (relative) gene-expression levels in two different cell lines simultaneously. The technique covers a huge spectrum of applications such as identification of oncogenes (genes up-regulated in cancer), time-series studies of cell lines under different environmental stress conditions, clustering of expression profiles to identify related genes etc. The cDNA microarray technique is also started to be used in clinic trials for diagnosis on a patient-to-patient basis. With the vast amount of
biological data produced an increasing need for new statistical and computational methods is seen.

Robotically, using a so called *arrayer*, up to 40000 unique DNA (deoxyribonucleic acid) spots are printed onto a microscope glass slide. RNA from the two samples are purified separately, amplified and then turned into cDNA by reverse transcription. During reverse transcription the two solutions of cDNA are labeled with two different dyes, commonly the fluorescent dyes Cy3 and Cy5. Equal amounts of the two labeled cDNA solutions are co-hybridized to the DNA spots on the glass slide. The large number of cDNA strands and the competitive nature of hybridization make it possible to approximate the relative gene expression of a certain gene with the relative amount of Cy3 and Cy5 in the corresponding spot. The amounts of Cy3 and Cy5 in each spot are estimated by scanning the hybridized microarray slide at wavelengths 632 nm and 535 nm, which excite the Cy3 and Cy5 fluorescent dyes, respectively. The foreground and background signals of the spots are extracted and used in downstream analysis.

In the above process, there are several sources of error that introduce artifacts in data. In addition to problems of such steps as selecting sample tissues and extracting RNA, transcribing it into cDNA and so on, there are several other quality related problem. Dust sticking to the slides produces false signal peaks inside some spots, non-linearity between the two channels may favor one of the cell lines, spatial variation across a slide or systematic variation between replicated slides could additionally result in false conclusions. The list of artifacts is long, where some are well understood whereas some are still to be explained or identified.

In this paper we discuss what we believe is an artifact due to systematic variation between the DNA clone libraries and/or systematic variation between spots printed during the several hours long printing process. We refer to these artifacts as *plate effects*. As explained later, it is not obvious how to distinguish, a priori, between plate effects and effects due to differences in the two fluorescent labels. To identify the most dominant sources of systematic variations, we use a technique where different sequences of normalization methods, referred to as *normalization strategies*, are applied to data. Introducing a measure of reproducibility, we then search for the sequence of normalization steps that gives the least variable signals across replicated slides.
2. Data

All methods discussed in this paper have been implemented using the R software [6] and the com.braju.sma package [1], which is an object-oriented extension to the sma package [3] and relies on the R.oo package part of the R.classes [2].

The data set analyzed is described in Section 2, followed by Section 3, which lists possible artifacts in data. Previously known systematic variations are discussed, but also a not commonly discussed effect, referred to as plate effects. Print-order plots are introduced. In Section 4 different methods to normalize for artifacts are discussed. Different methods for normalization of plate effects are given in detail. Section 5 defines 42 different normalization strategies to be compared as well as the Measure of Reproducibility (MOR) used to compare the different strategies. Results are given in Section 6. Section 7 concludes with a discussion of underlying reason for plate effects.

2 Data

Data for the eight cDNA microarray slides analyzed in this paper comes from the Matt Callow Lab at Lawrence Berkeley National Laboratories [4]. Their experimental setup was to compare gene expressions between apolipoprotein AI (apo AI) knockout mice and a reference of C57Bl/6 mice. Two main experiments were conducted. The first one compared eight different apo AI knockout mice with a reference pool of eight different C57Bl/6 mice on eight cDNA microarray slides. The second experiment compared the individual C57Bl/6 mice with the pool of them (same pool as above). In total 16 slides were hybridized. In this paper, we only present results based on the eight slides from the second experiment, although the same analysis on the eight apo AI slides gives similar results (indeed almost identical numbers). However, the first set of slides is believed to contain fewer differentially expressed genes compared to the other set. Hence, it is easier to argue that the underlying assumptions used in this analysis are true.

To further summaries the work by [4], part of the extracted RNA from the livers of the eight individual mice in each slide set was synthesized separately into cDNA together with Cy3-dUTP. A mix of the same extracted RNA was then used to produce the Cy5-dUTP labeled reference cDNA. A similar setup was used for the six unique apo AI knockout mice. For the production of the microarrays, approx-
imimately 5600 expressed sequence tags (ESTs) from the I.M.A.G.E. consortium
database [7] were selected. 257 of these ESTs represent genes with possible and
confirmed associations with lipid metabolism. The slides were spotted using a 4-
by-4 print-tip arrayer with a total of 7056 spots on each slide. After hybridization
the slides were scanned and the foreground and background signals were extracted
using the Spot software [10]. Of the 21 rows in each print-tip group the last two
were excluded resulting in a total of 399 spots per print-tip group or in total 6384
spots per slide. Among these spots, there were in total 5357 different genes and
840 blank spots. Among the 5357 genes, six of them were spotted trice, 175 of
them duplicated and the rest were spotted only once per slide. In Figure 1, the

log gene-expression ratios on slide 6 are shown. To the left, the logarithm of the
gene-expression ratios, \( M = \log_2(R/G) \), are plotted against the log-intensities,
\( A = 1/2 \cdot \log_2(R \cdot G) \), where in this case \( R \) and \( G \) are the non-background
subtracted signals for the Cy5 (red) and Cy3 (green) channels, respectively. To
the right, a spatial plot depicts the relative gene expressions as positioned on the
slide.
3 Artifacts in data

As seen in the left plot in Figure 1, the log-ratios depend strongly on the log-intensities. A low intensity spot tends to be redder (up regulated) than a high intensity spot, and this intensity dependency is non-linear. All slides show the same systematic effects. Due to the experimental setup, most genes are expected to be non-differentially expressed. For this reason, the observed non-linearity is an artifact and should be corrected for. There is a variety of methods available to correct for this effect, but, throughout this paper, we will rely on the intensity-dependent normalization methods suggested by [10]. Furthermore, investigating the spatial distribution of log-ratios, cf. the right plot in Figure 1, we find a repeated pattern of horizontal stripes for all slides in the data set. If we combine this plot with the $M$ versus $A$ plot and keep in mind that there are intensity-dependent artifacts, we conclude that these stripes may be due to different intensities between the stripes. The red areas with apparent up-regulated genes are darker, and the down-regulated and the non-differentially expressed genes tend to be located in brighter areas. We will return to this in the next section. As shown in Section 4.1, an intensity-dependent normalization removes most of the spatial artifacts, but our final findings indicate that the non-linearity between the Cy3 and Cy5 channels is not the only contributor to such systematic variations. Even if it would, it still has to be explained why the intensity varies spatially in such a regular pattern across the slides.

3.1 Systematic variation between plates

At each, so called, dip, the print-tips of the arrayer spot a number of (different) DNA clones onto the glass slide. In our case 16 spots are spotted at each dip. Depending on how the arrayer is programmed it either move on to the rest of the slides and spot the same set of clones (with possible intermediate refills of print-tips), or it retrieves a new set of clones from the microtiter plates (we say that the arrayer does a source visit) and finish the first glass slide before moving onto the next. The most commonly used approach is to spot all glass slides in the arrayer before spotting the next set of clones. The positions of all the spots printed at each dip are determined by the layout of the print-tips (here 4-by-4 print-tips approximately 200 µm apart) and the way the arrayer was programmed. Typically, the next set of clones (here 16) is spotted to the right of the ones from the previous source visit. This is repeated until one row is filled when the arrayer continues with the next
row (starting over at the very left). Most modern arrayers allow user to spot the clones in any order and at any position, for instance column by column or randomly, but in most cases (also in this current setup) the arrayer moves from left to right row by row. With the knowledge of the exact printing of the slides one can create a plot where, say, the log-ratios of the 6384 spots are plotted in the order that the corresponding spots were printed onto the slide. A print-order plot is shown in Figure 2, with the log-ratio on the y-axis and the time of printing on the x-axis. From this plot it is clear that the log-ratios vary with the time point when the clones were spotted onto the slides. The systematic variation of the log-ratios over time is exactly that found to be repeated over the rows within each print-tip group, cf. right plot in Figure 1. The horizontal and vertical lines and the box plot to the right will be explained below. At a first glance it looks like there are

![Plot](image)

Figure 2: Left: The log-ratios of the 6384 spots on slide 6 in the order of when they were printed onto the slide. At each print dip, 16 spots were spotted simultaneously. The vertical lines mark the different clusters of plates. The horizontal lines are the biases of each plate group, which are colored according to which plate cluster (source) they belong to. Right: Box plot showing the median and the variability for each plate using the same ordering and coloring as in the left plot. Plate 1 is at the top and plate 18 at the bottom.

four to six printing periods (separated by vertical lines for clarity) with different properties. It turns out that the clones can be grouped into these groups based on what lab produced them. The slides used here were spotted using 18 different 384-well microtiter plates. Each of the vertical lines coincides exactly with a
plate and lab switch. In Figure 2 the plate numbers are displayed along the x-axis. Thus, the graph may also be referred to as a plate-order plot. In order not to focus too much on different lab procedures, we refer to these different groups of plates as plate clusters. Plate 1 contains mostly of water (empty spots), plates 2-4 come from the I.M.A.G.E. clone library, plates 5-8 contain clones selected by the King-ley lab at Stanford University, plates 9-12 contain brain-tissue clones obtained by the Cheng Lab at Lawrence Berkeley National Laboratories, plates 13-17 were purchased from Research Genetics, and plate 18 comes from the Vulpe Lab. For plate 1, only 96 clones were spotted, and for plate 18, 144 clones were spotted. For all other plates all 384 clones were spotted. Furthermore, 95 out of the 96 spots (99%) from plate 1 are blanks, 336 out of 384 spots (88%) on plate 12, 326 out of 384 (85%) on plate 17, and 64 out of the 144 spots (44%) from plate 18 are blanks. In total there are 840 empty spots. In Figure 2 the plate clusters have been clarified by vertical lines. The horizontal lines, which are colored according to which plate cluster they belong to, are the median log-ratios for each plate. The box plot to the right depicts the distribution of the log-ratios for each plate. Plate 1 is at the top and plate 18 at the bottom. The same color scheme is used throughout the paper.

As suggested above, the systematic variations in data, especially the spatial effects seen in Figure 1 can be explained by a combination of systematic variation in the spot intensities across the slides and an intensity-dependent log-ratio effect. The print-order plot of the log-intensities in Figure 3 together with the print-order plot of the log-ratios in Figure 2 confirm this. Although differences in signal strengths (intensities) for different plates explain most of the spatial effects observed, the results show that there are additional plate effects of different origin.

### 3.2 Possible sources for different artifacts

The plate effects originate from the clones and the plates, that is, from a step before spotting the slides. In this sense, all slides should show approximately the same plate effects. The intensity-dependent effects, however, results from a non-linearity between the Cy3 and the Cy5 channels. There are at least two different possible sources for this non-linearity. First, the incorporation rate vary with the fluorescent dye used. For instance, the incorporation rate of Cy5-dUTP is about 60% (in moles) of that of Cy3-dUTP [5]. The second source of bias is introduced when the slides are scanned. Due to quenching, the Cy3 and the Cy5 dyes
Figure 3: The log-intensities on slide 6. The plates with a lot of blanks (1,12,17 & 18) do have low intensities. Furthermore, the clones from brain tissue on plate 9-12 are, as expected, not very responsive to the test and control samples, which are from liver RNA. Left: The log-intensities in print order. Right: Box plot displaying the median and the variability of the log-intensities for each plate using the same ordering and coloring as in the left Figure.

are not linearly responsive over the full intensity range [8]. Since quenching is added after plate and labeling effects, it should be corrected for before the latter. However, because quenching and labeling effects show similar symptoms, it is hard to distinguish between them. Even if it is possible to identify the exact amount of quenching, a remaining problem is to separate the plate effect from the labeling effect. This is especially hard since these two effects are most likely combined in the hybridization step in an unknown way. We will attack the problem by assuming that the amount of quenching is not significant enough to be considered. Since we cannot distinguish between plate and labeling effects, we will test different sequences of normalization methods, referred to as normalization strategies, and use a measure of reproducibility (Section 5.1) to find the best one. This approach will guide us to which artifact is the most dominant one.
4 Normalizing data

Plate effects can be taken into account in many different ways when data is normalized. One may either search for a physical model to explain the effects, or use a blindfolded black-box model to normalize the observed signals. Here we use the latter approach, because we do not know in what step(s) or combination of steps of the cDNA microarray process plate effects are introduced, but also because it is a more objective approach for arguing for existence of plate-dependent effects. Next, we will present a few alternatives to normalize plate effects, and in Section 5.1 we define a measure of reproducibility in order to compare them.

4.1 Intensity-dependent normalization

Figure 4 shows the log-ratios after scaled print-tip intensity normalization [10]. Spatial artifacts are less significant after the normalization, but the variation between the plates are still there. A possible strategy is to remove the plate biases after the intensity normalization, cf. Section 4.4. For details on intensity-dependent normalization methods, see [10].

Figure 4: Gene expressions (for slide 6) after scaled print-tip intensity normalization. Left: Print-order plot of the log-ratios with a horizontal box plot showing the distribution of the log-ratios within each plate group. Right: The spatial location of the log-ratios. Normalizing for intensity-dependent artifacts removes much of the spatial and some of the plate effects.
4.2 Platewise median normalization

An alternative way to normalize data is found if we assume that the average log-ratio on each plate or plate cluster is zero. This is similar to the assumption used for normalization where one assumes zero shift, either for all spots or for each print-tip group. These two assumptions look similar, but there is an intrinsic difference between them. The clones on the plates might be selected in such a way that some plates may contain many clones that are differentially expressed whereas such an \textit{a priori} selection of clones is less likely to affect the print-tip groups, because they contain clones from all plates and are therefore more random. The assumption of zero shift for each plate is violated if the expression levels of the clones on a plate differ significantly between test and control samples. For instance, this may be the case with the apo AI versus control pool data set. We do not believe this is true for the self-to-self mouse experiment investigated here, because it is based on eight control mice versus the pool of them. Hence, we assume that all plates should have an average log-ratio of zero, which satisfies the assumption that the overall shift on the slide is zero. In Figure 2, the median log-ratio of each plate is depicted by horizontal lines. The result from normalizing data plate by plate assuming zero bias, that is, by shifting all the horizontal lines in Figure 2 to zero, is illustrated in the $M$ versus $A$ plot and the spatial plot in Figure 5. Comparing these with the corresponding plots for the non-normalized
4. Normalizing data

data (Figure 1) we find that intensity-dependent effects are partly removed by plate-bias normalization and that spatial effects are less significant. Remaining intensity-dependent biases are studied in Section 4.4.

4.3 Intensity-dependent normalization performed platewise

An alternative strategy is to apply intensity normalization for each microtiter plate individually. In Figure 6 the lowess-estimated intensity curve is shown for each plate. In this case we have to be careful, because the number of data points might not be large enough in order for robust smoothing estimates to be robust over the full intensity range. In such cases we have to look for larger groups, see Section 4.6, or find a clever way to combine different fitted curves, cf. [9]. This may be an issue for plate 1 and plate 18, but for simplicity we ignore such problems in this study. We do this because we do not believe it affects our findings in general. However, if we were searching for, say, differentially expressed genes, we would have to consider this problem more carefully. The platewise intensity normalized data for slide 6 is depicted in the print-order and spatial plots in Figure 7.

Figure 6: Intensity dependency for each plate (on slide 6). The top most curve (blue) of plate 12 is shifted upward because it has only one data point in the upper intensity range, $A = [12, 13]$, with a high value of the log-ratio.
4.4 Subsequent normalization strategies

When a platewise constant-shift normalization is applied some of the intensity effects are removed, but not completely as seen in Figure 5. Remaining effects can be removed by utilizing standard intensity normalization. However, care has to be taken since this will correct the plate-normalized signals too much and reintroduce “new” plate effects. A naive approach is to do another round of plate-dependent bias normalization. A similar strategy can be used to correct for remaining plate effects after an initial intensity-dependent normalization, or any other type of normalization. More importantly, the existence of both plate and intensity effects raises the question of which one of the two effects is the most dominant one, and which one should be corrected for first, or if both effects should be corrected for simultaneously. We leave the problem of simultaneous normalization to the future and focus on the simpler case, which is most likely the one that will be used in practice. Recall Section 3.2 for a short discussion on this problem.

4.5 About scale normalization

Since some plates have clones from different tissues and have been prepared by different labs and people, it is likely that their variabilities differ. For this reason, we do not assume that all plates have equal variability, and therefore we do not consider scale normalization of each plate group. For the same reason, we do not assume equal average intensity across plates. In addition, the noise and signal

Figure 7: Gene expressions (for slide 6) after platewise intensity-dependent normalization. Left: Print-order plot of the log-ratios with a horizontal box plot show the distribution of the log-ratios within each plate group. Right: The spatial location of the log-ratios.
levels in the two channels differ between plates, and therefore the variances of the log-ratios as well as log-intensities are expected to differ between plates. However, it is possible to assume equal variability across print-tip groups, because of “semi-random” positioning of clones. It is also reasonable to assume log-ratios to be zero on average.

4.6 Alternative ways to group data

Instead of focusing on plates, we may define other groups that are normalized separately. The main thing to keep in mind is, as for almost all normalization methods, that we must be confident that it is possible to assume that the average log-ratio of group is zero. For instance, in our case we could have grouped data into one group containing all the empty spots, and the rest of the clones grouped according to in which lab they were created. Another possibility is to estimate the average plate median shifts (or the plate intensity-dependent shifts) for all (standardized) slides, and use these platewise constants (or smooth curves) to normalize each slide individual. However, in this paper, we only consider platewise normalizations.

5 Comparison of different strategies

We compare seven classes of normalization strategies to each other and to the non-normalized case. In one group we normalize data for (slide, print-tip and scaled print-tip) intensity-dependent effects (labeled Sl(A), Pr(A) and sPr(A) in tables and figures). In a second group we normalize data for (constant) plate-dependent effects (labeled Pl), and in a third group we apply intensity-dependent normalization to each plate group (Pl(A)). In addition, we test different order of normalization sequences for each of them. In one group we first unshift the plate biases and then do standard intensity normalizations (labeled Pl-Sl(A), Pr(A) and sPr(A)). The reverse order is also considered (labeled Sl(A)-Pl, Pr(A)-Pl and sPr(A)-Pl). Furthermore, we do plate-intensity-plate normalization (Pl-Sl(A)-Pl, Pl-Pr(A)-Pl and Pl-sPr(A)-Pl). Finally, we combine intensity normalization over plates with standard intensity normalization methods, in that order (Pl(A)-Sl(A), Pl(A)-Pr(A) and Pl(A)-sPr(A)), and in the reverse order (Sl(A)-Pl(A), Pr(A)-Pl(A) and sPr(A)-Pl(A)). Each of the above is applied to both background (labeled bg) and non-background subtracted signals. In total, we compare $2 \cdot (1 + 3 + 1 + 1 + 3 + 3 + 3 + 3 + 3) = 42$ different strategies. For each of the strategies, we rescale...
the log-ratios so that all slides have the same MAD (within each normalization strategy, not between). More precisely,

\[
Z_l \leftarrow \text{MAD}_{k=1:K} M_{k,l}, \forall l
\]

\[\bar{Z} \leftarrow \left( \prod_{l=1}^{L} Z_l \right)^{1/L}\]

\[M_{k,l} \leftarrow \frac{\bar{Z}}{Z_l} \cdot M_{k,l}, \forall k, l\]

where \(k = 1, \ldots, K, l = 1, \ldots, L, K\) is the number of spots on each slide, and \(L\) is the number of slides (not to be confused with the number of genes and the number of replicates).

5.1 Measure of reproducibility

In order to identify an optimal normalization strategy, we measure the variability of the gene-by-gene residuals. The average variability and the variability of the variabilities can be seen as a measure of reproducibility. Ignoring the bias, the smaller the variabilities are, the more reproducible data is. Since we do not know the true values of the expression ratios, we cannot test if the different normalization methods introduce bias. For this reason, we cannot give rigorous evidence that normalized data is more consistent. However, all normalization methods described above (except the scale normalization) use additive operators, not multiplicative. Therefore, we believe the risk for overfitting data is low. Furthermore, if we search for the most differentially expressed genes among several, a small bias in the average expression ratio has little impact on the identified genes.

The measure of reproducibility is defined as follows. Let \(d_i\) be the variability of gene \(i\) calculated as the median absolute deviation of the gene residuals:

\[d_i = 1.4826 \cdot \text{median}_{j=1:J} |r_{i,j}|,\]

where 1.4826 is a constant to make MAD a consistent estimator of the standard deviation for normally distributed data. The residuals are defined as

\[r_{i,j} = M_{i,j} - \text{median}_{j=1:J} M_{i,j}\]
5. Comparison of different strategies

Figure 8: The log-ratios for all the 24 replicates of gene 1540 found on the eight slides after first performing intensity normalization for each print-tip group and then for each plate group (red discs), compared to not doing normalization at all (blue circles). The left scale is for normalized log-ratios and the right one for non-normalized ones. Note that the unit lengths of the two vertical axes are equal, but their origins have been aligned to simplify visual comparison.

where $M_{i,j}$ is the log-ratios of gene $i$ and replicate $j$. In Figure 8 the log-ratios for the 24 replicates of the triplicated gene 1540 are plotted for two different strategies. For this specific gene, print-tip followed by plate intensity normalization (red discs) makes the measurements (of this gene) more similar compared to the non-normalized approach (blue circles). The former has a genewise MAD of 0.115 (sample standard deviation is 0.0994) whereas the latter has a MAD of 0.190 (0.264). Obviously, this may only be true for this specific gene. Therefore, we look at the overall distribution of the gene-by-gene variabilities. The latter can be summarized by, say, the 95% quantile, or the mean, of the genewise variabilities $\{d_i\}_i$. The mean is not only a measure of the most common variability, but it is also sensitive to genes with high variabilities;

$$MOR. = \frac{1}{N} \sum_{i=1}^{N} d_i$$

where $N$ is the number of genes. Note that, the smaller MOR is the more reproducible data is. In general, rescaling log-ratios across all slides increases (worsen) the MOR. However, rescaling is commonly small and its effect on the MOR and,
hence, on the final findings is therefore of subordinary order.

6 Results

Our investigations show that it is optimal to normalize the data set in hand for intensity-dependent effects both within each print-tip group (Pr(A) or sPr(A)) and within each plate group (Pl(A)). The overall optimal strategy, for this data set, is to start with a scaled print-tip intensity normalization (sPr(A)) followed by a plate-wise intensity normalization (Pl(A)). The first one reduces the genewise variability by approximately 55%, and the second one reduces it another 5% relative to the variability of the non-normalized data (or approximately 10% compared to the proceeding one). See Table 1 and the box plot in Figure 9. This indicates that, in addition to intensity-dependent effects, there exist plate-dependent effects. Moreover, we find that it is always better to apply intensity-dependent normalization to each print-tip group individually (Pr(A) or sPr(A)) than to apply it to all spots at once (Sl(A)). We believe this is because normalization print-tip by print-tip corrects for spatial effects, which the latter is not capable of.

We find that not doing background correction give in more coherent measurements across slides and across replicates. This was true all normalization strategies tested. The only case when background subtraction gave less variability was when no normalization was performed (labeled 'none' in the table and in the box plot).

Almost identical results are obtained if the sample standard deviation is used instead of the sample median absolute deviation in the calculation of the variability of the gene residuals.

Finally, performing the same analysis on the apo AI knockout mice data set, which is expected to have a few differentially expressed genes, also gives that a print-tip followed by a plate intensity normalization performs better than a print-tip intensity normalization alone. For the same data set, the authors of [10] identify eight genes (or ESTs) to be the most differentially expressed genes. The top ten genes (the eight listed in their paper plus two others) were confirmed by real-time quantitative polymerase chain reaction (QRT-PCR) to be differentially expressed. All normalization strategies that we found to be the best ones do all rank these ten genes as the most differentially expressed genes. However, these genes are so
Figure 9: Comparison of sample distributions of the gene log-ratio variabilities after (2 × 21) different normalization strategies have been applied. Note that the tails of the sample distributions have been cut off at 0.75 and that they do extend (approximately equally far) beyond this level. Legend: none - No normalization, Pl - Constant plate bias removed, Pl(A) - Intensity-dependent plate bias removed, Pr(A) - Print-tip intensity-dependent bias removed, sPr(A) - Scaled print-tip intensity-dependent bias removed, Sl(A) - Slide intensity-dependent bias removed. Normalization strategies that start out with background subtracted signals are, in addition, labeled with 'bg'. For example, Pl-sPr(A) refers to a normalization sequence with a plate normalization followed by a scaled print-tip normalization.

extreme that almost any normalization strategy applied will identify them to be the most differentially expressed ones. Hence, we may only conclude that our best methods will not remove the signal for these top ten genes.
7 Discussion

The actual source of the plate effects is unknown, but one reason may be that there are different concentrations in the spots, which in turn may be due to different levels of evaporation for different plates.

The large difference between performing background correction and not is interesting. The subtraction of background from the foreground signals introduces noise, and the effect is enhanced by the log-ratio log-intensity transform. However, we do not believe it is of the same order of magnitude as observed for almost all of the normalization strategies tested. We believe that the background estimates, which are based on measurements from the surrounding regions of the spots, are not the same as the background noise that affects the internals of the spots. A physical explanation for this may be that the background noise found in the proximity of the spots do not stick to the spots themselves, simply because where the spots are the glass is already covered by cDNA and nothing but matching cDNA sequences can hybridize to the foreground areas. Furthermore, some (not all) of the background estimates from the Spot image analysis software show clear print-order effects too. This was surprising and has to be investigated further.

Finally, an argument against our measure of reproducibility is that it is only including the variance. The reason for this is that we do not know the true gene-expression levels for any genes. However, recently it has become more common to add control spots to cDNA microarray slides such as spike-ins, negative and positive controls etc. We are currently working on methods for comparing normalization strategies using both the bias and variance information of such control spots.

Acknowledgments

I am in big debt to professor Terry Speed at the Statistics Department, University of California, Berkeley, for supervising me while I was at the Statistics Department at UC Berkeley in 2000/2001. I am very thankful to Jean Yang and the rest of the Speed Group for taking the time to explain the cDNA microarray technique and for fruitful discussions. I would also like to thank Matt Callow at the
Lawrence Berkeley National Laboratory for providing data and answering all my questions about details in the experimental setup. Also, if it would not be for the discussion that I had with Karen Berger at the Monica Moore's Lab, Ernest Gallo Clinic & Research Center (Emeryville) in July 2002, I probably would not have come up with the print-order plot in the first place. I would also like to thank my supervisors Ola Hössjer and Jan Holst for feedback and proof reading. Finally, thanks to all the contributors to the R project, which provides me with an important foundation for applied research.

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Table 1: Comparison of different normalization strategies. Legend: none - No normalization, Pl - Constant plate bias removed, Pl(A) - Intensity-dependent plate bias removed, Pr(A) - Print-tip intensity-dependent bias removed, sPr(A) - Scaled print-tip intensity-dependent bias removed, Sl(A) - Slide intensity-dependent bias removed. Normalization strategies that start out with background subtracted signals are, in addition, labeled with 'bg'.
References


The R.oo package - Object-oriented programming with references using standard R code

Henrik Bengtsson

Mathematical Statistics at the Centre for Mathematical Sciences, Lund University, Box 118, SE-221 00 Lund, Sweden. Email: hb@maths.lth.se.

ABSTRACT

When designing and implementing object-oriented applications in R, problems concerning generic functions and reference variables often occur. It is currently not clear how to create generic functions in a robust way such that a new package will be compatible with existing or future packages. This becomes an important issue as more and more packages are made available. As a functional language, R does not provide methods for programming with references as all arguments to functions are copied by value. However, there are situations where it is useful or even necessary to pass arguments by references. In this paper we present the package R.oo, which overcomes these problems. In addition to a way for automatically creating generic functions in the background and a transparent way of using references, it also provides a foundation for designing and implementing object-oriented applications in a robust way. In this context, we also suggest a draft of a coding convention intended to bring additional structure to the source code. The package also extends the current exception handling mechanism in R such that exception objects can be caught based on their class.

Keywords: Object-oriented programming, generic functions, reference variables, coding conventions, exception handling, root class, S3, UseMethod.

1 Introduction

In R there are currently two ways for programming with classes, which are commonly referred to as the S3 (or S3/UseMethod) style [12] and the S4 style [6].
The S3 style has been supported by R from the very beginning and support for the S4 style was added with the release of the methods package [7], which became part of the core distribution as of R v1.4.0 and is loaded by default from R v1.7.0. The package presented in this paper is currently relying exclusively on the S3 programming style, meaning that all classes defined using the package become S3 classes. However, future versions are likely to migrate to the S4 programming style. Having said this, it should be clear that the intention of the R.oo package is not to replace S3 or S4, but to extend them with a layer such that object-oriented design and programming in R become easier and more robust.

For beginners, but also for experts, it can be quite tedious to implement an object-oriented design in a robust way using the standard S3 (or S4) schema. It is easy to forget to add a generic function or, worse, to overwrite a non-generic function by a generic function. As a developer one has to stay up to date with all generic functions and all functions implemented in the latest R distribution to be sure that core functionalities in R are not overridden or removed when one's package is loaded. To be certain that the package works anywhere one must stay up to date with all other packages that the end user might load in parallel with your package. This approach is not scalable as more and more packages are added to the CRAN.7

In R arguments are supplied to functions by a pass-by-value semantic, also known as call-by-value. Inside the function, the arguments behave like local variables and any change made to a value inside the function is not reflected in the original value. The rational for this is that a function by definition takes one or several input values, returns something else and it should never modify the input values. When passing large data objects to a function a true pass-by-value approach would be too expensive in terms of time and memory. To overcome this, an argument in R is in practice passed by reference as long as the variable is not modified by the function. As soon as the function changes the value of the argument, a local copy of it is created to obtain pas by value. Occasionally there are requests for adding a true pass-by-reference semantic so that functions can modify the original variable. From now on we refer to such functions as methods and reserve the word function for its original meaning. There are different ways to emulate a pass-by-reference

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7The current effort of adding namespaces [11] to the R language will remove some of the problems related to generic functions. A possible future support for multiple generic function will most likely solve the problem completely.
semantic by letting regular R variables represent reference variables or shortly references. With references it is possible to write more memory and time efficient code. Another advantage is that it is possible to write more user-friendly methods where fewer arguments need to be specified by the end user.

For reasons like these the R.oo package was developed. Its purpose is to relieve the developer from implementation details to make it possible to focus on the object-oriented design. The utility functions setConstructorS3() and setMethodS3() create constructor functions and class specific methods and at the same time make sure that required generic functions are created (or not) and throw exceptions if illegal method names are used, e.g. reserved words. Furthermore, the package provides a totally transparent way of using references by defining a new “data type”, namely the class Object. Any object of a class that inherits from this root class will be passed to functions as a reference. Using a single root class, which all classes inherit from, improves the overall design and structure of any package developed. Finally, the R.oo package also provides an extended exception handling mechanism where an exception can be thrown and then caught depending on its class.

The R.oo package was written using standard R code (“100% R”) and runs on all platforms. It was designed and implemented in an object-oriented style.

This document will not discuss what object-oriented programming and design are about. For an extensive case study on how to use the R.oo package see [2]. In Section 2, we describe how to use classes that are defined using the R.oo package and that inherits from the root class Object. In this section some additional object-oriented features that come with the R.oo package are also described. To further improve the structure of an object-oriented implementation, we suggest an R Coding Convention (RCC) in Section 3. The utility functions for defining constructors and methods, which are introduced in Section 4, assert that the RCC is followed. The root class named Object is described in detail in Section 5, and the usage of references, which is provided by the Object class, is discussed in Section 6. Section 7 is about exception handling and Section 8 describes some other utility functions that is part of the package. Other classes defined in the package are briefly mentioned in Section 9. Section 10 explains how to download and install the package. Conclusions are given in Section 11.
2 Using classes

Throughout this document we make use of a simple case study example to describe the major parts of the package. Let the class SavingsAccount represent a bank account, which in the simplest case can be described by its balance. To secure against illegal modifications of the balance we represent the balance with a private field named .balance (see Section 3). To obtain the balance of the account the function getBalance() is provided. Using setBalance() it is possible to modify the account balance directly, but it is not possible to set it to a negative balance. More commonly used are the methods for withdrawal and depositing, i.e. withdraw(amount) and deposit(amount), respectively. The withdrawal method will not accept withdrawals if the balance becomes negative. The class inherits from the root class Object (Section 5). When a SavingsAccount object is created, the balance will by default be set to zero. A Unified Modeling Language (UML) model of the SavingsAccount class is depicted in Figure 1.

<table>
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<td>.balance: double</td>
</tr>
<tr>
<td>getBalance(): double</td>
</tr>
<tr>
<td>setBalance(newBalance)</td>
</tr>
<tr>
<td>withdraw(amount)</td>
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<tr>
<td>deposit(amount)</td>
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</table>

Figure 1: UML representation of the SavingsAccount class, which extends the Object class. Private fields have a . (period) as a prefix. 'Object:' in the header means that the class extends the Object class.

2.1 Creating an object

The implementation of the class is described in Section 4, but for now assume that the usage of the constructor is SavingsAccount(balance=0) and that

account <- SavingsAccount(100)

creates a SavingsAccount object with initial balance 100. The object is referred to by the reference variable account.
2. Using classes

2.2 Accessing fields

The fields of an instance of class inheriting from root class Object, shortly an Object, can be accessed similar to how elements of a list are accessed. For example, the balance field of the account object can be retrieved by either account$.balance or account["balance"]. To set the balance of the account either account$.balance <- newBalance or account["balance"] <- newBalance will do.

Note that there is no way to prevent the access to private fields. However, if one follows the RCC rule [3] that private fields and only private fields should have a . (period) prefix, it should be clear which fields should be accessed from outside and which should only be accessed from inside the SavingsAccount class. Moreover, private fields named this way will, by default, not be listed by the functions getFields() and ll(), which are described further in Section 5, and neither by ls() [10].

2.3 Calling methods coupled with a class

Under the S3 schema a method coupled with a class is called in the same way as a regular function, but with (the reference to) the object as the first argument. When specifying the withdrawal and depositing methods above, we excluded the object argument for simplicity, e.g. withdraw(amount). However, when calling the method one has to include it, e.g. withdraw(account, amount). The method dispatching mechanism in S3/UseMethod then makes sure that the method of the correct class is called. For detailed information on how method dispatching is done in S3, see [10].

Similar to how fields are accessed, methods that are coupled with any Object derived class, can be accessed via the $ (or the [ ] operator, e.g. account$ withdraw(amount). In many other object-oriented languages such as Java and C++, but also the Omegahat's OOP project [8], this is the format used for methods calls. However, we do not recommend this style, except for static methods. Methods in R should be though of as belonging to generic functions [6] and not to a specific class per se.
2.4 Calling static methods

A static method of a class is invoked using only the class name and it does not require an instance of a class. All classes extending the Object class may define static methods. The most readable way to call a method of a class is via the $ operator, e.g. Object$load(file) and Exception$getLastException(), even though load(Object(), file) and getLastException(Exception()) are also possible.

2.5 Accessing virtual fields

For Object instances, there is a third way of calling methods. Methods with a name of format get\(<\text{Field}\>(\text{object})\) or set\(<\text{Field}\>(\text{object, value})\) can be accessed by what we denote as virtual fields, i.e. as object$\(<\text{field}\>$>. For instance the methods getBalance(account) and setBalance(account, newBalance) will be called whenever account$balance and account$balance <- value are evaluated, respectively.\(^8\)

There are at least three real advantages of using virtual fields. First, it is possible, as the name suggest, to make it look like a class has a certain field whereas it internally might use something else. For instance, a Circle class can have the two redundant fields named radius and diameter where one is a virtual field and the other is the actual field. Indeed, both might be declared virtual at the same time. We find that the use of virtual fields reduces the redundancy, which in turn reduces the risk for inconsistency. It also reduces the memory usage. Another advantage is that it is possible to restrict what values can be assigned to a field. For instance, we can prevent the user from setting a negative radius, e.g. circle$radius <- -20. Finally, virtual fields can prevent direct access to private fields or modification of constants, that is, they provide a mechanism for encapsulation (data hiding).

\(^8\)By default, virtual fields have higher priority than regular fields, meaning that if a virtual field exists it will be accessed first, but it is possible on a reference-to-reference or an object-to-object basis to change the order which fields, virtual fields, methods and static methods are accessed.
2.6 Accessing class fields

A class field, also known as a static field, is a field associated with the class itself, not with a particular instance of the class. A class field of a class is shared by all objects of that class. A common role of a class field is that of a global variable (except from the important difference that it is not a global variable), e.g. `Colors$RED.HUE`.

A class field is accessed as a regular field, except that the object is now the static class object, e.g. `SavingsAccount$count <- SavingsAccount$count + 1`. Any class extending the `Object` class can have static fields.

3 Coding conventions

An important part of object-oriented design and implementation is to follow a standard for describing the design and for implementing it. There are several standards for describing object-oriented design of software, but one that has become the major standard is the Unified Modelling Language (UML) [9]. For implementation standards, also referred to as coding conventions, some languages have a well defined specification to follow whereas others do not. Unfortunately, there is no explicit and official coding convention for R. A well defined coding convention is useful because it helps to make the code more structured and more readable, and it reduces the risk for mixing up field names with class names or reassign fields that are supposed to be constants etc. It is also fundamental in order to efficiently share source code between developers and over time. For this reason we are working on a R Coding Convention (RCC) draft [3]. Next we will present an excerpt of its naming conventions.

3.1 Naming conventions

Some of the naming convention rules of the RCC apply to object-oriented design and programming. One of the most important is how classes, fields and methods should be named. According to the RCC, names that represent classes must be nouns and written in mixed case starting with upper case, e.g. `SavingsAccount`. Both field and method names must be in mixed case starting with lower case, e.g. `balance` and `getBalance()`. Private fields should have a . (period) as a prefix to make it clear that it is a private field, e.g. `.balance`. Reserved keywords [10] and unsafe method names must also be avoided according to the RCC. The methods `setConstructorS3()` and `setMethodS3()`, described next, en-
force these naming rules and if not followed, an `RccViolationException` is thrown. Not all rules are enforced in order to be backward compatible with some basic R functions that (for obvious reasons) do not comply with the RCC. As a last resort, it is always possible to turn off the test against RCC by using the argument `enforceRCC=FALSE` when calling the above functions.

4 Defining new classes

Under the S3 schema there is no way to formally define a class and there is no way to enforce that an instance of a class has the correct format or contains the correct fields, or to assure that the inheritance structure is valid. One reason for this is that the class of the object and the inheritance structure of the class is solely specified by the class attribute of the individual objects. This attribute may be modified in any way, at any time, which makes the object-oriented implementation vulnerable to programming mistakes, but also to misuse. The S4 schema overcomes some of these lack-of-robustness drawbacks. The way the `Object` class is designed, the idea is that the fields (and hence the class) are defined inside the constructor function, and that the class attribute is never accessed by the programmer. This minimizes the risk for errors.

The two utility functions `setConstructorS3()` and `setMethodS3()`, introduced next, help the programmer to create constructors and methods without having to worry about generic functions. These functions can be used to define any method or classes, not only classes derived from `Object`.

4.1 Defining constructors

The `setConstructorS3()` sets the constructor function and automatically creates any necessary generic function (there are cases where this might be necessary). When defining a class descending from the `Object` class, the constructor function also plays the role of defining the class (its fields) and specifying which class to extend. For example, to create the SavingsAccount class we write:

```r
setConstructorS3("SavingsAccount", function(balance=0) {
  if (balance < 0) {
    throw("Trying to create an account with negative balance: ", balance)
  }
})
```

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4. Defining new classes

```r
extend(Object(), "SavingsAccount",
    .balance = balance
)
```

The declaration of the inheritance is done via the `extend()` method of the `Object` class, which will be called recursively throughout all the superclasses. The first argument to `extend()` should be the object returned by the constructor of the superclass. In the above example, the `SavingsAccount` inherits directly from the `Object` class, which is done by calling its constructor. The second argument to `extend()` should be the name of the class to be defined, e.g. `SavingsAccount`. According to the RCC, the name of the class should be the same as the name of the constructor function. Any other arguments to `extend()` are optional, but they must be named value arguments, e.g. `.balance=balance`, which then declare the fields of the class and their default values. Finally, all classes derived from `Object` must comply with the rule that it should be possible to create an instance of it by calling its constructor with no argument, e.g. `account <- SavingsAccount()`, cf. prototypes in S4.

### 4.2 Defining methods

The `setMethodS3()` method creates methods for S3 classes and at the same time encapsulates a lot of details that the programmer should not have to think about. One such thing is if a generic function should be created or not, and if so, how it should be created. For a detailed discussion on how generic functions are automatically created if missing, see Section 4.4. To create the `setBalance(newBalance)` method for the `SavingsAccount` class, the only thing needed is:

```r
setMethodS3("setBalance", "SavingsAccount",
    function(this, newBalance) {
      if (newBalance < 0) {
        throw("Trying to create an account with negative balance: ",
            balance)
      }
      this$.balance <- newBalance
    })
```

The reason for this is that static class objects are created by calling the constructor with no arguments.
The complete usage of `setMethodS3()` is:

```r
setMethodS3(name, class="default", definition, private=FALSE, protected=FALSE, static=FALSE, abstract=FALSE, trial=FALSE, deprecated=FALSE, envir=parent.frame(), createGeneric=TRUE, enforceRCC=TRUE)
```

where `name` is the name of the method, `class` is the name of the class, and `definition` is the definition, i.e. the function itself. If `class` == "default" (or "ANY"), a default function [10] is created, meaning that `setMethodsS3()` can be used whenever a function is defined. For all other arguments, see the help page of the function.

### 4.3 Details

The `setMethodS3()` method creates a standard S3 method and at the same time makes sure that a generic function for that method is available. For instance, the evaluation of

```r
setMethodS3("getBalance", "SavingsAccount", function(this) {
  this$.balance
})
```

creates the S3 method for the class and the S3 generic function, i.e.

```r
getBalance.SavingsAccount <- function(this) {
  this$.balance
}
getBalance <- function(...) UseMethod("getBalance")
```

It also makes sure that if there already exists a non-generic function called `getBalance()`, then it will be renamed to `getBalance.default()`. If the latter already exists, `setMethodS3()` cannot solve the conflict, and therefore an `Exception` will be thrown explaining this. A generic function is not created if a generic function (or an internal function that works as such) already exists. For instance,

```r
setMethodS3("as.character", "SavingsAccount", function(this) {
  paste(data.class(this), ": balance is ", this$.balance, ",",
  sep="")
})
```
5. The root class Object

creates only the S3 method and not the generic function. In addition to this, 
setMethodS3() will, by default, assert that the (most important) RCC naming rules 
are followed. If not, it throws an RccViolationException informing that an RCC 
rule was violated.

4.4 Safely creating generic functions

When writing a package it is important to make sure that the package does not 
overwrite preexisting functions. If a preexisting function exists that is not a generic 
function, in most cases, the conflict can be solved by redefining the function to 
turn into a default function. The test whether a function already exists or not is 
commonly done manually by the programmer. However, new functions might 
be added when a new version of R is released and more seriously, other packages 
might be loaded before or after our package is loaded and there is no way we can 
know which functions will be defined or not. A much safer approach is to check 
for conflicts and solve them when the package is loaded. Furthermore, it is impor-
tant to make sure that the generic function will work with all packages and not just 
the methods in our package. Moreover, complete object-oriented programming 
requires that methods can have the same name for different classes, but with an-
other set of arguments. By not specifying the arguments of the generic functions, 
but only the special ... argument, the generic function is “as generic as possible”, 
e.g. getArea <- function(...) UseMethod("getArea")\textsuperscript{10}. For a fur-
ther discussion on how to create generic functions safely, see [4]. These problems 
are all taken care of automatically by setMethodS3().

5 The root class Object

By enforcing that all classes are derived (directly or indirectly) from a common 
root class, we know that there exists a set of methods that are applicable to all such 
classes. This idea already exists to some extend in R, but by using a common root

\textsuperscript{10}If we do specify any arguments, we restrict the corresponding methods for all classes in all 
packages loaded at the same time to have the exactly the same set of arguments. Under the S4 
style, it is required and enforced that all methods for all classes have exactly the same arguments 
as the corresponding generic function. This is one of the reasons why we currently not use the S4 
style of programming with classes, but we hope to overcome this problem soon by making use of 
namespaces.
class it will be more explicit to the end user. The R.oo package defines the root class *Object*, which has the following methods coupled to it. See also Figure 2.

<table>
<thead>
<tr>
<th>Method</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>$(name): ANY</td>
<td>ANY is a data type that refers to any type or class.</td>
</tr>
<tr>
<td>$&lt;-(name, value)</td>
<td></td>
</tr>
<tr>
<td>[[(name): ANY</td>
<td></td>
</tr>
<tr>
<td>[&lt;-&lt;(name, value)</td>
<td></td>
</tr>
<tr>
<td>as.character(): character</td>
<td>Returns a string with short information about the object. This is the same string that by default is displayed by <code>print()</code>.</td>
</tr>
<tr>
<td>attach(private=FALSE, pos=2)</td>
<td></td>
</tr>
<tr>
<td>clone(): Object</td>
<td></td>
</tr>
<tr>
<td>detach()</td>
<td></td>
</tr>
<tr>
<td>equals(other): logical</td>
<td></td>
</tr>
<tr>
<td>extend(this, ...className, ...): Object</td>
<td></td>
</tr>
<tr>
<td>finalize()</td>
<td></td>
</tr>
<tr>
<td>getFields(private=FALSE): character[]</td>
<td></td>
</tr>
<tr>
<td>hashCode(): integer</td>
<td></td>
</tr>
<tr>
<td>ll(...): data.frame</td>
<td></td>
</tr>
<tr>
<td>static load(file): Object</td>
<td></td>
</tr>
<tr>
<td>objectSize(): integer</td>
<td></td>
</tr>
<tr>
<td>print()</td>
<td></td>
</tr>
<tr>
<td>save(file=NULL, ...)</td>
<td></td>
</tr>
</tbody>
</table>

Figure 2: UML representation of the root class *Object*, which all classes should be derived from directly or indirectly through other classes. ANY is not a defined data type, but refers to any data type or class.

The method `as.character()` returns a string with short information about the object. This is the same string that by default is displayed by `print()`.

The `print()` method prints information about the object. By default, the string returned by `as.character()` is printed. For convenience in R, *any* object of any data type or class whose name is typed at the command line followed by ENTER, the `print()` of that object will be called. Example:

```r
> 1+2 # gives the object '3', i.e. print(3)
[1] 3
> account # same as print(account)
SavingsAccount: balance is 100.
```

The method `getFields(private=FALSE)` returns the name of all fields in the object. By default, only names of non-private fields are returned.
The method `ll(...)` returns a data frame with detailed information about the fields of the Object. By default, only non-private fields are listed. For more information see section 8.

The `hashCode()` method returns an integer hash code for the Object.

The `objectSize()` method returns the (approximate) size of the Object.

The `equals(other)` method compares one Object with another. If they are equal, the method returns `TRUE`, otherwise `FALSE`. If argument `other` is `NULL`, then `FALSE` is always returned. The default implementation of `equals()` is comparing the `hashCode()` values of both objects.

The `clone()` method creates an identical copy of the Object\(^{11}\).

When an Object is deallocated from memory by the garbage collector the `finalize()` method is first called. Subclasses can override this method to make sure that any instances of those classes clean up after themselves. For instance, objects that allocate shared resources such as connections should make sure that these resources are closed and deallocated upon deletion.

The methods `attach(private=FALSE, pos=2)` and `detach()` attach and detach an Object to the search path, respectively. By default, only public fields (private=FALSE) of an Object are attached, and, by default, they are attached to the beginning of the search path (pos=2) immediately after the global environment. Any modification to such attached fields will not be reflected (saved) in the actual Object. The attach-detach mechanism is intended for read-only purposes.

The method `save(file=NULL, ...)` saves an Object to a file (or a connection), and the static method `load(file)` loads a previously saved Object and returns a reference to it.

The somewhat special method `extend(...className, ...)` extends an Object class into a subclass named according to the string `...className`\(^{12}\) and which contains

\(^{11}\)Doing `ref2 <- ref` will only create a new reference to the same instance.

\(^{12}\)The second argument to `extend()` has three dots as a prefix to make it possible to name fields.
all fields as given by the ... arguments. This method is not intended to be
overridden by any subclass.

Finally, as explained in the next section, the functionality for references is hidden
inside the Object class. Hence, all subclasses will support references automatically
and the programmer does not have to think about how reference variables should
be implemented. They are always provided and they always behave in the same
way.

6 Reference variables

All instances of the Object class (or one of its subclasses) are accessed via reference
variables or shortly references\textsuperscript{13}. In standard R, where reference variables are not
provided, each instance of a class is accessed by one single variable, the object
itself. With references, however, it is possible for several variables to access the
same object. Here is an example where a list contains several references to the
same Object:

\begin{verbatim}
person <- Person("Dalai Lama", 68)
l <- list(a=person, b=person, c=clone(person))
setAge(l$a, 67)
print(person)
[1] "Dalai Lama is 67 years old."
setAge(l$c, 69)
print(person)
[1] "Dalai Lama is 67 years old."
\end{verbatim}

If person would not be a reference, the two elements a and b would be another
two copies (clones) of the Person object and a modification of one of them
would not affect the other instance and neither the original variable person. It
is possible to create a copy of an Object by using clone() as illustrated in the
above code.

\textsuperscript{13}For those who are not familiar with references but with pointers, references can be thought of
as safe pointers to objects that cannot by mistake be made to point to the wrong part of the memory.
Object-oriented programming where objects are passed by references does not differ much from the
case when objects are passed by value. However, there are differences that are important to be aware
of.

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Furthermore, references make it possible to implement software that otherwise would not be possible, or would be very tedious, to implement. Using references, more details can be encapsulated and thereby the package will be more user-friendly. We believe that a well designed object-oriented method interface based on references can serve as a base for, but also be a good complement to, a graphical user interface (GUI). For more complete real-world examples, see [2] and [1].

6.1 Garbage collector

The use of references requires a memory management. Many languages, including R, provide a built-in garbage collector, which removes obsolete objects from the memory that are not referred to by anyone. Since objects inherited from the Object class are standard R objects they are recognized by the garbage collector. For example, an Object created inside a function and for which no reference is returned, will be deleted by the R garbage collector. In summary, objects do not have to be deleted explicitly, but for an Object to be deleted it is important that all references to it are removed, for instance by rm(), or set to NULL. It is a good custom to do this as soon as an Object is not needed.

6.2 Details

R does not support references, but references can be emulated using so called environments [10]. However, using environments explicitly will quickly fill the source code with several get(name, envir=ref), assign(name, value, envir=ref) and/or eval(..., envir=ref) statements. This makes the code hard to read and increases the risk for errors. By encapsulating all calls to get() and assign() in the operator methods $(), $[(), $<-(), $[-()] of the root class Object, all fields can be accessed as if they were elements in a list. There are other ways for emulating references and some are more and some are less memory and time efficient than others. The R.oo package use the first approach, where each object lives in its own environment. One reason is that the garbage collector recognizes environment variables.
7 Exception handling

In addition to methods for defining classes and support for references, the package provides an improved exception handling mechanism. The core functionalities for exception handling is done by the `Exception` class (see Figure 3). It provides methods to create and throw exceptions and together with its companion `trycatch()` complete exception handling is provided.

7.1 Creating and throwing exceptions

The easiest way to create and throw an exception is by calling `throw()`, e.g.

```R
throw("Division by zero.")
```

which is equivalent to calling

```R
throw(Exception("Division by zero."))
```

An object of any class that inherits from `Exception` contains information about the error and when it occurred. Any `Exception` object can be thrown using the `throw()` method and then optionally be caught by either `trycatch()` or `try()`. If an `Exception` is thrown, the last exception thrown can be obtained by the static method `getLastException()` of class `Exception`. The `as.character()` method for the `Object` class is overridden by the `Exception` class and the default print message of an `Exception` has the format:

```R
> throw("Division by zero.")
```

![Figure 3: UML representation of the Exception class, which extends the Object class.](image-url)
7.2 Catching exceptions depending on class

The `trycatch()` method can catch exceptions based on what class they belong to. Like the `try()` function, the first argument to `trycatch()` is the expression to be evaluated, which might throw an exception. Any further arguments must be named arguments where the name specifies the `Exception` class to be caught and the value the code to be evaluated if such an exception is thrown. An argument with name `ANY` will catch any kind of Exception (including `try-error` thrown by `stop()`). If an exception is caught, and no further exceptions are thrown, then `trycatch()` returns safely. The following code will generate and throw an exception, which will be caught by the `ANY` clause, preventing the R session from being interrupted.

```r
trycatch(
  x <- log(2)
  y <- log("a")
), ANY=
  x <- 0
  y <- 0
  print(Exception$getLastException())
)
```

```r
print("trycatch() did indeed catch the exception.")
```

Moreover, code defined by an argument named `finally` is guaranteed to be evaluated immediately before `trycatch()` returns. This is, for instance, useful if a connection needs to be closed regardless of whether an exception is thrown or not.

8 Utility functions

In addition to the already mentioned methods, the package also defines some useful utility functions, which are applicable to objects of any class or data type. The default method of `ll()` lists detailed information about objects (variables and functions) found in an environment. The returned data frame will by default contain information about the member (name of the variable or function), `data.class`, `dimension` and `object.size`, which are the values returned by the functions with the same names.
C

> 11()

<table>
<thead>
<tr>
<th>member</th>
<th>data.class</th>
<th>dimension</th>
<th>object.size</th>
</tr>
</thead>
<tbody>
<tr>
<td>analyze</td>
<td>function</td>
<td>NULL</td>
<td>248</td>
</tr>
<tr>
<td>ma</td>
<td>MAData</td>
<td>1</td>
<td>452</td>
</tr>
<tr>
<td>raw</td>
<td>RawData</td>
<td>1</td>
<td>452</td>
</tr>
<tr>
<td>gpr</td>
<td>GenePixData</td>
<td>1</td>
<td>460</td>
</tr>
<tr>
<td>y</td>
<td>numeric</td>
<td>100</td>
<td>828</td>
</tr>
</tbody>
</table>

For information about other utility functions provided by the package, see the help of the package.

9 Other classes

Other classes that are loaded with this package are Class, Package and Rdoc. The class Class provides an interface for querying classes about methods, fields etc. The class Package represents any kind of package, e.g. Package("base"). Given a Package object, it is possible to query it for its classes, its author, check for updates (see below for an example) etc. The Rdoc class provides a compiler for Rdoc documentation, which is an extension of the Rd language that minimizes the need for having to update the documentation when the source code is updated. For instance, the tag @synopsis generates a correct \usage (or a \synopsis) markup given the other information in the Rdoc code. The Rdoc documentation can be standalone files similar to Rd files (the simplest Rd file is a plain Rd file) or it can be part of the source files in form of comments. The Rdoc code is compiled into standard Rd files, which are then converted into help pages etc. by R CMD build. We intend to extend the Rdoc compiler to recognize S4 classes too.

10 Installation

Since the package was written in 100% R, no native code needs to be built and installation is straightforward. The R.oo package is part of a bundle of packages called R.classes [5]. To download and install the R.classes bundle, do

install.packages("R.classes", contriburl="http://www.maths.lth.se /help/R")

from within R. By default, the R library directory is the directory named library/ in the directory where R is installed. The bundle can be in-
stalled in a private directory by setting the environment variable \texttt{R_LIBS}, e.g., \texttt{setenv R_LIBS $HOME/R/}. By loading the package, e.g. \texttt{library(R.oo)}, the correctness of the installation can be verified. To install on a Macintosh that does not have OS X, or to manually install the bundle, see [5]. For future updates, load the package and do 

\texttt{update(R.oo)}

\section{Conclusions}

The R.oo package is open source, it is designed in an object-oriented style and implemented using plain and richly commented R code (100\% R). Moreover, it is designed and implemented such that any future migration from S3 to S4 will be as smooth as possible for the end user.

For over two years we have used the R.oo package and the \texttt{R.classes} bundle in a project developing a cDNA microarray analysis package (com.braju.sma [1]). We have found that, by using the R.oo package, we never have had problems with conflicts related to generic functions and we never have had to create a generic function explicitly. In cases when we by mistake tried to use a reserved word as method name, \texttt{setMethodS3()} immediately notified us. We have also found the Rdoc compiler to be a valuable tool for maintaining nearly 200 Rd files. Due to the huge memory load and the large amount of redundancy in microarray data, the use of reference variables has been a natural and successful choice. Moreover, since methods can change the state of objects if references are used, we have been able to decrease the number of arguments that has to be specified in the method calls and therefore we can provide a cleaner and more user-friendly method interface. For a further discussion on how the R.oo package has been used in the development of our microarray package, see [1]. To install the com.braju.sma package see [1].
References


[10] R Development Core Team. R Language Definition (v1.7.0, draft), April 2003.


Paper D

Methodological study of affine transformations of gene expression data with proposed normalization method

Henrik Bengtsson¹ and Ola Hössjer²

¹ Mathematical Statistics at the Centre for Mathematical Sciences, Lund University, Box 118, SE-221 00 Lund, Sweden. Email: hb@maths.lth.se.
² Mathematical Statistics at the Mathematics Department, Stockholm University.

ABSTRACT

A detailed methodological study of affine models for gene expression data is carried out. We focus on two-channel comparative studies, although the findings are not restricted to such. We find that an affine model is capable of explaining many of the commonly observed discrepancies and systematic effects in gene-expression levels and observed log-ratios. Focus is also on data obtained by the two-color spotted microarray technology, but most of the discussion applies equally well to other gene-expression techniques such as single-channel hybridization methods and quantitative real-time PCR. An affine model can also explain non-linear systematic effects commonly observed when log-ratios obtained by different gene-expression technologies are compared. A high-quality cDNA microarray data set is used to demonstrate the power of the affine model. In the light of the affine model, the strengths and the weaknesses of the most commonly used normalization methods are discussed. Based on the affine model, we propose a novel method to normalize one or multiple arrays simultaneously, where each array has been hybridized with one, two or more samples. A package with all necessary methods to read and normalize data have been written in the R language and is made available for free.

Keywords: microarrays; affine transformation; bias; logarithm; non-linear systematic effects; robust normalization; background correction; principal component analysis; iterative reweighted PCA.
1 Introduction

The general idea behind most gene-expression measurement technologies is to assess the expression levels of (all or a subset of) genes in one or several cell populations. The ensemble of gene-expression levels is often referred to as the gene-expression profile of the sample. As it is not possible to measure the expression level of a gene in practice, techniques have been developed to measure closely related properties such as the amount of mRNA (messenger RNA) or the amount of proteins produced. Although different techniques to measure protein levels have evolved and improved recently, they are still not as cheap and widely used as those to measure mRNA levels. The amount of mRNA is either measured directly or, as it is much less stable than DNA, indirectly by first translating it into cDNA (complementary DNA). One of the technologies measuring cDNA is Southern blot, which is based on electrophoresis in gels and measures the DNA levels of a small set of genes simultaneously. A succeeding technology, Northern blot, is similar, but measures the RNA levels instead. Another technology that measures mRNA levels indirectly via cDNA is QRT-PCR (quantitative real-time polymerase chain reaction). QRT-PCR is a low-throughput technique, but, because of its high accuracy (closer to the truth) and high precision (more similarity between replicated measurements), it is often taken as the gold standard. For this reason, QRT-PCR is often used to verify the correctness of other technologies such as the microarray technology.

In addition to the above methods, recently developed microarray techniques [30] provide a way of measuring the expression levels of many more genes, in the order of $10^3 - 10^5$ or more. This has made it possible to take a snapshot of the activity of a person’s complete genome in one single experiment. Microarrays have well defined and immobilized regions that each consists of clones or synthesized sequences of DNA specific to a unique gene. We refer to these (non-hybridized) regions or spots as probes [13]. A cocktail of cDNA created from the RNA extract, referred to as the target, from the cell population in study is then hybridized to the DNA on the microarray for a few hours after which excess cDNA is washed off. The result is that each region of the microarray contains a certain amount of hybridized DNA unique to the corresponding gene. By first labeling the cDNA strands in the sample cocktail with a radioactive or a fluorescent probe, the amount of hybridized DNA can be measured using radioactive sensitive film or a color-sensitive scanner, respectively.
1. Introduction

When measuring gene expressions for an individual gene we try to assess how active (measured on some scale) that gene is. Because it is hard to identify an absolute scale to measure on, often, but also for various other reasons, a reference is used to obtain a relative scale. As even genes from the same sample are not directly comparable to each other, each gene gets its own reference, which is typically the same gene from a reference sample. With this approach, we can obtain gene-expression ratios for every gene, which for instance can be used to test the hypothesis if a gene (in the test sample) is differentially expressed or not (compared to the gene in the reference sample). This is the core idea behind the two-channel microarray technology, in which the test and the reference cDNA cocktails are hybridized to the same array simultaneously and in a competitive way. The same idea has been adopted by single-channel hybridization technologies where the comparison is instead done numerically in the data analysis step.

Even if gene-by-gene references are used, the measurements are not perfect and are likely to contain systematic errors, which possibly vary from measurement to measurement, and the obtained gene-expression ratios may still be biased and not comparable to each other. What we ultimately would like to do is to measure all control and all reference samples under identical conditions. The aforementioned two-color microarray technology tries, in some sense, to do this by measuring the control/reference pairs for each gene in one hybridization (although it is not clear if the gain from co-hybridizing two samples with different labels is larger than hybridizing twice with identical label molecules and then scanning the samples separately).

In this paper we present an affine model, which can explain many of the systematic effects often observed when gene-expression levels from two (or more) samples are compared. The main contributors to such systematic effects are the biases in each channel, which give non-linear systematic effects in ratios etc. Here we will not give an error model, but instead only a deterministic model. The main reason for this is that we believe an error-free model makes us better understand the impact that channel biases have on the downstream analysis regardless of gene-expression technology used. This is especially of interest as these are often implicitly assumed to be small and of no effect, which we believe is a too strong assumption. Although we do focus on the microarray technology
and although some results have been published [28], we believe much more research is needed before we can understand and correctly model the various error sources introduced in the microarray process.

The outline of this report is as follows. In Section 2, a general model that incorporates all steps of any gene-expression technology is given. By dissecting the generic model and focusing more on the microarray technologies, an affine model is introduced. In Section 3, we study the properties of the widely adopted and accepted log-ratio log-intensity transform under the affine model. An investigation of how the affine transform introduces biases in the log-ratios is carried out. In Section 4, we revisit some of the normalization methods available in literature, to which background correction may also be counted. We conclude the section by suggesting a novel robust normalization method that can be applied to single-channel as well as multi-channel microarray data. The method will correct for both systematic effects and scale differences in the log-ratios. We end the report with a discussion on similarities with other normalization method, which gives us some future research directions.

2 Model

2.1 A general model

Consider an experiment that involves genes $i = 1, 2, \ldots, I$ from RNA extracts $c = 1, 2, \ldots, C$. For example, in oligonucleotide microarrays each slide measures the gene-expression levels of exactly one RNA extract whereas for a two-color microarrays each slide measures two RNA extracts, one in each channel. From now on, we refer to the RNA extracts or replicates of such as channels. Let $x_{c,i}$ be the true gene-expression level of gene $i$ in channel $c$ and let $y_{c,i}$ be the corresponding observed gene-expression level. The relation between the observed and the true expression levels can be written as

$$y_{c,i} = f_c(x_{c,i}) + \varepsilon_{c,i}$$  \hspace{1cm} (1)

where $f_c(\cdot)$ is a channel specific function, which we refer to as the measurement function, that includes all steps in the gene-expression acquisition process. Most generally, we have that $E[\varepsilon_{c,i}] = 0$ and $V[\varepsilon_{c,i}] = \sigma_{c,i}^2$, where the variance can take any form. Importantly, the properties of $\varepsilon_c$ are still not well known and
2. Model

varies depending on if the overall or a submeasurement function is considered. For instance, under certain circumstances it can be shown that the standard deviation of the noise is proportional to the signal, but in other cases the relationship might be more or less complicated. For this reason and because of the many interesting effects that the affine transformation (presented below) generates by itself, we have decided to conduct this study under the assumption of noise-free data.

To continue, since we want to do inference based on $x_{c,i}$, the inverse of $f_c(\cdot)$ has to be identified, something that is possible if it is strictly increasing. Violation of this constraint has been observed in, for instance, two-color microarray data. This can be due to too high concentrations of fluorophore molecules, which quenches the signal, sometimes so much that the signal decreases when the concentration increases [27, 24]. Extreme saturation in the scanner, which is commonly observed when the PMT gain is set too high, results in censored signals, which in turn prevents us to find a unique inverse of the measurement function. The above relationship can be specific for certain subsets of genes or spots such as print tip [35], microtiter plate or clone library [4], spatially dependent or even gene specific, but in the general discussion that follows we will avoid such details for simplicity.

2.2 Dissection of the overall measurement function

Gene-expression levels are not measurable directly and any technique used transforms the expression level into a measurably quantity, commonly a fluorescent intensity or a voltage level. Mathematically, each step in the microarray process can be seen as a function that takes a set of input objects and outputs another set of objects. The sequential nature of the process allows us to think of the measurement function $f_c(\cdot)$ as a composite function (function of functions): $f_c(\cdot) = (f_{c,K} \circ f_{c,K-1} \circ \cdots \circ f_{c,1})(\cdot)$, where $K$ is the number of steps in the process. For instance, and of course simplified, it could be that $f_{c,1}(\cdot)$ models the extraction of the RNA from the cell, $f_{c,2}(\cdot)$ models the reverse transcription of RNA into cDNA etc., and $f_{c,K}(\cdot)$ models the scanning and the signal quantification of the hybridized array. We realize that some of these submeasurement functions are shared by several channels and that others are channel specific or even gene specific. Moreover, there is also one or several joining subfunctions, e.g. the hybridization of labeled cDNA sequences to the
probes on the array. Thus, the overall measurement function $f_c(\cdot)$ consists of a complex and to a large extent unknown ensemble of subfunctions $\{f_{c,k}(\cdot)\}_{k=1}^K$.

A first order Taylor series expansion of an arbitrary measurement function $f_c(x_c)$ has the form

$$f_c(x_c) = a_c + b_c x_c + R_c(x_c), \forall c.$$  \hspace{1cm} (2)

Next, we will argue that $a_c$ cannot be assumed to be zero and $b_c$ cannot be assumed to be one. In addition, in order to focus on the effect of $a_c$ and $b_c$, we will ignore higher order terms ($R_c(x_c)$) assuming they are small. Thus, we do not take into account the possibility that saturation might occur. However, as we believe saturation can be avoided, we will not discuss function of higher order in this report, but instead we focus on measurement functions that are affine.

### 2.3 Existence of biases in each of the channels

From the above dissection of the measurement functions it is not hard to believe that some of the subfunctions introduce bias and therefore we ought to have an overall bias in $f_c(\cdot)$. For instance, the bias terms can be due to non-uniformity of the reverse transcription, the labeling [24] or the hybridization, due to dark noise in the PMT [19] or laser scatter light in the scanner, background noise, or non-uniformity of the scanned glass slide [16], threshold effects etc. In [3] it is shown how various background estimates based on different image analysis methods may introduce bias. Similarly, we have observed that different scanners, but also different image analysis methods, introduce a bias in the observed signals [6].

### 2.4 Existence of scale factors in each of the channels

It is not hard to imagine that there is also an overall scale factor (different from one) in each $f_c(\cdot)$. This can be due to differences in labeling or hybridization efficiency, laser power, PMT gain, wavelength dependent optical effects, bleaching of fluorophores, and so on. As in the case for the biases, the scale may vary between different subsets of genes or spots as well as with spatial position. Different scale factors have also been observed between clone libraries and spike-in controls [32].

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2.5 The affine measurement function

From the above discussion the simplest possible parametric measurement function reasonable to consider is an affine transformation of the RNA concentration $x_{c,i}$ as

$$f_c(x_{c,i}) = a_c + b_c x_{c,i}, \forall c, i,$$  \hspace{1cm} (3)

with unique inverse

$$x_{c,i} = f_c^{-1}(y_{c,i}) = \frac{y_{c,i} - a_c}{b_c}, \forall c, i,$$  \hspace{1cm} (4)

where $a_c$ is the overall bias and $b_c$ is the overall scale factor of channel $c$. Both the bias and scale parameters are commonly positive, but under certain circumstances, for instance, as demonstrated later, when two different measuring techniques are compared, the effective bias may be negative. Recall that especially $a_c$, but also $b_c$, may vary between different subset of genes etc., but for simplicity we focus on the case where all spots have the same bias. The left plot in Figure 2 shows, on a log-log scale, an example of an affine transformation of two-color microarray data where the offsets in the red and the green channels are 20 and 200, and the scale factors are 0.8 and 1.4, respectively. Modeling microarray data by an affine transform is not novel [28, 17, 21, 12], but the reasons for such a model might have been different in those papers.

3 The log-ratio log-intensity transform

Especially in two-color but also in oligonucleotide microarray experiments, it is convenient to do statistical analysis on the log-ratios and the log-intensities [8] of the gene-expression levels in two channels instead of on the expression levels directly. For gene $i$ we have that

$$M_i = \log_2 \frac{y_{R,i}}{y_{G,i}} = \log_2 \frac{f_R(x_{R,i})}{f_G(x_{G,i})}$$  \hspace{1cm} (5)

$$A_i = \frac{1}{2} \log_2 (y_{R,i} \cdot y_{G,i}) = \frac{1}{2} \log_2 (f_R(x_{R,i}) \cdot f_G(x_{G,i})).$$  \hspace{1cm} (6)

For clarification (and for simplicity) we denoted the two channels by $R$ and $G$ instead of 1 and 2, which are mnemonics for the red and the green dyes commonly
Figure 1: Log-log plot of the transformed red (dashed) and green (solid) signals as a function of the true signals under the affine transformation \((a_G, a_R) = (200, 20)\) and \((b_G, b_R) = (1.4, 0.8)\). The light blue solid line represents the \(x_R = x_G\) line. Since \(b_G > 1\), \(f_G(x_G)\) will be saturated at high intensities.

used in two-color microarray data. It may also be of interest to compare log-ratios within the same channel, which is the case for instance in single-channel normalization and calibration methods [7, 6, 37, 32]. One of the rationales for this bijective transform (if the observed signals are positive) is that the main measure of interest, the fold change, is contained in one variable. However, since the transform is based on observed expression levels and not the true ones, the conclusion that all information about the biological fold change is contained in \(M\) only is not necessarily true. This can be seen if we consider the true fold change for an arbitrary gene \(i\)

\[
r_i = \frac{x_{R,i}}{x_{G,i}}
\]

where \(r_i > 0\). If we drop the gene index \(i\) in (5) and (6), \(M\) and \(A\) can be written as functions of \(x_G\) and \(r\), i.e. \(M = g_r(x_G)\) and \(A = h_r(x_G)\). Now, we can write

\[
M = m_r(A) = g_r(h_r^{-1}(A)).
\]

It is not hard to see that the log-ratios \(M\) are independent of the log-intensities \(A\) for all fold changes if and only if \(f_R(\cdot) = f_G(\cdot)\). Hence, and discussed thoroughly
below, commonly observed intensity-dependent effects in the log-ratios may contain valuable information, and consequently, applying normalization methods without care may result in loss of information and introduced bias.

3. Log-ratios as a function of log-intensities

Under an affine transformation one can show (appendix A.2) that the relationship between the observed log-ratios and the observed log-intensities for fold change $r$ is

$$M = m_r(A) = \log_2 r + M_{\text{bias}}$$

$$M_{\text{bias}} = \log_2 b + \log_2 \left( \frac{1}{2} (a_R - r ba_G) + \sqrt{\frac{1}{4} (a_R - r ba_G)^2 + rb 2^A} \right)$$

where $b = b_R/b_G$ is the relative scale factor between the two channels compared.

Recall that $\log_2 r$ (the true log-ratio) is the variable of interest. The derivative of $M$ with respect to $A$ for a fixed fold change $r$ is

$$\left. \frac{dM}{dA} \right|_{x_R=rx_G} (A) = - \frac{a_R - r ba_G}{\frac{1}{4} (a_R - r ba_G)^2 + rb 2^A}. \quad (11)$$

Regarding $r$ as fixed, there are only two parameters in (9)–(11) that determine the shape of the function $m_r(A)$. The first is $b = b_R/b_G$ and the second is $a_R - r ba_G$. Consequently, when $a_R, a_G \neq 0$, $M$ is independent of $A$ if and only if $r = (b_G a_R)/(b_R a_G)$. For this particular value of $r$, we have that the observed log-ratio is $M = \log_2 (a_R/a_G)$, which is independent of the scale factors. Moreover, for log-ratios of non-differential expressions, that is $\log_2 r = 0$, to be independent of $A$, it must be true that $b_G a_R = b_R a_G$ or, equivalently, $b_R/b_G = a_R/a_G$. It is also clear from Equation (9) and Equation (11) that the scale parameters cannot introduce any curvature themselves, which was incorrectly argued in [12], but only enhance or suppress curvature introduced by the bias. In addition to this, the relative scale will shift the log-ratios up or down. In the case of a linear transform ($a_R = a_G = 0$) we notice that $M$ is independent of $A$ for any choice of $r$ and that the bias in the log-ratios is $\log_2 (b_G/b_R)$. Moreover,
the size of the effect that the bias terms have on the log-ratios decreases as the intensity increases. At high intensities the only observable effect is that from the relative scale between the two channels. The observed log-ratio for non-differentially expressed genes at high intensity is $M \approx \log_2 b$. The weakest data point possible to observe is $(A_0, M_0) = (1/2 \cdot \log_2(a_R/a_G), \log_2(a_R/a_G))$, which is independent of both gene expression and scale parameters. If $a_R, a_G > 0$, all fold-change curves will converge to this point. In the left graph in Figure 2 the

![Graph](image)

Figure 2: Affine transformation of the red and the green signals where $(a_G, a_R) = (200, 20)$ and $(b_G, b_R) = (1.4, 0.8)$. **Left:** The observed log-ratios as a function of the observed log-intensities for different fold changes. The blue dot-dash curve corresponds to the non-differentially expressed genes and the thinner curves above and below this curve represent $\log_2 r = \pm 1, \pm 2, \ldots$ as labeled to the right of the curves. The lines in the gray grid, which is rotated 45 degrees, show the levels where the true signals $\log_2 x_R$ and $\log_2 x_G$ are equal to $\ldots, -1, 0, 1, \ldots, 16$. These levels have been labeled to the left of the grid. No observations can lie outside this grid. **Right:** Real-world example of an affine transformation. The same slide was scanned four times at four different PMT settings. For each of the six scan pairs, the within-channel log-ratio and log-intensities were calculated. Data shown is from the red channel, which was estimated to have an offset of $a_R = 20.3$ for all scans.

The effect of the affine transform $(a_G, a_R) = (200, 20)$ and $(b_G, b_R) = (1.4, 0.8)$ at different fold changes is depicted. The different curves plotted are the functions $M = m_r(A)$ for different fold changes. Non-differentially expressed genes
3. The log-ratio log-intensity transform

$\log_2 r = 0$ lie along the blue dot-dash curve, two times up and down regulated genes ($\log_2 r = \pm 1$) are the thinner curves above and below the non-differentially expressed curve. The curves outside these represent $\log_2 r = \pm 2, \pm 3, \ldots$. Note the asymmetry in curvature between up and down regulation. From the above discussion we know that the observed log-ratios are independent of the log-intensities for $\log_2 r \approx -2.51$ with the value $M_0 \approx -3.32$. The log-ratios for non-differentially expressed genes at high intensities is $M_\infty \approx -0.81$. A real-world example taken from [6], where the same array was scanned four times at various scanner PMT (sensitivity) settings, is shown in the right plot in Figure 2. Observed within-channel log-ratios $M_c = \log_2(y_c^{(v)}/y_c^{(w)})$ are plotted against the within-channel log-intensities $A_c = \log_2(y_c^{(v)}/y_c^{(w)})/2$ for the red ($c = R$) channel where $y_c^{(v)}$ and $y_c^{(w)}$ are observations at two different scanner PMT settings. In this case it turned out that all scans share the same bias. For more details see [6].

3.2 Bias in the log-ratios

From Equations (9)-(10), we see that the bias in the log-ratios introduced by the affine transform is intensity dependent. This non-linearity can be observed as a propeller shaped graph in Figure 3, where the log-ratios under the affine transform $(a_G, a_R) = (200, 20)$ and $(b_G, b_R) = (1.4, 0.8)$ are plotted against the true log-ratios at different log-intensity levels. When doing a linear regression between the affinely transformed log-ratios and the true log-ratios, the slope will always be less than one. Moreover, this will be true for all normalization methods that do not overcompensate for bias. This can explain why some studies show that cDNA microarrays tend to compress the absolute log-ratios compared to oligoarrays and QRT-PCR [38, 39, 2]; the channel biases in cDNA microarrays are probably larger. When [23] compared cDNA microarray log-ratios with Northern blot log-ratios for their background correction method they found similar behavior, which emphasizes the close relationship between bias and background estimates. We will return to this later. Furthermore, note that the relationship between observed and true log-ratios does not become one to one for large fold changes. The reason for this is partly explained by the fact when the fold change increases the signal in one channel increases, but it could also be that the signal in the other channel decreases. In such cases, the latter signal will be affected relatively more by the bias, which introduces a non-linear relationship. This effect can be observed at
the extreme fold changes in the left graph in Figure 3. The same patterns can be seen if we do an $M$ versus $M$ scatter plot for non-normalized versus (affinely) normalized data. See right scatter plot. To visualize the intensity dependency of the log-ratios, only data points at certain log-intensity levels are plotted. For details on data, see [6].

![Figure 3: Left: Bias in the log-ratios introduced by the affine transform $(a_G, a_R) = (200, 20)$ and $(b_G, b_R) = (1.4, 0.8)$. Each line displays the relationship between the observed and the true log-ratios at a certain (observed) log-intensity $A$. Each curve is marked with the value of $A$. We have chosen to truncate the curves when the signals become saturated and the labels for those curves are positioned approximately where they have been truncated. For low intensities there is a great bias (deviance from the diagonal line), especially for large fold changes. At higher intensities the bias is smaller. The curves intersect at the one fold-change level that is independent of the intensity. Right: Real-world example of log-ratios for non-normalized versus (affinely) normalized signals. To clarify the intensity-dependent effect only data points close to $A = 5.0, 5.5, \ldots, 16$ are shown.]

### 4 Normalization

Depending on the design of the microarray experiment different types of patterns in data are expected to be observed. A typical example is where a subset of the genes studied is expected to be non-differentially expressed in a test sample
4. Normalization

compared to a reference sample. It is common that the meaning of subset is “most
genes”, “house-keeping genes”, or similar. Furthermore, some genes are expected
to not be expressed at all, in one or both channels. However, it is common that
the patterns of the observed expression levels are not in line with the expected
patterns of the true expression levels. Whenever this happens different approaches
can be taken to make the observed data meet the expectations. Normalization of
microarray data is about identifying and removing such artifactual variations that
are not due to noise or natural variability. An example is the intensity-dependent
log-ratio artifact, which we already have shown can, to a large extent, be explained
by biases in the different channels.

In this section we will, with the affine model in mind, revisit various more
or less well known normalization methods that directly or indirectly remove
intensity-dependent artifacts. The most well known is the lowess or curve-fit
normalization method that, based on the assumption that most genes are
non-differentially expressed, fits a curve through data and shifts the signals such
that the log-ratios lie along the $M = 0$ line. Other models that rely on the same
assumption, but normalize data differently, are the perpendicular and parallel
translation normalization methods. These are discussed below. Although they
are design oriented, we will also discuss various so called dye-swap normalization
methods. More sophisticated are the quantile normalization methods, which rely
on the assumption that the distribution of the gene-expression levels should be
the same for all channels and arrays. These can also be understood in the light
of the affine measurement function. With the gained knowledge we will then
propose a generic and robust normalization method based on the affine model.

To be more precise in what follows, we will refer to methods that correct for
differences in observed and expected data, that is, conform the signals to a
standard or a norm, as normalization methods, where normalization has the
meaning of conforming to expectations. Sometimes calibration data, also known
as control data, which contains true relative or absolute expression levels, is
available. Such data can be used to correct for discrepancies between the observed
and the true expression levels. We refer to methods that use calibration (read
known) data points to correct for artifacts as calibration methods. To this category
we also count methods that are based on models for which we can find the inverse
of the measurement function. We will not discuss calibration methods here.
In general, a normalization method is only capable of estimating \( a_R - b a_G \) in Equation (10) and not the individual bias terms. This is because the assumption that most genes are non-differentially expressed (and/or that there is an equal amount of up and down regulated genes) will only help us identify one fold-change curve, namely \( \log_2 r = 0 \). For a normalization method, like most calibration methods, to be able to estimate both \( a_R \) and \( a_G \) more constraints are needed and without known data this can only be done based on more assumptions. As more research is needed, we will not elaborate with such additional assumptions in this report. Thus, the rest of this report will only discuss normalization methods based on the commonly accepted assumption that it is possible to identify a set of genes that can be used to normalize the non-differentially expressed genes.

4.1 Curve-fit normalization revisited

When [35] first observed the intensity-dependent effects on the log-ratios they suggested a curve-fit normalization method that is often referred to as lo(we)s normalization. The simplest version of this assumes that the majority of the genes are non-differentially expressed and for this reason the log-ratios are expected to be centered around zero for all intensities. Under the above assumption curves estimated using robust local regression methods such as lowess [9, 10] or loess [11], or curves modeled by smoothing splines [15] will be good approximations of the \( m_r=1(A) \) function, which then can be subtracted from the observed log-ratios

\[
M \leftarrow M - m_{r=1}(A) = m_r(A) - m_{r=1}(A).
\]  

(12)

Under an affine transform, \( m_r(A) \) and \( m_{r=1}(A) \) are as in Equation (9), but we do not know of a closed form expression for (12). An example of a curve-fit normalization under the affine transform is depicted in Figure 4. Note that the asymmetry between up- and down-regulated genes is not corrected for. Moreover, if we look at the overlaid \((\log_2 x_G, \log_2 x_R)\) grid in the left graph of Figure 4, we find that the curve-fit normalization warps it and removes the otherwise orthogonal relationship between \(\log_2 x_R\) and \(\log_2 x_G\) (if the \((2A, M)\) plane is considered).

4.2 Perpendicular translation normalization revisited

The (perpendicular) shift-log method proposed in [21] is a normalization method that corrects for differences in the channel biases. It normalizes the log-ratios us-
4. Normalization

Figure 4: Curve-fit normalization of affinely transformed data where \((a_G, a_R) = (200, 20)\) and \((b_G, b_R) = (1.4, 0.8)\). Left: Log-ratios as a function of log-intensities for different fold changes. Note that the distance between up and down-regulated genes at any intensity is the same before and after the normalization. Right: Normalized log-ratios versus true log-ratios. We see that intensity-dependent artifacts have been removed for the observed and true log-ratios where all curves intersect (here at \((0, 0)\)).

A \(= \log_2(y_R y_G)\)

\[
M = \log_2(y_R y_G)
\]

\[
r = \log_2(x_R x_G)
\]

\[
y_{R,i} \leftarrow a_R + b_R x_{R,i} + a
\]

\[
y_{G,i} \leftarrow a_G + b_G x_{G,i} - a
\]

for all spots \(i\). We refer to this translation normalization transform as the perpendicular translation normalization, because it moves data \((x_G, x_R)\) perpendicular to the \(x_R = x_G\) line. From Equation (10), we get that the observed log-ratios \(m_r(A)\) can be made independent of the intensities if and only if

\[
a = \frac{r b_R a_G - b_G a_R}{b_G + r b_R}, \quad r > 0.
\]

As this is a function of \(r\), it is only for a single fold change at the time this method can make \(M\) independent of \(A\). The most common choice is \(r = 1\) for which
the optimal perpendicular shift is

\[ a = \frac{b_R a_G - b_G a_R}{b_G + b_R}, \]  

which is the weighted difference between \( a_R \) and \( a_G \) with weights \( b_G/(b_G + b_R) \) and \( b_R/(b_G + b_R) \), respectively. The distance from the \( r = 1 \) curve to the \( M = 0 \) curve for the optimal perpendicular shift is \( \log_2 b \). In other words, the perpendicular shift normalization will not remove an overall bias in the log-ratios (although it is not hard to estimate \( b \) afterward). The optimal shift for \((a_G, a_R) = (200, 20)\) and \((b_G, b_R) = (1.4, 0.8)\) is \( a = 60 \). The distance between the \( r = 1 \) and the \( M = 0 \) curve is 0.57. The result of this normalization is depicted in Figure 5. Note that \( m_r(A) \) after normalization is constant for \( r = 1 \).

Figure 5: Perpendicular translation normalization of affinely transformed data where \((a_G, a_R) = (200, 20)\) and \((b_G, b_R) = (1.4, 0.8)\). The optimal amount of normalization shift in the raw data is \( a = 60 \), which corresponds to \( a'_R = 80 \) and \( a'_G = 140 \). Left: Log-ratios as a function of log-intensities for certain fold changes. The \( r = 1 \) curve (dot-dash blue) is horizontal, that is, for this specific value of \( r \) and \( a \) the log-ratios are independent of the log-intensities. Right: Normalized log-ratios versus true log-ratios. From this graph it is clear that we obtain the minimum error in log-ratios at zero-fold change. The dotted curves correspond to the minimum and maximum log-intensities possible to observe.
As suggested by [21], one way to find the optimal shift $a$ is to minimize the curvature by minimizing the variation of the log-ratios after applying the shift $a$. To do this robustly, the median absolute deviation (MAD) can be used as measure of variation:

$$
\hat{a} = \arg\min_a \text{MAD}_{1\le i \le I}(M_i(a)).
$$

A problem with the perpendicular translation normalization methods is that the optimal shift can result in non-positive signals making a huge number of expression ratios invalid. The normalization method discussed next does not have this problem, but on the other hand, it will not work or work badly under certain conditions.

### 4.3 Parallel translation normalization revisited

Another very similar normalization method that corrects for differences in biases in two channels is what we refer to as the *parallel translation normalization* method. For historical reasons, but also because it will contribute to our discussion about background correction, the shift-log method proposed by [26] for stabilizing (read decreasing or shrinking) the variance of the measured log-ratios is of interest. A side effect of their method is that it can correct for intensity-dependent curvature. It is based on a translation transform where the same constant is added to the signals in both channels:

$$
\begin{align*}
y_R &\leftarrow a_R + b_R x_R + a \\
y_G &\leftarrow a_G + b_G x_G + a
\end{align*}
$$

where in their paper $a \geq 0$, but it is possible that $a$ is negative. We refer to this special affine translation normalization as the *parallel translation normalization*, since it moves data $(x_G, x_R)$ parallel to the $x_R = x_G$ line. Again, as this is a function of $r$, $M$ can only be made independent of $A$ for one unique $r$ at the time. See Equation (10). For $r = 1$ the optimal parallel shift is

$$
a = \frac{b_R a_G - b_G a_R}{b_G - b_R}, \quad b_G \neq b_R,
$$

which can be estimated as in (17). For example, if $(a_G, a_R) = (200, 20)$ and $(b_G, b_R) = (1.4, 0.8)$, the optimal parallel shift is $a = 220$ with the $r = 1$ curve
Figure 6: Parallel translation normalization of affinely transformed data where $(a_G, a_R) = (200, 20)$ and $(b_G, b_R) = (1.4, 0.8)$. The optimal amount of normalization shift in the raw data is $a = 220$, which corresponds to an effective shift of $(a'_G, a'_R) = (420, 240)$. Left: Log-ratios as a function of log-intensities for certain fold changes. The $r = 1$ curve (dot-dash blue) is horizontal, that is, for this specific value of $r$ and $a$ the log-ratios are independent of the log-intensities. Right: Normalized log-ratios versus true log-ratios. From this graph it is clear that we obtain the minimum error in log-ratios at zero-fold change.

0.57 units below the $M = 0$ line. The result of this normalization is depicted in Figure 6. From the above expression, we also see that an optimal value of $a$ indeed can be negative. For example, if $(a_G, a_R) = (200, 140)$ and $(b_G, b_R) = (1.4, 0.8)$, the optimal parallel shift is $a = -60$, which corresponds to an effective shift of $(a'_G, a'_R) = (140, 80)$. However, it can also result in non-positive signals and therefore undefined log-ratios. For example, if $(a_G, a_R) = (200, 20)$ and $(b_G, b_R) = (1.4, 0.8)$, the optimal parallel shift is $a = -440$, which corresponds to an effective shift of $(a'_G, a'_R) = (-420, -240)$. Moreover, from (20) we see that when the scale parameters are equal there is no solution. For $r = 1$ the derivative is independent of $a$. This is because in such cases data is moved in parallel to the $x_R = x_G$ line making it impossible to get closer. As in the case of the perpendicular shift normalization, the distance between the $r = 1$ curve and the $M = 0$ curve is $\log_2 b$. Hence, the parallel shift normalization will not remove an overall bias in the log-ratios either and rescaling is necessary.
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4.4 Single-channel translation normalization

The perpendicular shift normalization method has the problem that it can introduce non-positive signals and the parallel shift normalization method works badly or not at all under certain conditions. A hybrid of the two methods is a normalization method that translates the signals in one of the channels at the time according to

\[ y_R \leftarrow a_R + b_R x_R + a \cdot I(a \geq 0) \]  
\[ y_G \leftarrow a_G + b_G x_G - a \cdot I(a < 0), \]  

where \( I(\cdot) \) is the indicator function. This will not generate non-positive signals as only positive translations are applied. Moreover, because only one channel is shifted an optimal shift will always be found. The same objective function as before can be used to find the optimal shift. However, the derivative of the objective function will be discontinuous at \( a = 0 \), but in practice this is not a problem and \( a \) can be estimated as done previously.

4.5 Rescale normalization

The above translation normalization methods removed the curvature by adjusting the bias parameters in \( a_R - b a_G \) keeping the relative scale \( b \) fixed. Similarly, if we keep the bias parameters fixed, we can remove curvature by adjusting the relative scale \( b \). We have shown that the scanner introduces scale (PMT) insensitive biases in both the red and the green channel [6]. Thus, by adjusting the PMT settings such that the curvature of the pre-scanned data is as small as possible one minimizes \( |a_R - b a_G| \). Indeed, this strategy is in practice used by many. However, from above we know that this can equally well be done numerically. It is much more important to adjust the PMT (and laser) settings such that the dynamical range of the signals is as large as possible. Furthermore, as scanner settings are often adjusted for each array separately, there will be a discrepancy between arrays, which in any case has to be normalized for.

4.6 Dye-swap normalization revisited

The dye-swap normalization, also known as the reverse labeling and the paired-slides normalization, is a balanced experimental design for two-color microarrays that can be used whenever two technically replicated hybridizations are available.
Assume we have two different sets of cell populations, \( A \) and \( B \), for which we would like to compare the relative gene expressions \( r = \frac{x_A}{x_B} \) for all genes. After cDNA is obtained through reverse transcription the two samples are each split into two identical parts, one which is labeled with a red fluorescent dye and one which is labeled with a green fluorescent dye. We mix the red cDNA cocktail from the sample \( A \) with the green ditto from the sample \( B \) and let them co-hybridize to the DNA on the first array. After scanning we observe \( (f_{G1}(x_B), f_{R1}(x_A)) \). We do the same for the remaining red-green pair for which we obtain the observed expression levels \( (f_{G2}(x_A), f_{R2}(x_B)) \). Now, applying the dye-swap normalization suggested by [36] we get that

\[
M = \frac{1}{2}(M_1 + M_2) = \frac{1}{2} \left( \log_2 \frac{f_{R1}(x_A)}{f_{G1}(x_B)} - \log_2 \frac{f_{R2}(x_B)}{f_{G2}(x_A)} \right)
\]

and similarly for the log-intensities

\[
A = \frac{1}{2}(A_1 + A_2) = \frac{1}{4} \left( \log_2 [f_{R1}(x_A)f_{G1}(x_B)] + \log_2 [f_{R2}(x_B)f_{G2}(x_A)] \right)
\]

Thus, the result of a dye-swap can be written as the average of two “virtual” hybridizations \( (A_1', M_1') \) and \( (A_2', M_2') \). Moreover, if (and only if) the measurement functions are equal for each array, that is, \( f_{R1}(\cdot) = f_{R2}(\cdot) \) and \( f_{G1}(\cdot) = f_{G2}(\cdot) \), then the observed ratios will be identical to the true ratios for non-differentially expressed genes. For this to be true for differentially expressed genes we already know that they also have to be linear, that is, affine with zero intercept.

Several authors [25, 22] have reported that dye-swap normalization does remove curvature, but less successful results have also been reported [33]. To better understand the reasons why and when dye-swap normalization works and not, we dissect the measurement functions \( f_c(\cdot) \) of the four channels \( c = R_1, G_1, R_2, G_2 \) into \( (v_c \circ u_c \circ t_c \circ s_c)(\cdot) \) where \( s_c(\cdot) \) models the process of all steps up to the step where the (not yet labeled) cDNA sample is obtained, \( t_c(\cdot) \) models the labeling, and \( u_c(\cdot) \) models the following steps including the hybridization, and \( v_c(\cdot) \)
models the scanning etc. As channel $R_1$ and $G_2$ are from sample $A$ and the other two are from sample $B$, we know that $s_{R_1}(\cdot) = s_{G_2}(\cdot) = s_A(\cdot)$ and $s_{R_2}(\cdot) = s_{G_1}(\cdot) = s_B(\cdot)$. Furthermore, if the labeling process is well controlled we can assume that $t_{R_1}(\cdot) \approx t_{R_2}(\cdot) \approx t_R(\cdot)$ and $t_{G_1}(\cdot) \approx t_{G_2}(\cdot) \approx t_G(\cdot)$. When hybridizing channel 1 and 3 to array one and the other two to array 2 we approximately have that $u_{R_1}(\cdot) \approx u_{G_1}(\cdot) \approx u_1(\cdot)$ and $u_{R_2}(\cdot) \approx u_{G_2}(\cdot) \approx u_2(\cdot)$. The properties of the glass arrays can be incorporated into these two measurement functions. Moreover, if we keep the same scanner settings for both arrays and assume everything else equal, we have that $v_{R_1}(\cdot) \approx v_{R_2}(\cdot) \approx v_R(\cdot)$ and $v_{G_1}(\cdot) \approx v_{G_2}(\cdot) \approx v_G(\cdot)$. The overall measurement function for the channels are then approximately

\begin{align}
    f_{R_1} &= v_R \circ u_1 \circ t_R \circ s_A \\
    f_{G_1} &= v_G \circ u_1 \circ t_G \circ s_B \\
    f_{R_2} &= v_R \circ u_2 \circ t_R \circ s_B \\
    f_{G_2} &= v_G \circ u_2 \circ t_G \circ s_A.
\end{align}

For the dye-swap normalization to be efficient, we conclude that we must control the process of extracting the RNA etc. to an extent such that we can expect $s_A(\cdot) \approx s_B(\cdot)$. Moreover, we must also be able to reproduce hybridizations well on high quality arrays such that $u_1(\cdot) \approx u_2(\cdot)$. If these requirements are met, then any differences in physical properties of the dyes or the red and the green channels in general will be self-normalized.

Turning to the affine model, from Equation (23) we get that a dye-swap normalization of affinely transformation data will result in the two virtual hybridizations

\begin{align}
    M'_1 &= \log_2 \frac{a_R + b_R x_A}{a_R + b_R x_B}, \\
    M'_2 &= \log_2 \frac{a_G + b_G x_A}{a_G + b_G x_B},
\end{align}

and similar for $A'_1$ and $A'_2$. For both of them, the signals in both channels have undergone identical affine transformations. We know from before that identical transformation in both channels does not introduce curvature for the non-differentially expressed genes and that there is a symmetry between up- and down-regulated genes, cf. perpendicular and parallel shift normalization. Moreover, optimally there is no bias and the above simplifies to $M = \log_2 (x_A / x_B)$ and $A = \frac{1}{2} \log_2 (x_A x_B)$ as in [36], which are the true expression levels. If the
biases in any of the two replicated channels are not equal ($a_{R_1} \neq a_{R_2}$), the dye-swap normalization will not work.

The above discussion assumed that the same cell samples have been replicated. If biological replicates are used, an additional source of variability is introduced. However, as long as we can assume that for most genes $x_{A_1} \approx x_{A_2}$ and $x_{B_1} \approx x_{B_2}$, dye-swap normalization should still perform well.

4.7 Alternative dye-swap normalization

An alternative to the above dye-swap normalization method is to average the observed expression levels before taking the logarithm

$$M = \log_2 \left( \frac{f_{R_1}(x_A) + f_{G_2}(x_A)}{f_{R_2}(x_B) + f_{G_1}(x_B)} \right)^2 = \log_2 \frac{f_{R_1}(x_A) + f_{G_2}(x_A)}{f_{R_2}(x_B) + f_{G_1}(x_B)},$$

(30)

and analogously for $A$. This approach uses the arithmetic mean of the observed signals whereas the previous dye-swap method used the geometric mean. To be able to say more about the difference between the two approaches, we turn to the affine transformation for which we have

$$M = \log_2 \left( \frac{(a_R + a_G) + (b_R + b_G)x_A}{(a_R + a_G) + (b_R + b_G)x_B} \right),$$

(31)

$$A = \frac{1}{2} \log_2 \left[ \frac{(a_R + a_G) + (b_R + b_G)x_A}{(a_R + a_G) + (b_R + b_G)x_B} \right].$$

(32)

Again, we note that the dye-swap method makes the transforms in the resulting two virtual channels equal. However, comparing the bias in log-intensities between the geometrical and the arithmetical approaches, we see that for the latter we have

$$A_0 = \frac{1}{2} \log_2 \left( \frac{1}{4} (a_R + a_G)(a_R + a_G) \right) = \log_2 \frac{a_R + a_G}{2},$$

(33)

whereas for the former we have

$$A_0 = \frac{1}{2}(A'_{1,0} + A'_{2,0}) = \frac{1}{2} \left( \frac{1}{2} \log_2 a_R^2 + \frac{1}{2} \log_2 a_G^2 \right) = \log_2 \sqrt{a_R a_G}. \quad (34)$$
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Because \((a_R + a_G)/2 \geq \sqrt{a_R a_G}\), we conclude that \(A_0\) is always larger for arithmetic dye-swap than for the geometric one, and from previous results we know that the larger \(A_0\) is the more curvature we have for the differentially expressed genes.

However, there are other differences too. For instance, if each microarray glass array (the \(u_c(\cdot)\) functions above) introduces the same additive bias to both channels and this bias is different between arrays, but otherwise everything else is the same, that is, \(a_{R_2} = a_{R_1} + a\) and \(a_{G_2} = a_{G_1} + a\), then the arithmetic dye-swap would do better

\[
M_a = \log_2 \frac{(a_{R_1} + a_{G_1} + a) + (b_R + b_G)x_A}{(a_{R_1} + a + a_{G_1}) + (b_R + b_G)x_B}
\]  

(35)

than the geometric dye-swap

\[
M_g = \frac{1}{2} \left( \log_2 \frac{a_{R_1} + b_Rx_A}{(a_{R_1} + a) + b_Rx_B} + \log_2 \frac{a_{G_1} + a + b_Gx_A}{a_{G_1} + b_Gx_B} \right),
\]

(36)

which in such case fails to remove curvature.

4.8 Two-channel quantile normalization

Two-channel, or in general multi-channel quantile normalization [7, 37] is based on and relies on the assumption that the true gene-expression levels in the two biological samples are approximately equally distributed. If the measurement functions in the two channels, say \(f_R(\cdot)\) and \(f_G(\cdot)\), are different, then the distributions of the measured signals in the two channels are different even if the underlying distributions of the true expression levels would be identical. By estimating the distributions of the two channels and making them equal, for instance to an average distribution, the log-ratios for the non-differentially expressed genes will be unbiased and independent of the intensities. Thus, making the density functions of the measured data equal for the two channels is the same as making their transformation functions equal, say to \(f_{RG}(\cdot)\), and equal transformation functions will make \(M\) independent of \(A\) for the non-differentially expressed genes. If \(f_{RG}(\cdot)\) could be made linear too, this would be true for all fold changes.

If we turn to the affine transform, two-channel quantile normalization succeeds to remove intensity-dependent effects, because the bias \(a_R\) and \(a_G\) in the two channels are made identical when the quantiles of the signals are made the same. The
elimination of the bias in log-ratios is because $b_R$ are made equal to $b_G$. Hence, two-channel quantile normalization can be considered to be both a method that corrects for differences in bias between two channels, but also a method that corrects for biases in the expression ratios. In Figure 7, the two-channel quan-

tile normalization of affinely transformed data where $(a_G, a_R) = (200, 20)$ and $(b_G, b_R) = (1.4, 0.8)$ is depicted. To the left are the affine transformations for the red and the green channels shown. The (upside-down) curve at the bottom shows a hypothetical density function of the true (log) gene-expression levels expected to be equal in both samples. The distributions of the affinely transformed signals are shown in the (rotated) density functions at the left (red and green curves). The average signal density (middle gray curve) to be normalized toward corresponds to a common measurement function (gray function in the main plot). The two-channel quantile normalized data is shown in the $M$ versus $A$ plot to the right. We see that the curvature of the non-differentially expressed genes is removed.

Figure 7: Equalizing the signal densities of the two channels remove the intensity dependency of the log-ratios of non-differentially expressed genes. Left: Equal gene-expression distributions in both channels will under the non-channel balanced affine transform where $(a_G, a_R) = (200, 20)$ and $(b_G, b_R) = (1.4, 0.8)$ turn into two different densities for the measured data. Right: Normalizing the non-equal densities of the two channels makes the log-ratios of the non-differentially expressed genes zero for all intensities. All axes are logged.
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4.9 Background subtraction as a normalization or calibration method

We have observed that the log-ratios of the background signals show the same intensity-dependent effects as the foreground signals, which suggests that the background signals undergo the same transformation as the foreground signals. An example of this is shown in Figure 8, where the background and the foreground estimates from one hybridization are plotted in the same M versus A scatter plots. This may be explained by the results in [6]. A widely adopted rationale for background correction is the assumption that the region that defines the spot is contaminated with the same physical noise that can be observed in the surrounding regions. Background noise is commonly explained to be due to dust particles, DNA contaminated buffers, failed washing during printing or hybridization, cross hybridization etc. [23, 29]. This type of background noise is often assumed to add to the foreground signal. Thus, to obtain the true signal the background is subtracted from the foreground signal

\[ y_{c,i}^{\text{(fg)}} - y_{c,i}^{\text{(bg)}} \]  

(37)
where \( y^{(f)\text{c},i} \) is the estimated foreground signal and \( y^{(bg)\text{c},i} \) is the estimated background signal for channel \( c \) and spot \( i \). Under a transformation that is dominated by an affine function for lower intensities (of the same order as the background), subtracting the background from the foreground will shift the biases toward zero and background subtracted signals will have less curvature in the \((A, M)\) plane than non-background subtracted signals (not shown). In this sense we can consider background subtraction to be a normalization (or a calibration) method. However, just because the log-ratios as a function of the log-intensities becomes more flat, it does not mean that the foreground estimates are contaminated by the same signal that is found in the background area (commonly the regions surrounding the spots). Instead, the explanation could be that the background estimates happen to be approximately equal to the amount of shift in the foreground signals. Different image analysis software estimate the background signal differently based on different algorithms such as fixed-size circles, adaptive circles, morphological estimates, and pixel intensity distributions. Although comparative studies have been conducted [34, 3], it is still not clear which background estimate is most correct. Some methods give higher background estimates than others, which means that they all correct for channel biases by different amounts, which is another argument for why we most likely do have biases in our signals. Moreover, subtracting the background from the foreground signals commonly results in non-positive signals, which makes data harder to analyze, but also to interpret. Even though the background estimates were known to be unbiased, it is not surprising to get non-positive signals under the assumption that the foreground region is contaminated by the same noise as found in the background region. This is because both the foreground and the background signals are estimates. A pioneering work that makes use of this is [23], which emphasizes that the true signal can \textit{not} be negative and uses a Bayesian approach to correct for this.

4.10 Result of a (relative) negative translation

If, for instance, too much background is subtracted or a threshold has to be passed before the reverse transcription takes place, one can imagine that \( a_G, a_R < 0 \). With negative shift the curves in Figure 2 bend downward instead of toward the left. Negative bias also applies if the observed signals are compared, not to the true signals, but to the signals obtained by another measuring technique that has a higher bias. Examples of such comparisons can be two-color microarray data.
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compared to oligonucleotide (Affymetrix) data or two-color microarray data compared to QRT-PCR data. Negative bias can also be observed when control clones, spike-ins, negative and positive controls etc. are compared with the genes/ESTs of interest. The effect of a negative translation is depicted in Figure 9. The fan-out effect in the fold-change curves for the lower intensities is due to the negative translation. Note that this should not be mistaken for the fan-out effect due to decreasing signal-to-noise levels.

\begin{align*}
A &= \log_2(y_R/y_G) \\
M &= \log_2(y_R/y_G) \\
r &= \log_2(x_R/x_G)
\end{align*}

Figure 9: Affine transformation of the red and the green signals with negative translation where \((a_G, a_R) = (-87, -24), (b_G, b_R) = (1.4, 0.8)\). Left: Log-ratios as a function of log-intensities for certain fold changes. Right: Translated log-ratios versus true log-ratios. The slope of a line fitted in the \(M\) versus \(M\) plot will be larger than one, which is due to the negative translation. Note that we have truncated both fold-change curves and the grid in the left graph and the intensity curves in the right graph such that \(x_R, x_G \geq 1\).

4.11 Robust affine normalization

It is clear that it is essential to correct for biases when two or multi-channel cDNA microarray data is normalized. We can obtain estimates of \(a_R, a_G\) and \(b\) as follows. Define \(\alpha = a_R - b a_G\) and \(\beta = b = b_R/b_G\). For non-differentially expressed genes (without noise) we have that

\[ y_{R,i} = \alpha + \beta y_{G,i}; \ i = 1, 2, \ldots, I. \]
Define \( y = \{ y_i \}_{i=1}^{I} \) where \( y_i = (y_{G,i}, y_{R,i}) \) and let

\[
Q(\alpha, \beta; y) = \sum_{i=1}^{I} w_ir_i(\alpha, \beta; y_i)^2
\]

be our objective function where \( r_i(\alpha, \beta; y_i) > 0 \) is the orthogonal Euclidean distance between \( y_i \) and the line \( L(\alpha, \beta) \) with intercept \( \alpha \) and slope \( \beta \). The estimate of \( \alpha \) and \( \beta \) is then

\[
(\hat{\alpha}, \hat{\beta}) = \arg \min_{(\alpha, \beta)} Q(\alpha, \beta; y).
\]

With \( w_i = 1 \) for all observations we obtain standard principal component analysis (PCA), which minimizes the orthogonal distances in the \( L_2 \) norm. With weights \( w_i = 1/(r_i(\hat{\alpha}, \hat{\beta}; y_i) + \delta) \) we can minimize the distances in the \( L_1 \) norm, if we let \( \delta \to 0^+ \). Other robust estimators can be obtained by choosing other weight functions, but we choose to optimize in \( L_1 \). Moreover, if one suspects a non-symmetric distribution of data points around the line, then a trimmed version of the weight function should be considered. In practice, the above optimization can be done by an iterative reweighted principal component analysis (IWPCA) scheme. For iteration \( l = 1, 2, \ldots \), minimize (39) using weighted PCA where \( w_i^{(1)} = 1 \) and \( w_i^{(l+1)} = 1/(r_i(\alpha^{(l)}, \beta^{(l)}; y_i) + \delta) \) with \( \delta \) being a small positive number to avoid infinite weights.

As a last step, in order to get estimates of the three parameters \( a_R, a_G \) and \( b \) from the two parameter estimates \( \hat{\alpha} \) and \( \hat{\beta} \), we introduce an additional constraint. Let \( y_{c,\min} = \min_i y_{c,i} \) for \( c = \{ R, G \} \). We then choose

\[
\hat{b} = \hat{\beta}
\]

\[
\hat{a}_G = \max\{a_G; a_G < y_{G,\min} \land \hat{\alpha} + \hat{b}a_G < y_{R,\min}\}
\]

\[
\hat{a}_R = \hat{\alpha} + \hat{b}\hat{a}_G
\]

to be the estimates of the bias and the scale parameters in model (3). This constraint is only correct in the error free model. If we allow noise, say

\[
y_{R,i} = a_R + b_Rx_{R,i} + \varepsilon_{R,i}
\]

\[
y_{G,i} = a_G + b_Gx_{G,i} + \varepsilon_{G,i},
\]

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where $E[\varepsilon_{c,i}] = 0$ and $V[\varepsilon_{c,i}] = \sigma^2_{c,i}$ for $c = \{R, G\}$, it is possible that the bias terms $a_R$ and $a_G$ are indeed larger than the smallest observed value in the respective channel. This is especially important if the distributions of $\varepsilon_{c,i}$ for $c = \{R, G\}$ have heavy negative tails. However, as already mentioned, we do not know enough about the various noise sources, and to avoid the problem with non-positive signals, we have decided to keep the above constraint. We are aware of the fact that the estimates in such cases are inconsistent, but we believe that the error we do by subtracting as much bias as possible without introducing non-positive signals, as in (41)-(43), is much smaller than not correcting for the bias at all. An alternative, which introduces negative estimates, is to replace $y_{c,\text{min}}$ in (42) with $y_{c,(i)}$, where order index $(i)$ is chosen such that $i$ non-positive signals are obtained in channel $c$.

Furthermore, it has been observed that the noise in each channel is roughly proportional to the signal strength, that is, $\sigma_{c,i} \propto x_{c,i}$. Thus, a positive side effect of the above estimation algorithm is that, contrary to having equal weights for all spots ($w_i = 1$), more weight will be given to the low-intensity spots compared to the high-intensity ones. This will make the method more robust to saturation and other non-linear effects that might occur at high intensities, effects for which classical line fits, which rely on homoscedasticity, would fail.

Finally, with backward transformation (4) based on estimates for $(\hat{a}_R, \hat{a}_G, \hat{b})$, we translate and rotate data such that it falls around the diagonal line that goes through $(0, 0)$ and $(1, 1)$. In Figure 10 an example is given where the above affine normalization method has been applied to a two-color microarray data set. It is clear that the curvature in the $M$ versus $A$ scatter plot is removed by the normalization method. For a comparison of the bias in the log-ratios, see also Figure 3. Data is the same as in [6] and a more detailed analysis of differentially expressed genes etc. will be published elsewhere.

Furthermore, if more than one array is normalized, we suggest that the signals from all arrays and both channels are normalized together. For this to be realistic, the assumption that most genes are non-differentially expressed for all possible hybridization/channel pairs must be added. However, this is not a problem since the experimental design is often set up such that this is true. For instance, in two-channel microarrays experiments it is common to hybridize one test sample
Figure 10: Log-ratios versus log-intensities before (left) and after (right) a robust affine normalization. The intensity-dependent trend of the log-ratios in the raw data is removed by the affine normalization. The affine parameters were estimated to \((a_R, a_G, b) = (21.5, 29.3, 0.163)\), which for instance gives that \((A_0, M_0) = (4.65, -0.45)\) and \(M_{\infty} = -2.62\).

and one reference sample, which is selected such that it does not differ too much from the test sample, to the same array. The same reference is then used between arrays (in either channel). Thus, since each test-reference pair is “close” to each other, all test-test pairs should approximately be “close” to each other also.

The multi-channel algorithm can be summarized as follows. Say there are \(N\) arrays each hybridized with two \(K = 2\) samples (colors). The observations \(y_i = (y_{R_{i,1}}, \ldots, y_{R_{N,i}}, y_{G_{i,1}}, \ldots, y_{G_{N,i}})\) for genes \(i = 1, 2, \ldots, I\) then span a \(KN\)-dimensional space. Analogously to the above two-dimensional procedure, we can fit a robust line \(L\) through data in \(\mathbb{R}^{KN}\) and constrain the estimate of \(a = (a_{R_1}, \ldots, a_{R_N}, a_{G_1}, \ldots, a_{G_N})\) by enforcing that \(a \prec y_i; \forall i\) where \(\prec\) is the component-wise inequality. Backward transformation will again translate and rotate data such that it lies along the diagonal line. Moreover, normalizing all arrays at once will bring the signals from all hybridizations onto the same scale and no further, so called, between-slide scale normalization is needed. This multi-dimensional affine normalization method with slightly different constraints was successfully used in a multiscan calibration two-color microarray experiment [6].

An implementation of the above algorithm is made available in the R [31, 18] package named aroma [5] (formerly known as the com.braju.sma package). The only parameter in the algorithm that has to be specified is $\delta$. However, its value is not critical and we have found that for instance $\delta = 0.02$ works well in general and is therefore the default value. Thus, in practice the algorithm is completely automatic. For instance, \texttt{normalizeAffine(rg)} will normalize all arrays and all channels in the microarray object \texttt{rg} at once. Moreover, the method can be applied to any subsets of genes separately such as print-tip groups, clone groups and spike-ins.

## 5 Discussion

In previous sections, we did not discuss the variance stabilizing methods described by [17, 14], which are based on error models that also contain channel-specific bias terms. These bias terms are estimated and corrected for. Thus, those methods do indeed correct for intensity-dependent effects. Because they are based on specific error models, but also because they stabilize the log-ratio variances, they do not fit well into the above deterministic discussion. However, we do believe that the directions drawn up by their underlying error models are promising.

If we compare the robust affine normalization method in Section 4.11 with for instance the perpendicular and the parallel translation normalization methods optimized by minimizing the curvature, we find that there are similarities, because minimizing the curvature is identical to finding estimates of the bias parameters along the line $L(\alpha, \beta; y)$. Assuming a pure affine transformation, there are also similarities to the curve-fit method, which fit approximately the same line (curve) through data. The only difference then is how data is transformed to meet the assumptions. The affine method translates and rescales data in the original domain whereas the curve-fit method operates in a rotated and log-transformed domain.

Moreover, the translation and the curve-fit methods rely on two-dimensional data (log-ratios) and it is not clear how to generalize them to multi-dimensional data, although re-iterative versions such as cyclic loess [7] and the (multi-dimensional)
contrast based method [1] have been suggested. Our affine normalization method is not limited to two-dimensional data, but can be applied to any number of channels, which means that for instance three and four-color microarray data can be normalized as easily as two-color data.

It is also interesting to note the close relationship between the quantile and affine normalization method. In the case of an affine transformation, both will fit roughly the same line (curve) through data and translate it into the original domain. The difference is how data is normalized to this line. In quantile normalization the data points are shifted such that the sample densities of both channels are made identical. This results in new measurement functions, which are not necessarily linear, but still such that for non-differentially expressed genes the ratios between them are one. However, the affine normalization model can be thought of as a quantile normalization method with special constraints on the underlying densities. An interesting continuation of the affine model and quantile normalization method would be how to relax the affine constraint by using other parametric or semi-parametric models. One possibility is to add smoothness constraints to the transformation functions using smoothing splines [15], which then also would provide a probabilistic framework. Moreover, in the spirit of [23], it would be interesting to incorporate an empirical Bayes component to “allow” non-positive signals too.

Acknowledgments

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5. Discussion

Blanceflor Boncompagni-Ludovisi née Bildt, The Royal Swedish Academy of Sciences, and The Royal Physiographic Society in Lund. Finally, the microarray data from [6] was kindly provided by the SWEGENE DNA Microarray Resource Center at the BioMedical Center B10 in Lund, supported by the Knut and Alice Wallenberg foundation through the SWEGENE consortium.
References


[31] R Development Core Team. *R Language Definition* (v1.7.0, draft), April 2003.


A Detailed calculations on the affine transform

A.1 \( x_G \) and \( rx_G \) as a function of \( A \)

We want to find an expression for \( x_G = a_{G,R}^{-1}(A) \). First, doing the variable substitution \( x_g = b_G x_G \) we have

\[
A = \frac{1}{2} \log_2 [(a_R + r b x_g)(a_G + x_g)]
\]

(46)

where \( b = b_R/b_G \) is the relative scale and \( r = x_R/x_G \) is the true fold change. From this expression we can calculate the expression for \( x_g = a_{G,R}^{-1}(A) \) as follows:

\[
\begin{align*}
(a_R + r b x_g)(a_G + x_g) &= 2^2 A \\
\iff r b x_g^2 + (a_R + r b a_G) x_g + a_R a_G - 2^2 A &= 0 \\
\iff x_g^2 + ((r b)^{-1} a_R + a_G) x_g + (r b)^{-1} (a_R a_G - 2^2 A) &= 0 \\
\iff x_g &= -\frac{1}{2} ((r b)^{-1} a_R + a_G) \\
&\pm \sqrt{\frac{1}{4} ((r b)^{-1} a_R + a_G)^2 - (r b)^{-1} (a_R a_G - 2^2 A)}
\end{align*}
\]

(47)

where we in the last step used the fact that \( x_g \geq 0 \). From this it immediately follows that

\[
x_G = b_{G,R}^{-1} x_g
\]

\[
= b_{R}^{-1} r^{-1} \left( -\frac{1}{2} (a_R + r b a_G) + \sqrt{\frac{1}{4} (a_R - r b a_G)^2 + r b 2^2 A} \right). \tag{48}
\]

which in turn gives that

\[
rx_G = b_{R}^{-1} \left( -\frac{1}{2} (a_R + r b a_G) + \sqrt{\frac{1}{4} (a_R - r b a_G)^2 + r b 2^2 A} \right). \tag{49}
\]

We note that in the special case when \( a_R = a_G = 0 \) and \( b_R = b_G = b_c \) we have that \( x_G = (1/\sqrt{r})(2^A/b_c) \) and \( rx_G = \sqrt{r}(2^A/b_c) \). 

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A. Detailed calculations on the affine transform

A.2 \( M \text{ as a function of } A \)

The following equations

\[
a_G + b_G x_G = a_G + x_g
\]

\[
= (rb)^{-1} \left( \frac{1}{2} (a_R - rb a_G) + \sqrt{\frac{1}{4} (a_R - rb a_G)^2 + rb 2^{2A}} \right)
\]

\( a_R + b_R x_R = a_R + rb x_g \)

\[
= \frac{1}{2} (a_R - rb a_G) + \sqrt{\frac{1}{4} (a_R - rb a_G)^2 + rb 2^{2A}}
\]

(50)

(51)

gives us the relation between \( M \) and \( A \) for a given fold change \( r \)

\[
M = m_r(A) = \log_2(rb)
\]

\[
+ \log_2 \left( \frac{\frac{1}{2} (a_R - rb a_G) + \sqrt{\frac{1}{4} (a_R - rb a_G)^2 + rb 2^{2A}}}{-\frac{1}{2} (a_R - rb a_G) + \sqrt{\frac{1}{4} (a_R - rb a_G)^2 + rb 2^{2A}}} \right).
\]

(52)
By scanning the same spotted oligonucleotide microarray at different photo-multiplier tube (PMT) gains we have identified a channel-specific bias present in two-channel microarray data. The observed bias was very stable between subsequent scans of the same array although the PMT gain was greatly adjusted. This indicates that the bias does not originate from a step preceding the scanner detector parts. Between arrays the estimated bias varies slightly. When comparing bias estimates based on data from the same array, but from different scanners, we have found that different scanner models introduce different amounts of bias. So do various image analysis methods. We propose a scan protocol and a constrained affine model that allows us to identify and estimate the bias in each channel. Backward transformation removes the bias and brings the signals to the same scale. The result is that systematic effects such as intensity-dependent log-ratios are removed, but also that signal densities become much more similar. Average scan signals, which have a larger dynamical range and greater signal-to-noise ratio than the individual scans, can then be obtained. An implementation in R of the above calibration method is made available for free.

Keywords: microarray scanner; bias; dark noise; offset; zero level; affine model; multiscan calibration; normalization; dynamic range; photomultiplier tube; analog-to-digital converter; image analysis; intensity-dependent effects; background subtraction.
1 Introduction

The microarray technology provides a way of simultaneously measuring transcript abundances of $10^3 - 10^5$ genes from one or more individual cell or tissue samples. A microarray, also known as a gene chip, has well defined regions or features that each consists of immobilized sequences of DNA, which each is unique to a specific gene. These regions are referred to as probes [11]. When fluorophore labeled cDNA, referred to as targets, obtained by reverse transcription of mRNA extracted from the samples of interest, is let to hybridize to the probes for a few hours, each region on the microarray will specifically bind a certain amount of hybridized DNA unique to the corresponding gene. Depending on if a two-channel or single-channel microarray platform is used, either several and differentially labeled targets are hybridized to the same array, or different targets are each hybridized to an exclusive array using identical labels. Next, the array is scanned at different wavelengths to excite the fluorescent molecules using a light source, for instance a laser. Shortly after the fluorophores have been excited they emit photons, which are registered and quantified in each position by the scanner, which results in a high-resolution digitized image for each channel. Using image analysis methods, the pixels that belong to the regions that contain the probes are identified and averaged, and an estimate of the transcript abundance for each gene is obtained. Since these estimates are obtained from a complex measurement process of several steps, it is likely that the observed signals contain not only measurement noise, but also systematic variations of different kinds [5].

In this report we give evidence for a channel-specific bias that is introduced by the scanner and most likely its detector parts. In addition to this, our results indicate that the image analysis also contributes with a small bias. The effects channel-specific biases have on the downstream microarray analysis are many [8, 5]. Probably the most well known is the non-linear intensity-dependent bias in the log-ratios. We suggest a scan protocol and a model that will allow us to estimate the biases and calibrate the observed signals accordingly. The result will be that the intensity-dependent effects are removed, but also that the effective dynamical range of the scanner is increased several times.

The outline of this report is as follows. In Section 2, a model for the observed signals as a function of the probes/target concentration and the scanner settings is derived. In the same section, details on the two-channel microarray data set used
are also given. Results are given in Section 3, followed by a discussion of possible reasons for the observed biases in Section 4. Conclusions are given in Section 5.

2 Materials and Methods

2.1 General model

Consider a microarray experiment involving genes $i = 1, 2, \ldots, I$ from RNA extracts $c = 1, 2, \ldots, C$. In single-channel microarrays each array measures the gene-expression levels in one RNA extract, whereas in two-color microarrays each array measures two RNA extracts, one in each channel. From now on, we will refer to each set of signals from each RNA extract as a *channel*. Let $\mu_{c,i}$ be the true gene-expression (transcription) level of gene $i$ in channel $c$. Ideally, statistical analysis is done on these quantities. For instance, by comparing the relative abundances in two channels, that is $r_i = \mu_{1,i}/\mu_{2,i}$ for all genes $i$, it is possible to identify genes that are significantly differentially expressed ($r_i \neq 1$). However, in reality we do not observe the true expression levels, but only the quantified feature intensities $y_{c,i}$. The general relation between the observed and the true expression levels can be written as

$$y_{c,i} = f_c(\mu_{c,i}) + \varepsilon_{c,i}, \tag{1}$$

where $f_c(\cdot)$ is an unknown channel-specific function, which we refer to as the *measurement function*, that includes all steps in the microarray process. Moreover, we assume independent intensity-dependent error terms $\varepsilon_{c,i}$ such that $E[\varepsilon_{c,i}] = 0$. Because we want to do inference based on $\mu_{c,i}$, it must be possible to find the inverse of $f_c(\cdot)$, which is possible if it is strictly increasing.

To be able to find the form of $f_c(\cdot)$, high-quality calibration data from several stages along the microarray process is required. Here we will consider a simpler case. Split the overall measurement function into two parts. The first part, $x_{c,i} = g_c(\mu_{c,i})$, models the amount of light from spot $i$ in channel $c$ that enters the photomultiplier tube (PMT) [14] as a function of the transcription level of clone $i$ in channel $c$. The second part, which is studied in this report, is $y_{c,i} = h_c(x_{c,i})$ and models the observed signal as a function of the amount of photons in channel $c$ and spot $i$ that enters the PMT. That is, it captures the nature of the scanner’s light detector, but also the image analysis methods. We want
to emphasize that the light from one spot does not necessarily originate solely from the fluorescent molecules that are attached to the hybridized target DNA. Light from other sources such as cross-hybridized target, intrinsic fluorescence from spot buffer, and scatter light may also contribute with photons of similar wavelengths.

Continuing, we will show that $h_c(\cdot)$ is almost perfectly affine. This measurement function depends also on the scanner settings, especially the scanner sensitivity, which is indicated below with the superscript $(k)$. In other words,

$$y_{c,i}^{(k)} = a_c^{(k)} + b_c^{(k)} x_{c,i} + e_{c,i}^{(k)}, \ \forall i, c$$

(2)

where for each fixed scanner setting $k$, $a_c^{(k)}$ and $b_c^{(k)}$ are channel-specific bias and scale parameters, respectively. Assume that $x_{c,i}$ is fix for all PMT voltages.

Note that the above relationship is not necessarily linear ($a_c = 0$), which has important implications on downstream analysis. For instance, when spotted as well as in-situ synthesized microarray are used, it is common to do statistical analysis on the log-ratios $M_i = \log_2(y_{R,i}/y_{G,i})$ and the log-intensities $A_i = \frac{1}{2} \log_2(y_{R,i}y_{G,i})$ for all genes $i$ [10], where we for convenience have denoted the two channels to be compared by $R$ and $G$, which are mnemonics for the red and the green dyes commonly used in two-channel microarray data, although such comparisons are not limited to within-array measurements. One of the rationales for this bijective transform is that under ideal conditions the main measure of interest, the fold change, is contained in one variable only, i.e. in $M_i$. However, even if we could identify $g_c(\cdot)$ and calibrate data for that part accordingly, a channel specific bias introduced by $h_c(\cdot)$ will introduce so called intensity-dependent dye bias in the observed log-ratios. Commonly observed intensity-dependent effects in the log-ratios [23] can be explained by the fact that the logarithm is taken on affinely transformed signals [12, 8, 5].

2.2 Data

2.2.1 Arrays and hybridization

The results presented here are based on data from nine different hybridizations performed using spotted oligonucleotide microarrays, referred to here by the letters A to H. Array A and B were hybridized in October 2003. Arrays C-G were
2. Materials and Methods

hybridized the following day and array H was hybridized seven weeks later. All arrays contain the same human oligonucleotide set (QIAGEN) and all have an identical layout of 12-by-4 print-tip groups each containing 34-by-34 (1156) spots. In total there are 55488 spots on each array. The average (GenePix) spot area is 45-50 pixels and the average center-to-center distance between the spots is approximately 12-13 pixels (120-130 \( \mu m \)). Arrays were produced by the SWE-GENE DNA Microarray Resource Centre, Department of Oncology at Lund University using a MicroGrid II 600R arrayer fitted with MicroSpot 10K pins (BioRobotics). All arrays, except array H, were spotted in the same print batch on UltraGAPS\textsuperscript{TM} coated slides (Corning Incorporated) during August 2003. Array H was spotted in October the same year. Printing was performed in a temperature (18-20\(^\circ\) C) and humidity (44-49\% RH) controlled area. After printing was completed, arrays were left in a desiccator to dry for 48 hours, rehydrated for 1 second over steaming water, snap dried on a hot plate (98\(^\circ\) C), UV-cross-linked (800 mJ/cm\(^2\)) and subsequently hybridized with various test and reference RNA samples. Samples were labeled, purified and hybridized using Pronto\textsuperscript{TM} Plus System 6 (Corning Incorporated) according to manufacturer’s instructions.

2.2.2 Scanning

Each array was scanned at four different PMT settings on two different types of scanners. First the arrays were scanned on an Agilent G2505A DNA microarray scanner (Agilent Technologies) at PMT gains 100\%, 30\%, 50\%, and 80\% (in that order). The so called dark offset, intentionally added by the Agilent scanner to all signals [1, p. 18], has been uninstalled and is therefore not the reason for the findings in this report. Arrays were then re-scanned on an Axon GenePix 4000A scanner (Axon Instruments) at PMT gains 600, 700, 800, and 500 volts (in that order), except for array A, which was scanned at 700, 800, 500 and 600 volts, and array H, which was scanned at 600, 400, 500 and 700 volts. Thus, the images obtained by the Axon scanner were bleached more than the preceding ones obtained by the Agilent scanner. For both scanners, the power of the 532 nm and the 635 nm lasers was set to 100\% and the scan resolution to 10 \( \mu m \)/pixel. Moreover, a one-pass (both channels scanned simultaneously) and one-sample-per-pixel (Lines To Average or LTA equals one) procedure was used for all scans. The Agilent scanner has a special loading mechanism for microarrays, which allows automatic scanning of subsequent arrays without human intervention. However, due to limitations in the software or the scanner, each such batch of arrays
can only be scanned at a single PMT gain. To scan at more PMT gains with the Agilent scanner, it was therefore necessary to eject and reload the arrays between different PMT settings, which means that the alignment between the scanned images may not be perfect. Contrary, for the Axon scanner the arrays were put in the scanner one by one, then scanned at all PMT settings without being moved.

2.2.3 Image analysis - spot segmentation and registration

To quantify the foreground and the background signals, the scanned images (65536 gray scales and approximately 2000-by-5600 pixels) were analyzed using both the Axon GenePix Pro v4.1.1 software (Axon Instruments) and the Spot v2 software [22, 2]. We first analyzed each image with GenePix. For each of them, the grid and spot positions were manually set and then the alignment was optimized by GenePix. These positions were re-entered and re-optimized in Spot with visual inspection to verify the correctness. Moreover, for each individual scan the image analysis software was let to find the optimal spot segmentation. Thus, what is defined as a foreground pixel may vary with PMT setting although the images are from the same array. We decided on this schema for various reasons. The first reason was that the Agilent arrays are loaded and unloaded between subsequent scans and therefore require separate spot segmentations. To be able to compare the results from the Axon and the Agilent scanner, we choose the same procedure for the images scanned on the Axon scanner, even though, the optimized segmentation for the strongest image could have been reused. We further believe that this allows us compare Spot and GenePix more fairly. For both Spot and GenePix the median spot pixel intensity was used as for the foreground signal. Background estimates were not considered in this analysis. No spot signals were discarded in analysis.

2.2.4 Data analysis

All further analysis was carried out using R [20, 13] and the aroma package (formerly known as the com.braju.sma package) [4] in which the methods presented here have been included.

2.3 Constrained model

Model (2) is not identifiable. Consider the case where the same array has been scanned at \( K \) different PMT settings. Let \( \mathbf{y}_{c,i} = (y_{c,i}^{(1)}, y_{c,i}^{(2)}, \ldots, y_{c,i}^{(K)}) \) be
the vector of the $K$ quantified signals for gene $i$ and channel $c$. In the noise-free case it follows from (2) that $y_c = \{y_{c,i}\}_{i=1}^{L}$ lies on the line $L(a_c, b_c)$ in $\mathbb{R}^K$, which has direction $b_c = (b_c^{(1)}, b_c^{(2)}, \ldots, b_c^{(K)})$ and goes through the point $a_c = (a_c^{(1)}, a_c^{(2)}, \ldots, a_c^{(K)})$. The $2K$ parameters of $a_c$ and $b_c$ are not identifiable, since $L$ has only $2K - 2$ degrees of freedom. In fact, any transformation $b_c \leftarrow k \cdot b_c$, and $a_c \leftarrow a_c + l \cdot b_c$, where $k$ and $l$ are scalars, will leave $L$ intact. In this paper, we make $a_c$ and $b_c$ identifiable by choosing $k$ and $l$ so that $b_c^{(1)} = 1$ and $a_c$ is the point on $L$ closest (in Euclidean norm) to the (diagonal) line $L' = \{e_i(1, \ldots, 1); e \in \mathbb{R}\}$. The choice of $a_c$ can be motivated by looking at the observed data. By a first inspection we observe that the bias $a_c^{(k)}$ in model (2) is not varying much when the PMT gain is changed. To demonstrate this, $(y_{R}^{(k)}, y_{L}^{(l)})$ have been plotted in a unique color together with the correspond-
ing fitted line for each of the six possible PMT pairs in Figure 1. The scatter plot to the left shows all observations and the one to the right zooms in on the low-intensity spots. First, the close fit of data to the lines is evidence that the scanner is linear in its dynamical range. Second, all lines go through approximately the same point, lets call it \((e_c, e_c)\), which suggests that there is a common PMT-independent bias \(e_c\) to all scans. More precisely, split the bias term into two parts, one dependent and one independent on the PMT gain according to

\[ a_c^{(k)} = d_c^{(k)} + e_c; \quad k = 1, 2, \ldots, K, \]  

and define \(d_c = (d_c^{(1)}, d_c^{(2)}, \ldots, d_c^{(K)}) \in \mathbb{R}^K\). Then, data indicates that \(\|d_c\| \approx 0\), where \(\|\cdot\|\) is the norm in, say, \(L_2\) (Euclidean distance). Let \(d = y - e(1, \ldots, 1)\) where \(y \in L\) and \(e \in \mathbb{R}\). The constraint that \(a_c\) is the point of \(L\) closest to \(L'\) can be formulated as

\[ d_c = \arg \min_d \|d\| \]  

where minimization is with respect to \(y\) and \(e\). Equivalently, this means that \(d_c\) is orthogonal to \(b_c\) and \((1, \ldots, 1)\). The above can be interpreted geometrically as follows. By definition, \(a_c\) is a point on the line \(L(a_c, b_c)\). Similarly, \(e_c = e_c(1, \ldots, 1)\) is a point on the diagonal line that goes through \((0, \ldots, 0)\) and \((1, \ldots, 1)\) in \(\mathbb{R}^K\), i.e. \(L'\). Now, minimizing \(d_c\) according to (4) is the same as finding the shortest distance between the line \(L\) and the diagonal line, which is also the distance between the two points \(a_c\) and \(e_c\). From the geometric interpretation it is also clear that, in order for the parameters to be uniquely identifiable, the line \(L\) must not be parallel to the diagonal line, that is, \(b_c^{(k)}\) must be different from \(b_c^{(1)}\) for some \(k\). A robust estimate of \(L\) was proposed in [5], using iteratively reweighted principal component analysis (IWPCA). This estimate of \(L\), together with the above parametrization of \(a_c\) and \(b_c\), give us estimates \(\hat{a}_c\) and \(\hat{b}_c\) of all \(2K - 2\) parameters of \(a_c\) and \(b_c\), as well as an estimate of \(\hat{e}_c\) of \(e_c\). Let us illustrate the parametrization and estimation procedure for \(K = 2\). Since two (non-parallel) lines will always intersect, the constraint (4) degenerates to the assumption that \(d_c = 0\), or equivalently, that \(a_c^{(1)} = a_c^{(2)} = e_c\). In the noise-free case the line \(L\) is described by

\[ y_{c,i}^{(2)} = \alpha_c + \beta_c y_{c,i}^{(1)}; \quad i = 1, 2, \ldots, I, \]  

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where \( \alpha_c = e_c(1 - \frac{b_c(2)}{b_c(1)}) \) and \( \beta_c = \frac{b_c(2)}{b_c(1)} \). By setting \( y_{c,i}^{(2)} = y_{c,i}^{(1)} \) in (5) and applying the constraint \( b_c(1) = 1 \), we get that \( a_c = (e_c, e_c) \) and \( b_c = (1, \beta_c) \) where \( e_c = \alpha_c/(1 - \beta_c) \). To further illustrate the stability of the PMT independence, the parameters \((e_c, \beta_c)\) have been estimated for each of the six PMT pairs independently based on data from array A scanned by the Axon scanner and quantified by GenePix. The various estimates for both channels are listed in Table 1. The average estimate of the bias across all PMT pairs in the red channel was \( \bar{e}_R = 18.0 \) (with standard deviation 1.12). For the green channel the average bias estimate was \( \bar{e}_G = 20.3 \) (with standard deviation 0.80). The small standard deviations confirm that \( d_c \) is indeed small.

<table>
<thead>
<tr>
<th>PMT pair</th>
<th>( \hat{e}_R )</th>
<th>( \hat{\beta}_R )</th>
<th>( \hat{e}_G )</th>
<th>( \hat{\beta}_G )</th>
</tr>
</thead>
<tbody>
<tr>
<td>(800,500)</td>
<td>18.9</td>
<td>32.4</td>
<td>20.7</td>
<td>32.5</td>
</tr>
<tr>
<td>(700,500)</td>
<td>19.0</td>
<td>13.0</td>
<td>20.9</td>
<td>12.8</td>
</tr>
<tr>
<td>(800,600)</td>
<td>17.3</td>
<td>7.1</td>
<td>19.8</td>
<td>7.9</td>
</tr>
<tr>
<td>(600,500)</td>
<td>19.0</td>
<td>4.5</td>
<td>20.8</td>
<td>4.1</td>
</tr>
<tr>
<td>(700,600)</td>
<td>17.6</td>
<td>2.9</td>
<td>20.9</td>
<td>3.1</td>
</tr>
<tr>
<td>(800,700)</td>
<td>16.3</td>
<td>2.5</td>
<td>18.9</td>
<td>2.6</td>
</tr>
<tr>
<td>mean</td>
<td>18.0 ± 1.12</td>
<td>20.3 ± 0.80</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Table 1: Red and green channel estimates of the bias and the slope for each PMT voltage pair together with the mean and the standard deviation of all estimates. The small standard deviation is evidence that the bias to a large extent is independent of the PMT settings. Data shown originates from array A scanned on Axon and analyzed with GenePix.

2.4 Calibration

Given estimates of \( a_c^{(k)} \) and \( b_c^{(k)} \), data can be calibrated using backward transformation. Let

\[
y_{c,i}^{(k)} = \frac{y_{c,i}^{(k)} - a_c^{(k)}}{b_c^{(k)}}, \tag{6}
\]

\[
\varepsilon_{c,i}^{(k)} = \frac{\varepsilon_{c,i}^{(k)}}{b_c^{(k)}} \tag{7}
\]
be the backward transformed observed signal and the rescaled error terms, respectively, for all $c, i$ and $k$. The affine model (2) can then be rewritten as

$$
\tilde{y}_{c,i}^{(k)} = x_{c,i} - \tilde{\varepsilon}_{c,i}^{(k)}, \ \forall c, i, k.
$$

(8)

Moreover, let

$$
\bar{\tilde{y}}_{c,i} = \frac{1}{K} \sum_{k=1}^{K} \tilde{y}_{c,i}^{(k)}
$$

(9)

be the average backward transformed signal for gene $i$ in channel $c$. Now, if $E[\varepsilon_{c,i}^{(k)}] = 0$, then $E[\tilde{\varepsilon}_{c,i}^{(k)}] = 0$ and hence

$$
E[\bar{\tilde{y}}_{c,i}] = x_{c,i}
$$

(10)

when all $a_{c}^{(k)}$ and $b_{c}^{(k)}$ are known. Thus, if (6) is applied with estimates of $a_{c}^{(k)}$ and $b_{c}^{(k)}$ that are consistent as $I \to \infty$, and the error terms have zero mean, the mean of the backward transformed signals will converge to $x_{c,i}$ as $I$ grows. Even though $E[\tilde{y}_{c,i}]$ is not observable, we can estimate it consistently by increasing the number of scans $K$. Inspection of the residuals of calibrated signals (not shown) indicates that the variance of the calibrated noise is independent of PMT setting, that is $V[\tilde{\varepsilon}_{c,i}] = \sigma_{c,i}^2$. Assuming independent noise terms, the variance of the sample mean is

$$
V[\bar{\tilde{y}}_{c,i}] = \frac{1}{K} \sigma_{c,i}^2,
$$

(11)

or equivalent, the standard deviation of the calibrated mean spot signal decrease as $1/\sqrt{K}$ where $K$ is the number of scans. In summary, we obtain consistent estimates (up to a multiplicative constant) of all $x_{c,i}$ with increasing $I$ and $K$.

Finally, signals that are saturated by the scanner have to be excluded before calculating the average. If the quantified signal for a spot happens to be saturated in all scans, then that spot is marked as saturated, which still may be informative when compared to other non-saturated signals.
3. Results

3.1 Parameter estimates

For every possible combination of array, scanner and signal quantification method (image analysis or raw pixel intensities), we estimated the parameters $\hat{a}_c$ (including $\hat{e}_c$ and $\hat{d}_c$) and $\hat{b}_c$ in model (2)-(4) for both channels according to the algorithm described in Section 2.3. To better understand the properties of the estimates, we use a bootstrap approach to obtain not only bias corrected estimates, but also their standard deviations. For GenePix and Spot quantified signals a bootstrap sample of size 100 was used. For the estimates based on the raw pixel intensities a different approach was taken. Because the number of pixels for one scan is about $10^7$ (per channel) and we had four scans our computer system limited us to estimate the model based on a subset of $10^6$ pixel intensities. This was done for 100 random subsets and the mean and standard deviation of the parameter estimates were calculated, much like the above bootstrap method.

The mean and the standard deviation (within parentheses) of $\hat{e}_c$ and $\|\hat{d}_c\|$ for all possible setups are listed in Table 2 and Table 3. The mean and standard deviation of $\hat{e}_c$, and the corresponding median and median absolute deviation (MAD), across all arrays are shown in Table 4 and Table 5, respectively.

3.1.1 Comparison of arrays

The bias estimates for all bootstrap replicates in Table 2 and Table 3 have been depicted as box plots in Figure 2. Considered that the signals are in range is of 0 to 65535, the bias estimates are very stable between different arrays. The bias estimates span 9.8 units ($0.15\%$) in the red channel and 7.8 units ($0.12\%$) in the green channel. Next we will study the differences in bias estimates between scanners, image analysis methods, and channels.

3.1.2 Comparison of scanners

For the two scanners, we found that the estimated biases based on signals obtained by the Agilent scanner are consequently higher than the estimates from the Axon scanner. The box plots of their differences in the common bias $e_c$ (for each bootstrap sample) between the Agilent and the Axon scanner in Figure 3 confirm this. See also Table 4 and Table 5. The significant difference could be an effect of scan
Figure 2: Estimated biases $e_c$ for each array and for all arrays together from the Agilent scanner (left column) and the Axon scanner (right column). The top four graphs are for the red channel and the bottom four are for the green channel. For each channel the top two are estimates based on signals quantified by GenePix and the bottom two on signals quantified by Spot. See also Table 2 and Table 3.
No apparent trend was found.

To test for scan order effects, we could not simply balance the experimental design by letting some arrays be scanned in the reverse order. Instead, to test for scan order trends, we scanned one array (H) first on Agilent, then on Axon and then again on Agilent. The two Agilent-Axon comparisons are labeled H-1 and H-2. No apparent trend was found.

### 3. Results

**Table 2:** Bootstrapped parameter estimates for the red channel with standard deviations for each combination of array, image method (or pixel intensities), and scanner.

<table>
<thead>
<tr>
<th>array</th>
<th>image method</th>
<th>Agilent</th>
<th>Axon</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>$\hat{e}_R$</td>
<td>$\hat{d}_R$</td>
</tr>
<tr>
<td>A</td>
<td>Spot</td>
<td>17.9 ± 0.253</td>
<td>1.14 ± 0.148</td>
</tr>
<tr>
<td>A</td>
<td>GenePix</td>
<td>44.9 ± 0.036</td>
<td>22.85 ± 0.043</td>
</tr>
<tr>
<td>B</td>
<td>Spot</td>
<td>19.2 ± 0.255</td>
<td>1.05 ± 0.165</td>
</tr>
<tr>
<td>B</td>
<td>GenePix</td>
<td>21.2 ± 0.225</td>
<td>1.83 ± 0.203</td>
</tr>
<tr>
<td>B</td>
<td>pixels</td>
<td>42.8 ± 0.049</td>
<td>20.71 ± 0.051</td>
</tr>
<tr>
<td>C</td>
<td>Spot</td>
<td>17.6 ± 0.366</td>
<td>2.00 ± 0.192</td>
</tr>
<tr>
<td>C</td>
<td>GenePix</td>
<td>19.8 ± 0.284</td>
<td>0.94 ± 0.209</td>
</tr>
<tr>
<td>C</td>
<td>pixels</td>
<td>21.0 ± 0.138</td>
<td>1.48 ± 0.079</td>
</tr>
<tr>
<td>D</td>
<td>Spot</td>
<td>18.2 ± 0.301</td>
<td>1.56 ± 0.103</td>
</tr>
<tr>
<td>D</td>
<td>GenePix</td>
<td>21.1 ± 0.274</td>
<td>0.95 ± 0.139</td>
</tr>
<tr>
<td>D</td>
<td>pixels</td>
<td>25.0 ± 0.522</td>
<td>4.10 ± 0.358</td>
</tr>
<tr>
<td>E</td>
<td>Spot</td>
<td>18.0 ± 0.428</td>
<td>1.15 ± 0.109</td>
</tr>
<tr>
<td>E</td>
<td>GenePix</td>
<td>22.1 ± 0.268</td>
<td>1.77 ± 0.223</td>
</tr>
<tr>
<td>E</td>
<td>pixels</td>
<td>24.7 ± 0.144</td>
<td>5.51 ± 0.169</td>
</tr>
<tr>
<td>F</td>
<td>Spot</td>
<td>19.9 ± 0.423</td>
<td>0.48 ± 0.163</td>
</tr>
<tr>
<td>F</td>
<td>GenePix</td>
<td>23.8 ± 0.316</td>
<td>1.47 ± 0.258</td>
</tr>
<tr>
<td>F</td>
<td>pixels</td>
<td>24.2 ± 0.131</td>
<td>1.37 ± 0.101</td>
</tr>
<tr>
<td>G</td>
<td>Spot</td>
<td>16.0 ± 0.300</td>
<td>2.30 ± 0.166</td>
</tr>
<tr>
<td>G</td>
<td>GenePix</td>
<td>18.5 ± 0.208</td>
<td>1.88 ± 0.198</td>
</tr>
<tr>
<td>G</td>
<td>pixels</td>
<td>19.5 ± 0.090</td>
<td>1.48 ± 0.080</td>
</tr>
<tr>
<td>H-1</td>
<td>Spot</td>
<td>20.4 ± 0.161</td>
<td>2.12 ± 0.073</td>
</tr>
<tr>
<td>H-1</td>
<td>GenePix</td>
<td>22.1 ± 0.124</td>
<td>2.41 ± 0.097</td>
</tr>
<tr>
<td>H-1</td>
<td>pixels</td>
<td>44.9 ± 0.513</td>
<td>17.68 ± 0.528</td>
</tr>
<tr>
<td>H-2</td>
<td>Spot</td>
<td>20.1 ± 0.141</td>
<td>0.33 ± 0.040</td>
</tr>
<tr>
<td>H-2</td>
<td>GenePix</td>
<td>21.8 ± 0.087</td>
<td>0.22 ± 0.084</td>
</tr>
<tr>
<td>H-2</td>
<td>pixels</td>
<td>21.8 ± 0.047</td>
<td>0.26 ± 0.042</td>
</tr>
</tbody>
</table>
Applications use different spot segmentation algorithms [22, 3]. Because the

differences in estimates of the common bias (for each bootstrap sample) between

| array | image method | Agilent \( \hat{e}_G \) | || Agilent \( ||\hat{d}_G|| \) | Axon \( \hat{e}_G \) | || Axon \( ||\hat{d}_G|| \) |
|-------|--------------|----------------|----------------|----------------|----------------|----------------|
| A     | Spot         | 20.2 ± 0.106   | 0.51 ± 0.058   | 16.6 ± 0.165   | 6.31 ± 0.302   |
| A     | GenePix      | 21.8 ± 0.068   | 0.20 ± 0.045   | 20.4 ± 0.123   | 0.88 ± 0.242   |
| A     | pixels       | 33.6 ± 0.050   | 10.52 ± 0.048  | 22.8 ± 0.048   | 1.57 ± 0.093   |
| B     | Spot         | 19.6 ± 0.094   | 0.90 ± 0.055   | 17.2 ± 0.127   | 2.20 ± 0.231   |
| B     | GenePix      | 21.3 ± 0.046   | 0.94 ± 0.086   | 20.1 ± 0.152   | 1.20 ± 0.326   |
| B     | pixels       | 35.7 ± 0.153   | 12.85 ± 0.146  | 21.9 ± 0.041   | 2.04 ± 0.087   |
| C     | Spot         | 19.8 ± 0.050   | 1.21 ± 0.073   | 17.0 ± 0.127   | 2.75 ± 0.254   |
| C     | GenePix      | 20.8 ± 0.039   | 0.99 ± 0.086   | 19.4 ± 0.100   | 1.11 ± 0.218   |
| C     | pixels       | 21.3 ± 0.024   | 0.78 ± 0.035   | 22.0 ± 0.040   | 2.39 ± 0.082   |
| D     | Spot         | 19.4 ± 0.097   | 1.49 ± 0.073   | 17.0 ± 0.194   | 1.24 ± 0.376   |
| D     | GenePix      | 20.9 ± 0.059   | 0.80 ± 0.095   | 20.3 ± 0.126   | 2.31 ± 0.286   |
| D     | pixels       | 21.6 ± 0.105   | 0.21 ± 0.124   | 22.1 ± 0.046   | 3.10 ± 0.092   |
| E     | Spot         | 18.2 ± 0.121   | 3.29 ± 0.129   | 16.5 ± 0.132   | 2.88 ± 0.240   |
| E     | GenePix      | 20.2 ± 0.081   | 1.92 ± 0.128   | 19.2 ± 0.137   | 0.74 ± 0.264   |
| E     | pixels       | 21.2 ± 0.017   | 0.13 ± 0.046   | 21.0 ± 0.032   | 0.94 ± 0.058   |
| F     | Spot         | 21.4 ± 0.125   | 0.58 ± 0.103   | 16.5 ± 0.181   | 4.79 ± 0.327   |
| F     | GenePix      | 23.7 ± 0.084   | 1.32 ± 0.116   | 20.2 ± 0.132   | 0.34 ± 0.199   |
| F     | pixels       | 24.1 ± 0.019   | 1.57 ± 0.026   | 22.4 ± 0.039   | 1.87 ± 0.071   |
| G     | Spot         | 18.9 ± 0.088   | 1.62 ± 0.088   | 16.9 ± 0.149   | 3.72 ± 0.238   |
| G     | GenePix      | 20.4 ± 0.049   | 1.10 ± 0.088   | 20.0 ± 0.164   | 0.84 ± 0.262   |
| G     | pixels       | 21.8 ± 0.017   | 0.75 ± 0.024   | 22.2 ± 0.045   | 2.60 ± 0.085   |
| H-1   | Spot         | 21.3 ± 0.134   | 3.25 ± 0.080   | 18.5 ± 0.090   | 2.00 ± 0.170   |
| H-1   | GenePix      | 22.6 ± 0.100   | 3.47 ± 0.107   | 20.3 ± 0.043   | 0.63 ± 0.100   |
| H-1   | pixels       | 42.8 ± 1.417   | 16.68 ± 1.843  | 21.3 ± 0.033   | 1.82 ± 0.070   |
| H-2   | Spot         | 19.6 ± 0.229   | 0.16 ± 0.077   | N/A            | N/A            |
| H-2   | GenePix      | 21.7 ± 0.104   | 0.55 ± 0.076   | N/A            | N/A            |
| H-2   | pixels       | 21.5 ± 0.086   | 0.29 ± 0.021   | N/A            | N/A            |

Table 3: Bootstrapped parameter estimates for the green channel with standard deviations for each combination of array, image method (or pixel intensities) and scanner.

3.1.3 Comparison of image analysis methods

Furthermore, estimates of the common bias \( \hat{e}_c \) based on GenePix quantified signals are consistently greater than the corresponding ones based on Spot signals, cf. Tables 2 and 3. See also Tables 4 and 5. The box plots in Figure 4 show differences in estimates of the common bias (for each bootstrap sample) between GenePix and Spot. The difference may be explained by the fact that the two applications use different spot segmentation algorithms [22, 3]. Because the con-
3. Results

Table 4: Mean and standard deviation of the bias estimates of all arrays and for each signal quantification method.

<table>
<thead>
<tr>
<th>image method</th>
<th>$\bar{e}_{R}$</th>
<th>$\bar{e}_{G}$</th>
</tr>
</thead>
<tbody>
<tr>
<td>Spot</td>
<td>18.4 ± 1.49</td>
<td>19.7 ± 0.854</td>
</tr>
<tr>
<td>GenePix</td>
<td>21.3 ± 1.27</td>
<td>21.3 ± 0.822</td>
</tr>
<tr>
<td>pixels</td>
<td>24.6 ± 5.38</td>
<td>21.6 ± 0.817</td>
</tr>
</tbody>
</table>

Table 5: Median and MAD (median absolute deviation) of the bias estimates of all arrays and for each signal quantification method.

<table>
<thead>
<tr>
<th>image method</th>
<th>$\bar{e}_{R}$</th>
<th>$\bar{e}_{G}$</th>
</tr>
</thead>
<tbody>
<tr>
<td>Spot</td>
<td>19.8 ± 0.792</td>
<td>17.0 ± 0.633</td>
</tr>
<tr>
<td>GenePix</td>
<td>21.5 ± 1.065</td>
<td>20.0 ± 0.433</td>
</tr>
<tr>
<td>pixels</td>
<td>27.0 ± 7.76</td>
<td>22.0 ± 5.37</td>
</tr>
</tbody>
</table>

concentration of fluorophores is not homogeneous across a spot, the result is that the distribution of pixel intensities will vary with segmentation method. This effect can be more profound for spots with strong donut effects. Robust estimates such as the median pixel value will to some extend protect against this, but not completely. It has been suggested [7] that the median of (pixel) ratios is a better estimate of the ratio of hybridized cDNA than the ratio of median (pixels). However, the former requires that the images are perfectly aligned with respect to shift, rotation, shear and so on. Also, it applies exclusively to two sample comparisons. Because of this, we do not believe that pixel-ratio signals are useful in practice.

3.1.4 Pixel-based estimates

To better understand the underlying reasons for the observed channel biases, the proposed affine model was also applied to pixel intensities (instead of spot signals). The estimated biases for the red and the green channel for different arrays using multidimensional IWPCA based on pixel values are shown in Tables 2 and 3. Except for the green channel in the second scan round of array H, the pixel-based estimates are consistently higher than the estimates based on GenePix and the Spot foreground signals. As noted above, pixel-based estimates are very sensitive to image distortions. This is especially a concern for the Agilent scanner since it
Figure 3: Differences in the common biases \( e_c \) between the Agilent and the Axon scanners for each separate array and for all arrays together. For both the red (top) and the green channel (bottom), the biases estimated from GenePix quantified signals (left) are significantly greater than the biases estimated from Spot quantified signals (right). The exception is a small fraction of the bootstrap estimates from array F.

reloads the arrays between subsequent scans. To check this, we did a test where a test person with experience in microarray analysis was asked to subjectively rank how badly aligned the four images in the red channel with different PMT gains from the Agilent scanner were for each of the (unlabeled) nine arrays. The person rated the images from arrays A, B, D, and H-1 to be “extremely” misaligned. The images from array E were considered to be “quite” misaligned, and the images from array C to be “slightly” misaligned. For the rest of the arrays the images were considered to be aligned (less than one pixel off). This blind-folded test is perfectly in line with the discrepancies between pixel estimates and the others listed in Tables 2, which confirms our hypothesis. Another disadvantages of estimating the biases based on pixel intensities is that it is extremely memory and time consuming. For instance, estimating the parameters based on \( 10^6 \) pixels
3. Results

Figure 4: Differences in the common biases $e_c$ between the GenePix and the Spot image analysis method for each separate array and for all arrays together. For both the red (top) and the green channel (bottom), the biases estimated from Agilent signals (left) are significantly greater than the biases estimated from Axon signals (right).

took approximately 50 times longer than to estimate them based on 55488 quantified signals, which is because a greater number of data points are fitted, but also because the pixel intensities had to be reloaded and resampled before each estimation round due to memory constraints.

3.1.5 Comparison of channels

As Figure 5 shows, the common bias $e_c$ is greater in the green channel than the red channel, especially for GenePix quantified signals, when estimated based on data from the Axon scanner. For the Agilent scanner this trend is less clear, though the Spot quantified signals clearly seem to give higher bias in the green that the red channel. Furthermore, the biases in the red and the green channels seem to be stable between arrays, which give further evidence to our hypothesis that the bias originates from the scanner (and/or the image analysis methods).
Figure 5: Differences in the common biases $e_c$ between the red and the green channel for each separate array and for all arrays together. At the top is data from the Agilent scanner and at the bottom from the Axon scanner. Data in the left column by GenePix and data in the right column was quantified by Spot. For data from the Axon scanner the common bias is greater in the green channel than the red channel, especially for GenePix quantified signals. For the Agilent scanner this trend is less clear although the Spot quantified signals clearly seem to have a higher bias in the green than the red channel.

3.1.6 Deviation in bias estimates between PMT gains

In Figure 6 the distribution of the “bias residuals” $d^{(k)}_{ic}$ are depicted for different scans $k$ and channels $c$, for each separate array, but also all arrays together, and for both scanners and both image analysis methods. Each array is presented in increasing PMT order. The most apparent effect is that the estimates based on signals from the Axon scanner and especially those quantified by the Spot software are greater than for the others, cf. Table 2 and Table 3. Why this is so we do not know. For some arrays the estimates from the red and the green channels are strongly correlated, but it is not clear to us when this occurs. Although not in general, for some combinations of scanner and image analysis method, there is a
3. Results

The trend in the PMT order (or possibly scan order). Again, we do not know why.

To summarize, we have by means of exploratory data analysis (not shown) tried to understand what sometimes looks like patterns in the $d_c^{(k)}$'s, but we found no apparent relationships. However, systematic effects might indicate that $d_c^{(k)}$ could be modeled further.

3.2 Calibration

When data was calibrated according to the backward transformation in (6)-(9), estimates (up to a scale factor) of all $x_{c,i}$'s were obtained. Since we do not know the true values, we cannot verify the estimates directly. However, to some extent, we may do it indirectly by looking for remaining systematic effects in the log-ratios, but also by comparing the empirical densities of the calibrated scans. For a detailed study on various systematic effects an affine transformation introduces in the log-ratios, see [5]. For instance, here the amount of intensity-dependent curvature in the log-ratios is related to the bias and the relative scale factor via the product $e_c(1 - b_c^{(k)}/b_c^{(l)})$ assuming $\|d_c\| = 0$. To demonstrate this, we have for various PMT pairs compared the within-channel log-ratios and log-intensities

$$M_{c,i}^{(k,l)} = \log_2(y_{c,i}^{(k)}/y_{c,i}^{(l)})$$

$$A_{c,i}^{(k,l)} = \frac{1}{2} \log_2(y_{c,i}^{(k)}/y_{c,i}^{(l)})$$

respectively, with the corresponding ones for the backward transformed data, which we denote by $\tilde{M}_{c,i}^{(k,l)}$ and $\tilde{A}_{c,i}^{(k,l)}$. The log-ratios versus the log-intensities for the new signals of all six PMT pairs are shown in the left scatter plot in Figure 7. The corresponding plot for the backward transformed signals is shown to the very right. For each of the six data clouds, the curvature, but also the overall bias, in the log-ratios is removed. To further underline the effect that a channel-specific bias has, we have calculated the log-ratios for the bias-subtracted signals (no rescaling), which makes model (2) become linear. As seen in the middle scatter plot, the curvature introduced by the bias and the logarithm is removed. The overall bias in the log-ratios that remains is $\log_2(b_c^{(k)}/b_c^{(l)})$ and is removed when the signals are rescaled. Note that it is not correct to shift only the log-ratios toward zero, because then the log-intensities will be incorrect. The various $M$ versus $A$ scatter plots become very similar and so do the four empirical density functions of the signals, as seen in Figure 8. The small bumps at high intensities are due to the
Figure 6: Estimated bias residuals $d_c^{(k)}$ for each array and for all arrays together from the Agilent scanner (left column) and the Axon scanner (right column). For each array the distribution of the four of $d_c^{(k)}$ are shown in increasing PMT order. For details on PMT settings, see Section 2.2.2. The top four graphs are for the red channel and the bottom four are for the green channel. For each channel the top two are estimates based on signals quantified by GenePix and the bottom two on signals quantified by Spot.
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Figure 7: The affine transformation gives curvature in the M versus A plots, which is corrected for by the affine normalization method. The three scatter plots show the within-channel log-ratio versus the log-intensity for each of the six PMT pairs based on the same data as in Figure 1. Left: Observed signal for different PMT pairs. For each pair the estimated \((A_0, M_0) = \left(\frac{1}{2} \log_2(e_G^{(k)}) \cdot \log_2(e_G^{(l)})\right)\) has been marked with a circle, cf. Figure 1. Middle: Bias corrected signals. Right: Bias and scale calibrated signals. The range of the M axis is twice the range of the A axis so that (12)-(13) appear as a rotation in the plots.

saturated signals, which can also seen in for instance Figure 7. These bumps are not visible if the saturated signals are removed.

Figure 8: The affine normalization method makes the signal densities much more similar. Left: Density plots of the logarithm of the raw signals for each of the fours scans. Middle: Bias corrected signals. Right: Bias and scale calibrated signals. Data and colors are the same as in Figure 1.

3.2.1 Extended dynamical range

For the Agilent scanner the effective scale parameters \(b_c^{(k)}/b_c^{(1)}\) were estimated to be in the order of approximately 1 : 3.5. For the Axon scanner they were in the
order of approximately 1 : 40, cf. Table 1. Thus, the calibration method extends the effective dynamical range by a factor of 3.5 for data scanned on Agilent and a factor of 40 for data scanned on Axon. In both cases linearity was preserved.

4 Discussion

4.1 Sources of the bias

Because any bias introduced before the PMT detector would be registered and amplified differently by the PMT at different gains, we believe that the observed bias is due to the scanner and most likely its detector parts such as the analog-to-digital (A/D) converter after the PMT, but possibly also due to the image analysis method. The observed differences between the channels can be explained by the fact that there is one PMT and one A/D converter per channel, which may have slightly different properties. Although there are differences in bias between the two scanners, they are still of the same order, which we find remarkable. Another lab, which uses different arrays, reported biases also around 15-20 (personal communication). One possible reason for this is that the scanners could consist of similar parts although they are of different brands.

4.2 Other estimates

To rule out the obvious situation where all pixel intensities are biased, we compared the above estimates with the minimum pixel intensity. For example, for array A (scanned on Axon and analyzed with GenePix Pro), the minimum pixel intensities in the red channel were 9, 0, 8, and 9 for PMT 500, 600, 700 and 800 volts, respectively. In the green channel the minimum pixel intensity is 0 for all scans. It is not useful to use the minimum spot signals, ˆa(κ) c = min y(κ) c,i, either. For example, for the above scan the average minimum signal across all scans in the red channel is 19.8 (median 19.5, std. dev. 0.96), but in the green channel it is 34.8 (median 28.0, std. dev. 19.6). The estimates for the green channels have a large variation compared to the estimates based on the affine model, cf. Table 3.

4.3 On background subtraction

If the scanner is the main source for the observed bias, then the background estimates should be affected by this bias as well and subtracting the background from
4. Discussion

the foreground estimates will therefore not only correct for physical background noise from the array itself, but also for the scanner bias. The strong intensity-dependent effects of the log-ratios that are due to the bias are much less apparent if we apply background subtraction (not shown), giving more evidence to our hypothesis that the observed systematic effects originate from the scanner. Thus doing background correction might correct for the bias, but it will also introduce more noise at any given intensity (not depicted). Also, with the data set in hand background subtraction results in 4050 (7.3%), 6237 (11%), 7015 (13%) and 7349 (13%) negative values (in either channel), respectively whereas the bias subtraction results in no negative values. If we assume that the noise is additive such that the background is added to the foreground signals, then for probes with few or no fluorescent molecules the true foreground signal should be close or identical to the true background signal. As both are estimates, approximately half of the foreground signals for non-signal spots are less or equal to the corresponding background signals. Thus, about half of such spots results in negative signals. However, the different numbers of negative signals for different PMT voltages suggest that this cannot be the full explanation. One reason may be that the background estimates are likely to be biased [3]. An error model that incorporates different noise sources, but also different scan parameters, might give some answers to this. Some promising models in this context have already been suggested [17, 12, 8], but also other models based on empirical Bayes methods [15].

Another way to put it is that the background estimate is local and based on individual spots/pixels whereas the bias estimate is global, that is, there is one estimate for the whole array (although one could also imagine for instance spatially dependent estimates). Therefore, the background subtracted intensity estimates are noisier, resulting in more negative estimates for low intensity spots. Because remaining systematic effects due to the image analysis methods still have to be corrected for we find that the spot signals should be calibrated for biases.

4.4 Photo bleaching

Not considered in the above analysis is photo bleaching. We estimated the red dye (Cy5) to bleach about 2% and the green dye (Cy3) about 1% in a typical microarray experiment (not shown). Because the amount of bleaching is not too large, but also because it is a very complex phenomenon, we decided to not try to incorporate it in the above model. Some of the systematic variation seen in the bias estimates for the different PMT settings (Section 3.1.6) may be due to
bleaching.

4.5 Signal density normalization

As shown in Section 3.2, the empirical distributions of signals match each other remarkably well after calibration. It is interesting to compare this method with the quantile normalization methods proposed by [6, 24]. The latter is based on the “statistical” assumption that the signals in all channels (scans) should be equal whereas as the former is based more on a “physical” assumption that the signals should be linear in the dynamical range. For a further discussion on this see [5].

4.6 Related work

Another method that combines multiple scans is the masliner (Microarray Spot LINEar Regression) algorithm [9]. It works by combining one low-PMT scan and one high-PMT scan into a new virtual scan. If a signal in the high-PMT scan is within a specified linear range its value is used, otherwise the corresponding signal from the low-intensity range is used after being transformed affinely to fit the high-PMT scan. To combine three or more scans, the new virtual scan can be combined with another PMT scan and so on. The result is that the effective dynamical range is extended. However, there are several unnecessary drawbacks. First, although several observations of the same spot concentration exist, which all may be within the dynamical range of the scanner, only one observation is used. Statistically, the average of all (calibrated) signals would be a more precise (less noisy) estimate. Second, since the scans are combined pairwise, the estimate of the affine relationship between the scans is less robust. Third, although a sensitivity discussion is carried out in the supplementary materials, masliner fits the affine models in a non-robust fashion (in $L_2$). Also, classical linear regression is used, which assumes no error in the explanatory variable. Since masliner makes the signals from different PMT settings proportional to each other, it will indeed remove for instance curvature in within-channel $M$ versus $A$ scatter plots. However, masliner does not model the possibility of a PMT-independent bias and will therefore not correct for it. We believe this is the reason why the authors observe a “curvilinear effect” [9, supplementary material]. For these reasons, we believe that the robust multiscan calibration method presented in this paper is superior to the masliner algorithm and should be used instead.
4.7 Incremental robust estimates

It turned out to be infeasible to estimate the model parameters based on all pixel intensities, which limited us to use only on a 10% subset of all data points. As argued above, pixel-based estimates are not reliable and therefore not of interest. However, for spot-based estimates the same limitations may apply as larger data sets are made available. We wish to overcome such memory constraints. For this reason, we investigate the possibility to use (approximative) incremental re-weighted PCA methods [19, 16].

5 Conclusions

By scanning the same microarray at various PMT settings, we have shown that there exists a bias in the measurement of the concentration of fluorescent molecules in the spots on the microarray. Our analysis indicates that this bias is mainly due to the scanner, but also due to the image analysis methods. Moreover, by using a constrained affine model for the relationship between the obtained fluorescent intensities and fluorophore concentrations in the spots, we have been able to estimate the aforementioned bias. With estimates of the bias and scale parameters in each channel, back transformation gave estimate of the amount (up to a scale factor) of photons from each spot that enters the PMT, that is $x_{c,i} = h_{c}^{-1}(y_{c,i}); \forall c, i$. Although not all photons originate solely from fluorophores in the target DNA, this is still a far better estimate of the amount of hybridized target DNA in each spot than the corresponding signal quantified by the scanner and the image analysis alone.

Before calibration data show a strong intensity-dependent effect in the log-ratios whereas after calibration there is no apparent intensity-dependent trend. Furthermore, the distributions of signals from subsequent PMT scans are almost identical after calibration. In addition to this, with multiple scans, the signal-to-noise ratio is increased. Finally, scanning at both low and high PMT settings extends the dynamical range of data, which gives higher resolution at low intensities without having to pay the price of saturated signals.

The proposed method can be applied to other microarray technologies such as single-channel oligonucleotide arrays or nylon arrays, and possibly to other gene-expression technologies such as quantitative real-time polymerase chain
reaction (QRT-PCR). If an affine relationship is not observed, more sophisticated methods such as the promising quantile normalization methods suggested by [6, 21] may be used. More research is needed.

To conclude, we suggest that hybridized microarrays are scanned at two (preferably more) PMT gain levels to identify channel dependent bias terms. Knowing the exact PMT settings is not important, but it can be shown that the larger the differences are the more precise the estimates will be. We recommend that the scans are done in decreasing PMT-gain order (although we did not do so here). Given the estimates, data can then be calibrated easily.

For two-channel microarrays, after calibrating each channel separately, a similar strategy can be applied once more to bring differently labeled channels to the same scale as suggest in [5]. This would rely on the assumption that the amounts of hybridized DNA in all channels are approximately equal for the majority of the spots, which in turn is based on the commonly used assumption that most genes are non-differentially expressed. This also applies to normalization between arrays.

All necessary methods are made available in a free R package named aroma [4]. A typical usage is `calibrateMultiscan(rg)` where `rg` is the object containing the red and green signals. In addition, we are currently implementing the methods as a plug-in module for the BASE system [18].

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References


Paper F
aroma - An R Object-oriented Microarray Analysis environment

Henrik Bengtsson

Mathematical Statistics at the Centre for Mathematical Sciences, Lund University, Box 118, SE-221 00 Lund, Sweden. Email: hb@maths.lth.se.

ABSTRACT

We have implemented a self-contained package for DNA microarray analysis in R. The package is named aroma and is formerly known as com.braju.sma. The purpose of the package is to provide end-users with easy access to the latest statistical tools and methods, to provide other statisticians with a structured platform to develop and test new analysis methods, and to make our methods available to an audience as broad as possible. In turn, this gives us most valuable feedback for improving the package and its algorithms. The package provides methods for low-level analysis and pre-processing of spotted microarray data with two or more channels, although the recent methods are applicable to single-channel microarrays as well. Many of the commonly known and widely used normalization methods have been implemented as well as our own normalization and calibration methods in both early and mature versions. A few high-level analysis methods do exist. Import and export of microarray data is supported for most free and commercial file formats. Support for new ones can easily be added. Connectivity to other microarray analysis platforms based on relational databases is planned. For exploratory data analysis, the package provides a large set of methods for visualization of data with automatic graphical annotations. Textual and symbolic annotation of graphs is straightforward. Generated graphics can be exported for on-screen presentations and high-resolution printed publications. Report generators for simultaneously creating documents in plain text, HTML, and LaTeX are available, making it easy to create batch analysis scripts. Detailed documentation with code examples and references to the literature are included. In order to provide time and memory efficient algorithms, an object-oriented design and implementation utilizing reference variables has been used. In addition, to minimize the risk for programming mistakes, a consistent application protocol
interface (API) was developed by enforcing an R coding convention. This will make it possible for a new user to quickly become familiar with the structure of classes and methods enhancing overall productivity. To further minimize the risk for programming errors, method arguments are validated as far as possible and explanatory error messages are used. Priority is also on backward compatibility with previous versions, compatibility with other packages, and potential future packages. The package does not provide a graphical user interface (GUI) per se, but the object-oriented API and the fact that reference variables are used make it easy for third-party developers to implement a GUI for, say, everyday high-throughput normalization. The package is available online as open source written in 100% R for maximum accessibility on all platforms. Installation is straightforward.

Keywords: microarray analysis; single-, two-, and multiple-channel microarrays; low-level analysis; pre-processing of data; visualization; calibration; normalization; data import and export; object-oriented programming; reference variables; coding conventions; data hiding; encapsulation; virtual fields.

1 Introduction

Aroma is a package for low-level analysis (calibration and normalization) of two- or multiple-channel microarray data, but part of it may as well be applied to single-channel microarray data. The package is implemented in pure R [13]. This means that it runs out of the box and works identically on operating systems such as Windows, Mac OS X, Sun Solaris, and any Linux system.

It is not our intention of this paper to give a complete description of the aroma package, but to give an general overview and to provide enough details to become familiar with the structure of the package and its methods. Hopefully, the reader will be tempted enough to install the package and start use it in his or her own microarray analysis.

The outline of the paper is as follows. In Section 2, we give a brief introduction to object-oriented programming and reference variables in general, and emphasize the fact that the aroma packages pass variables by reference and not by value (or
2. Brief about object orientation and reference variables

For the novel user we give a very brief introduction to the object-oriented design used by us and reference variables as these two concepts are fundamental to the aroma package and might differ slightly from what is otherwise common in R. For a general introduction to R, see [15].

2.1 Object orientation

In object-oriented programming two concepts are of major importance, namely classes and objects. Simply speaking, a class describes and defines a structured data type, for instance, a data table containing signals and other spot attributes returned by the Axon GenePix software. An object is then an instance of a class, meaning that it contains the actual data on which analysis is to be done. Several instances can exist of the same class. A simple rule for understanding this is that a class exists as source code at all time whereas an object exists while running the analysis. Moreover, different classes (data structures) for different purposes may be defined. A more real-world analogy is on cooking. In a cookbook there are recipes. These recipes describe and define certain types of dishes and can be thought of as classes. For any given recipe one or several meals (of identical type)

copy), as is the default in R. Section 3 explains how microarray signals are stored and queried. After these introductory sections, a walk-through of the aroma package is provided. After describing how R and the package is started, it is described how typical microarray data generated by common academic and commercial microarray image analysis applications is imported and how foreground and background signals are extracted from such. A few of the methods for generating graphical representation of microarray data are then presented. In the following subsection, calibration and normalization is covered. The walk-through is ended by a short description on how normalized data is written to file. In the section 5, some additional useful features are described. Report generators are introduced, the help and documentation of the package is described, and compatibility with other packages and other scientific applications is discussed. The philosophy of the design and implementation of the package is given in Section 6. This part is recommended to anyone who wish to extend aroma with new classes or methods, but may be skipped by anyone not familiar with software development. Installation instructions are given in 7.
can be prepared. These can be thought of as the objects, that is, they are instances of the recipe.

Different data structures are often more or less related to each other. For instance, a class for Axon GenePix microarray data is related to a class for QuantArray microarray data by the fact that they both represent quantified signals for microarray data. In contrast, these two classes are not related to a class describing, say, the heartbeat of a mouse. Let's denote the two classes by `GenePixData` and `QuantArrayData`, respectively. A common superclass of these two classes would then be `MicroarrayData`, describing the fact that the two classes share similar properties (from the superclass). For instance, they both contain foreground and background estimates, but in addition have properties that are unique to each of them. We say that the `GenePixData` class extends (the properties) or inherits from the `MicroarrayData` class. Furthermore, we say that `GenePixData` is a subclass of the `MicroarrayData` class.

To continue our analogy to cooking, the recipes in a cookbook are often categorized into sections such as starters, main dishes, and desserts. These categories may be thought of as abstract (super-)classes of the actual dishes. Simply, you cannot ask someone to make just a dessert (the superclass), but you have to be more specific and say which type of dessert (the class) in order for someone to get the exact ingredients, make the dessert, and bring it to you (the object). In our microarray example, the class `MicroarrayData` is abstract, because it is “not specific enough”. The reason for having a hierarchy of classes in this way is to make it simpler for both the programmer and the user to become familiar with, and navigate between, all the data structures and their properties. For further details on object-oriented programming in R, see [11, 12, 4].

2.2 Reference variables

The language in R is a so-called functional language, which means that all methods (functions) take some input objects, performs calculations and operations based on these and returns a new object. This imitates the concept of a function in mathematics, which is the basic language for all scientific analysis. The main advantage is that functions cannot modify an object (just return a new one) leaving no side effects or surprises to the user. This has shown to be very useful. The disadvantage is that it is less memory efficient since often two
copies of the same data structure will live in the memory at the same time. This is of huge concern in microarray analysis because of the large amount of data processed. For instance, consider an *RGData* object containing red and green foreground signals for thousands of spots. Internally, R will then, for a function that subtracts background signals from these signals, first create a copy of the object containing all foreground signals and then subtract the background signals on this new object, which is then returned. In the aroma package we have chosen another strategy; we use *reference variables* [4]. The difference is basically that a function, or a *method* to differentiate it from the original meaning of a function, can change the values of any object passed to it. We say that objects are *passed by reference*, contrary to being *passed by value or copy*.

There are other advantages of using pass by reference than just being memory efficient. Because a method can *change the state* of an object, it can store information that succeeding analysis methods use. For instance, the user may specify that all signals for a certain set of genes in a specific data set (object) should be highlighted in red. This “request” will be stored in the object and recognized by all future plot methods that will highlight the data points without the user having to specify which data points to highlight in every call\(^\text{14}\). See ?Filter for more details. The *highlight()* method described below is another example that makes use of reference variables.

We ask the reader to keep in mind that aroma uses reference variables.

### 3 Representation of data

Microarray data in aroma is represented by objects of classes that are all subclasses of the *abstract* class *MicroarrayData*. Any such class contains probe (spot) signals from one or several microarrays. The signals are stored in a field that is a \(I \times K\) matrix, where \(I\) is the number of probes and \(K\) the number of arrays. Thus, the signals in row \(i\) are the signals for probe \(i\) on all \(k = 1, \ldots, K\) arrays, and the signals in column \(k\) are the signals from array \(k\) and all probes \(i = 1, \ldots, I\). If a *MicroarrayData* object\(^\text{15}\) contains signals from more than one channel, each

\(^{14}\)This can be done with functional programming too, but is much less memory efficient.

\(^{15}\)When we refer to a “*MicroarrayData* object” we actually mean an object of a class that extends the *MicroarrayData* class.
channel is stored in a separate field. For instance, the \textit{RGData} class is a basic class that represents two-channel microarray data with two matrix fields \( R \) (red) and \( G \) (green) of equal dimensions. Consider an object \( \text{rg} \) of class \textit{RGData}. Then the signal in the red channel for spot \( 4, 5, 14 \ldots 20 \) on the first arrays is obtained by

\[
\text{rg}\$\text{R}[\text{c}(4,5,14:20),1]
\]

and the same spots on all other arrays by

\[
\text{rg}\$\text{R}[\text{c}(4,5,14:20),-1]
\]

Note that field \( \text{R} \) of object \( \text{rg} \) is accessed by the \$-operator similar to how elements in lists are accessed. Another example is to calculate the log-ratios and log-intensities [20]

\[
M = \log_2 \frac{R}{G} \\
A = \frac{1}{2} \log_2 (RG)
\]

for all probes and arrays. \( M \) is for “minus” and \( A \) is for “add”, because the ratios and products become subtraction and addition under the logarithm. In \( R \) this can be done by

\[
M \leftarrow \log(\text{rg}\$\text{R}/\text{rg}\$\text{G}, \text{base}=2) \\
M \leftarrow \log(\text{rg}\$\text{R}\ast\text{rg}\$\text{G}, \text{base}=2)/2
\]

which will create a new \( I \times K \) matrix. Indeed, since these transforms are so common, it is not necessary to do the above every time. For any \textit{MicroarrayData} class for two-channel data, the log-ratios and log-intensities can be obtained by \text{rg}\$\text{M} and \text{rg}\$\text{A}, respectively. For instance,

\[
\text{rg}\$\text{M}[\text{c}(4,5,14:20),-1] \\
\text{rg}\$\text{A}[\text{c}(4,5,14:20),-1]
\]

These fields are in fact not stored in the object, but are \textit{calculated on-demand} and do therefore not take up valuable memory. This is unique to aroma. For more details, see Section 6.4 on virtual fields. Continuing, to assign new values to some probes do

\[
\text{rg}\$\text{R}[14:20,1:2] \leftarrow 0
\]
3. Representation of data

which will zero the signals in the red channel for probe $14 - 20$ on array one and
two. The MAData class is complementary to the RGData class, in the meaning
that it stores the log-ratios and the log-intensities, and calculates the red and the
green signals on-demand by the inverse transformations

$$
R = \sqrt{2^A + M}
$$

$$
G = \sqrt{2^A - M}.
$$

(2)

Conversion between these two classes is done by as.MAData() and
as.RGData(). For instance,

```r
ma <- as.MAData(rg)
rg2 <- as.RGData(ma)
```

will create an MAData object ma from to the RGData object rg and then another
RGData object rg2. From the above discussion (assuming no non-positive sig-
nals), all of `rg$M`, `ma$M`, and `rg2$M` will return equal values. Moreover, objects
rg and rg2 are equal, which can be tested by

```r
> equals(rg, rg2)
[1] TRUE
```

Note that these are two different objects, making identical(rg, rg2) to re-
turn FALSE. Compare this with

```r
> rg3 <- rg
> identical(rg, rg3)
[1] TRUE
```

This is because `rg` and `rg3` refer to the same object (instance). At this point we
have to admit that we have been sloppy with the words. When we say “the ob-
ject `rg`” we actually mean “the object that the reference variable `rg` refers to”, but
having to do this each time would be too lengthy and we will allow ourselves to
use the shorter form whenever it is non-ambiguous to do so. Thus, the refer-
ences variables `rg`, `rg2`, and `rg3` refer to two objects only. Hence, the following
outcome:

>=WWe will display the R prompt ‘>' whenever both commands and results are shown in the same
code example, otherwise not.
> rg3 <- rg
> rg$R[1,1] <- 2
> rg3$R[1,1]
[1] 2

To further emphasize the difference between objects and references, to query the size of an object use

> objectSize(rg)
[1] 48716

and compare this with the built-in R function

> object.size(rg)
[1] 452

which will return the size of a reference variable itself, which is only a few hundred bytes large regardless of the size of the object it is referring to.

4 A walk-through of the package

The R software is started by typing

R

at the command prompt. On Windows platforms, the Rgui environment may also be used. When in R, the aroma package is loaded by

library(aroma)

giving output similar to

Loading required package: R.basic
Loading required package: R.oo
R.oo v0.52 (2004-07-01) was successfully loaded.
Loading required package: R.io
R.io v0.52 (2004-07-01) was successfully loaded.
Loading required package: R.graphics
Loading required package: R.colors
R.colors v0.52 (2004-07-01) was successfully loaded.
R.graphics v0.52 (2004-07-01) was successfully loaded.

which confirms that it is loaded together with all necessary packages. For a first test-run, try

example(GenePixData)

To quit R, use quit(). For installation instructions, see Section 7.2.
4. A walk-through of the package

4.1 Importing microarray data

This section briefly explains how to read data into aroma. The package provides methods to read and write microarray data that is stored in one of the many file formats generated by some of the most well-known cDNA microarray image analysis applications such as GenePix, Imagen, QuantArray, ScanAlyze, Spot [1, 19], and Spotfinder. Most versions of these applications are supported. Consider an example where three Axon’s GenePix Result (GPR) files named “slide1.gpr”, “slide2.gpr”, and “slide3.gpr” are stored in the current working directory. Any one of these files may be read by the static method `read()` of class `MicroarrayData`;

```r
gpr <- MicroarrayData$read("Slide1.gpr")
```

Static (class) methods are methods that are specific to a class, but not necessarily to an instance of it. This is typical for read methods for which an object does not exist prior to the call. To call static methods, the `$` operator is used, indicating that the method belongs to a class much like a field belongs to an object, or an element belongs to a list. The above method will return an object of a subclass to `MicroarrayData` (recall that this class is abstract) that is specific for the data set read. Among the commonly used ones, aroma defines classes such as `GenePixData`, `ImagenData`, `QuantArrayData` etc. to represent the different data formats.

In addition to this, several files may be read at once by either giving a vector of filenames, or by specifying a string pattern matching filenames, to be read;

```r
gpr <- MicroarrayData$read(pattern="*.gpr")
```

which will read (in alphabetic order) all files that ends with “gpr” into one `GenePixData` object. For more details on filename patterns, see R help on regular expressions, e.g. `?regexp`.

In the above two examples, the read method returns a reference, which we denote by `gpr`, to an object of class `GenePixData` (a subclass of `MicroarrayData`, as can be seen if we write `print(gpr)` or, shortly, by typing `gpr` at the command prompt. For the latter example we get;

```r
> gpr
```
From this we see that the object is of class `GenePixData` and contains 43 different measurements from 3 microarrays (slides) with 5184 spots. The array was printed with a 4x4 print-tips head etc. To investigate the contents of the object further, `names(gpr)` will return the names of all 43 fields, and `ll(gpr)` will return a detailed list of all field names, their types and their sizes etc. Here is an excerpt:

<table>
<thead>
<tr>
<th>member data</th>
<th>class</th>
<th>dimension</th>
<th>object size</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>% &gt; B532+1SD</td>
<td>matrix c(5184,3)</td>
<td>63000</td>
</tr>
<tr>
<td>2</td>
<td>% &gt; B532+2SD</td>
<td>matrix c(5184,3)</td>
<td>63000</td>
</tr>
<tr>
<td>3</td>
<td>% &gt; B635+1SD</td>
<td>matrix c(5184,3)</td>
<td>63000</td>
</tr>
<tr>
<td>4</td>
<td>% &gt; B635+2SD</td>
<td>matrix c(5184,3)</td>
<td>63000</td>
</tr>
<tr>
<td>5</td>
<td>B Pixels</td>
<td>matrix c(5184,3)</td>
<td>63000</td>
</tr>
<tr>
<td>6</td>
<td>B532 Mean</td>
<td>matrix c(5184,3)</td>
<td>63000</td>
</tr>
<tr>
<td>7</td>
<td>B532 Median</td>
<td>matrix c(5184,3)</td>
<td>63000</td>
</tr>
</tbody>
</table>

Alternatively to the `$`-operator, fields can be accessed through the `[[`-operator as follows:

```
gpr["% > B635+2SD"][c(4,5,14:20),1]
```

which again is similar to how list structures in R work. To know what methods are available for a given class, `print(GenePixData)`, or shorter just the class name typed at the R prompt, will give a summary of the class:

```
GenePixData extends MicroarrayData, Object {
    public layout
    public version
    public anonymize(...)
    public append(other)
    public getAbscent(slides=NULL, include=NULL, ...)
    ...
}
```

Like for any class or method, further information can be found in the help pages installed with the package, e.g. `help(GenePixData)` or shorter `?GenePixData`. For more details about the help facilities in aroma, see Section 5.2.
4. Walk-through of the package

4.2 Extracting the foreground and background signals

As seen in previous sections, data output by microarray image analysis software contains a large number of measurements per probe, but in most analysis it is only the foreground and background signals that are used. Because of this, the RawData class exists. The estimates of the foreground and the background signals in a GenePixData object can be extracted by

```r
raw <- getRawData(gpr)
print(raw)
```

```
[1] "RawData: R (5184x3), G (5184x3), Rb (5184x3), Gb (5184x3), Layout: Grids: 4x4 (=16), spots in grids: 18x18 (=324), total number of spots: 5184. Spot names are specified. Spot ids are specified."
```

This object contains 5184 foreground signals R and G, and 5184 background signals Rb and Gb from all three arrays. Note how the array layout is included automatically. A GenePixData object contains a few different types of foreground and background estimates. By default, the median pixel values are returned. To retrieve other estimates, see ?GenePixData. Continuing, if the other measurements in the GenePixData object is not needed anymore, it can be deleted by standard R procedures, that is

```r
rm(gpr)
```

If no other reference variables refer to the same object, the object will be deallocated by the built-in R garbage collector. Thus, even if reference variables are used, a user or a programmer does not have to worry about memory management. Note the decrease in memory usage;

```r
> objectSize(raw)
[1] 258416
> objectSize(gpr)
[1] 5374740
> objectSize(raw)/objectSize(gpr)
[1] 0.04807972
```

Next step in a typical microarray analysis is to perform background correction or not. This is done by one of the following two alternatives:

```r
ma <- getSignal(raw, bgSubtract=TRUE)
ma <- getSignal(raw, bgSubtract=FALSE)
```
4.3 Visualization

In microarray analysis, as well as all other statistical analysis, it is most useful to initially display data in as many ways as possible. This will show the nature of data and reveal problems such as systematic effects in measurements etc. In addition to the huge number of exploratory data analysis (EDA) methods available in R in general, the aroma package provides a large set of graphical EDA methods that are special for microarray data. They have been designed such that they require minimal “interaction” by the end user. For instance, axes are labelled and scaled automatically, color annotation is automatic, and highlighting of certain data points is made simple. Here are two examples:

\[ \text{plotSpatial(gpr)} \]
\[ \text{plotSpatial(ma)} \]

Both of these commands will create a spatial plot of the 5184 observed log-ratios that by default are colored in a red-to-green and dark-to-bright scale. Other color scales may be chosen and color filters are available. Since a `GenePixData` object contains information about the spot positions and spot diameters, the first command will display the signals as discs of different sizes as shown in the left panel of Figure 1. For the second command no spot positions or sizes are available, but only the array layout, resulting in the image in the right panel.

![Spatial plots of two-color microarray data.](image-url)

**Figure 1**: Spatial plots of two-color microarray data. *Left*: Spatial plot from a `GenePixData` object. *Right*: Spatial plot for an `MAData` object with only array layout information.
For all *MicroarrayData* classes a generic `plot()` method is defined. Depending on which class the object belongs to, different types of plots will be created. For instance, calling `plot(rg)` on an *RGData* object `rg` will generate a scatter plot where the red signals are plotted against the green signals. Similarly, for an *MA-Data* object a log-ratio versus log-intensity plot will be generated. Moreover, the generic `highlight()` method will highlight certain spots in any type of plot generated by an aroma plot methods. For example, in

```r
idx <- (abs(ma$M) > 1.5)
plot(ma)
highlight(ma, idx, col="blue")
rg <- as.RGData(ma)
plot(rg)
highlight(rg, idx, col="blue")
```

spots with an absolute log-ratios above 1.5 are highlighted. The outcome is as in Figure 2. Note, if the `highlight()` method would be called after, say, `plotSpatial()`, the spots would be highlighted in the spatial plots. For the interested, this is another example where reference variables have been used; the objects remember what the last plot call was and this information is used when the highlighting is done.

In addition to spatial and X-Y scatter plots, there are various other plot method such as print-order [2], dip-order [2], and box-and-whisker plots. For example,

```r
plotDiporder(ma)
boxplot(ma, groupBy="printtip")
```

with results as in Figure 3. Additional plot styles will be given in the following sections. For a complete list of plot methods, see the help pages.

### 4.4 Calibration and normalization

Calibration and normalization of DNA microarray data is about removing artifacts in the observed signals that are due to systematic variations in the microarray process and not due to biological variation, which is the variability of interest. Removing systematic effects is necessary in order for higher-level analysis such as identification of differentially expressed genes and gene classification to be correct.
For a formal definition of the difference between calibration and normalization, see the introduction of [5]. Here is an example of a curve-fit normalization:

```r
plot(ma)
normalizeCurveFit(ma)
plot(ma)
```
4. A walk-through of the package

Note, normalization methods in aroma will update the object passed to it. This will save memory, but it means that the non-normalized data will be lost. If this is not wanted, the object can be cloned before calling the normalization method;

```r
maNormalized <- clone(ma)
normalizeCurveFit(maNormalized)
```

To normalize data such that the scales of the log-ratios becomes equal for all arrays, so called across-slide (scale) normalization can used;

```r
normalizeScaleAcrossSlides(ma)
```

Note that this should not be necessary for the quantile and the affine normalization methods that are described next. Affine normalization [7], which is performed in the (G,R)-signal space contrary to the (A,M) space, is done by

```r
normalizeAffine(rg)
```

The effect of the affine normalization in the M versus A plane is depicted in Figure 4. As a final example, quantile normalization [10] is done by

```r
normalizeQuantile(rg)
```

Quantile normalization will normalize data in each separate channel such that the empirical densities of the signals from all arrays become as similar as possible. See Figure 5, which compares the density functions from six arrays normalized in three different ways.

Several of the normalization methods can be applied spatially, in one dimension such as the smooth print-order normalization, or by stratifying on gene or spot group, e.g. by print-tip, microtiter plate, clone library etc. For instance, to apply affine normalization on each print-tip individually, do

```r
normalizeAffine(rg, groupBy="printtip")
```

and similar for the other normalization methods.

When several scans of the same array are available, the robust affine multi-scan calibration method described in [8] can be utilized. It will correct for scanner biases, which will remove intensity-dependent effects due to the scanner, and calculate the average signals so that the scanner's effective dynamical range is extended and
Figure 4: Left: Observed log-ratios versus log-intensities. The overlaid grid is the rotated \((\log_2 G, \log_2 R)\)-grid as it is after being deformed by (estimated) affine transformation. The left corner corresponds to \((G, R) = (1, 1)\) and the dashed curve is the where the normalized data satisfies \(G = R\). Right: The affinely normalized signals with grid deformations removed.

Figure 5: Density plots for the signals (logarithm base 2) in the red channels from six different arrays after various normalization methods (the labels on the axes read “R” (red channel) and “density (integrates to one”). Left: Curve-fit normalization and across-slide scale normalization. Middle: Affine normalization. Right: Quantile normalization.

the signal-to-noise level is increased. This is illustrated in Figure 6. Multiscan calibration, which should be applied before other normalization methods, is done by

calibrateMultiscan(rg)
where the columns of the RGData fields represent signals from multiple scans of the same array. Multiscan calibration is applied to each channel separately. For

![Image](image_url)

**Figure 6:** Example of a multiscan calibration of four scans from the same array. Left: Green versus green signals for all six possible pairs from the four different scans done at various sensitivity settings. Middle: The same data plotted as within-channel $M$ vs $A$ plots. Right: The same but now after calibration. Intensity-dependent effects have been removed and the signals can be averaged to decrease noise levels.

further details and for all available calibration and normalization methods, see the aroma help pages.

## 4.5 Exporting microarray data

Normalized microarray data can be exported for further analysis in other applications or to be saved to file for future usage. To save a MicroarrayData object to a tab-delimited file do

```r
write(ma, "ma.normalized.dat")
write(rg, "rg.normalized.dat")
```

and similar for objects of other classes.

## 5 Miscellaneous features

### 5.1 Report generators

For everyday analysis, a user will most likely write so called *batch scripts* that will perform many steps automatically. To support such batch jobs, a set of Reporter
classes that generate documents containing text, tables and figures, have been developed. Reports in plain text (no graphics), in HTML, and in LaTeX, are generated by the `TextReporter`, `HtmlReporter` and `LaTeXReporter` classes, respectively. For the records, these classes are implemented by the `R.io` package (part of the `R.classes` bundle), which is required (see Section 7.2) and loaded when aroma is loaded. Because all `Reporter` classes implement the same API, generating figures (of appropriate format), tables, lists, code examples, table of contents etc. is done in a uniform way regardless of the output format. Reports of multiple formats can be generated in parallel by utilizing the `MultiReporter` class. For an example of a microarray analysis report generated by the above classes, see Figure 7. As

![Figure 7: Example of an HTML page (left) and a LaTeX document (right) comparing different normalization methods. These were generated in parallel by an aroma script utilizing the MultiReporter class.](image)

a complement to `Reporter` classes, Sweave [14], which is part of the tools package, can be used to write a hybrid of LaTeX and R documents. Such documents may contain R code that generate figures, tables and other visual output, which is automatically inserted into the generated LaTeX document. The main difference between Sweave and the `Reporter` classes is that with the former, the report is written as a document from which the R code is extracted and evaluated whereas with the latter it is the R script that generates the document. Both can be used together.

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5.2 Help, manuals and other documentation

The aroma package is continuously updated with new algorithms and visualization methods. For this reason, the package’s help documentation, which comes with all installations and is updated equally often, should always be queried in the first place. The HTML version of these help pages can be accessed by typing `help.start()`, clicking the “Packages” link followed by the “aroma” link. At the top of this root help page for aroma there is a list of links to separate sections, e.g. “Introduction”, “Classes”, “Import and Export of Data”, “Normalization” and so on. All public methods and functions of the package are listed in these sections. Note that each method is also accessible via the page for the class for which it is coupled to. The “Classes” section lists all classes in the package with links to detailed help for each of them. For several classes and methods complete examples are given, examples which can be tried either by cut’n’paste or by

```r
example(GenePixData)
```

It is good to know that all examples are asserted to be executable when a new release of the package is built; invalid source code in the examples is not allowed. In addition to be available in HTML, the same help pages are available as a single LaTeX-compiled documentation for easy printing. At the moment of writing, this document consists of over 280 pages. See Figure 8 for an example. Additional help and examples are available online at http://www.maths.lth.se/help/R/.

5.3 Compatibility with other packages and software

Initially, this package was implemented as an object-oriented extension to the sma package [9], but is now a self-contained standalone package. The RawData and RGData classes are directly compatible with the structures in sma. In addition to this, classes that have virtual fields (see Section 6.4) that are named the same as the fields of the aforementioned two classes are compatible with sma. Similarly, converting sma data structures to aroma MicroarrayData objects is straightforward. See ?SMA for more details. For this reason, the two packages can be used in parallel and all the functions of the sma package can be called using aroma objects. However, many of the functions in sma have been re-implemented in this package. The aroma package is compatible with methods provided by microarray packages such as limma [16] and other Bioconductor [17] packages. Moreover, for
Note that the layout must be set for print-tip (method="p") and scaled (method="s") normalization. If layout is not set, an exception will be thrown. Normally, the layout is already set, such as when the data is read from for instance GenePix, ScanAlyze and Spot.

Author(s)
Henrik Bengtsson (http://www.braju.com/R/). Initial code for support of 'weights' by Jon McAuliffe, Statistics Dept, UC Berkeley. The original code was written by the sma authors Yee Hwa Yang ⟨yeehwa@stat.berkeley.edu⟩, Sandrine Dudoit ⟨sandrine@stat.berkeley.edu⟩ and Natalie Roberts ⟨nroberts@wehi.edu.au⟩.

References

See Also
For across-slide normalization see *normalizeAcrossSlides*. For more information see MAData.

Examples
# The option ,dataset, is used to annotate plots.
options(dataset="sma:MouseArray")

SMA$loadData("mouse.data")

layout <- Layout$read("MouseArray.Layout.dat", path=aroma$dataPath)

raw <- RawData(mouse.data, layout=layout)

ma <- getSignal(raw)

# Clone the data to get one non-normalized and one normalized data set.
ma.norm <- clone(ma)

# Normalize the data within slides using scaled print-tip normalization.
normalizeWithinSlide(ma.norm, "s")

# Plot data before and after normalization.
layout(matrix(1:4, ncol=2, byrow=TRUE))

plot(ma)

plotSpatial(ma)

plot(ma.norm)

plotSpatial(ma.norm)

6 Philosophy of design and implementation

The package has been designed in a true object-oriented manner with an implementation making use of references variables. As discussed more below, passing objects by reference is useful if the objects are huge and if the objects are “modified” frequently. This is undoubtedly the case for microarray data and analysis. Moreover, references make it possible to create a more user-friendly methods interface, which is something that we have taken advantage of in this package. The rest of this section contains details that are not necessary to understand in order to start using aroma.
6. Philosophy of design and implementation

6.1 Underlying classes

In order to implement the above design, the *R.oo* package [4] has been developed. It is part of a package bundle named *R.classes* and provides the core functionality for object-oriented design with reference variables. *R.oo* is based exclusively on the S3/UseMethod programming paradigm [18], although future versions may in addition use S4/methods [11]. The package provides a transparent way of using reference variables for both the user and the developer. Passing objects to methods as pass-by-reference makes it possible for the methods to change the state (and the contents) of objects directly without having to return a modified copy. That is, objects are *mutable*. The standard approach in functional languages such as R is that all objects are non-mutable and a new instance is created whenever an object is modified. For further discussion of philosophical and practical considerations between the functional and object-oriented approaches, see [4]. Instances of classes that are derived from the root class *Object*, which is defined in R.oo, will automatically be accessed by reference variables.

The object-oriented core was separated from the rest of the microarray package and put into the R.oo package. The major advantage is that the R.oo package is likely to be used in other projects and by other developers and end users. A larger user base means more CPU test time, which increases the likelihood for identifying programming errors that will otherwise not be found by standard test procedures. Hence, a separate core package will improve the overall quality of the microarray package.

6.2 User interface

The aroma package does not have a GUI per se, but instead a API for access to all methods directly at R commando prompt. As shown by the examples in previous sections, reference variables and mutable objects make it possible to reduce the number of arguments that the user needs to specify. This way, the API can be kept simple. However, provided mutable objects, it should be straightforward to add a light-weighted third-party GUI utilizing, say, Tcl/Tk, which is part of the default R installation.
6.3 Programming style

In order to provide a well defined API simple enough to be used by anyone, *The R Coding Convention* (RCC) document [3] was developed. The RCC, which is an open suggestion to the R community, contains naming conventions similar to those found in Java, where classes, variables and methods can easily be differentiated by their names. For instance, names of classes are with mixed-case letters, starting with an upper-case letter, methods and variables start with a lower-case letter, and constants consists only of upper-case letters, e.g. `GenePixData`, `normalizeAffine()`, and `RED.HUE`. By enforcing the RCC, the time needed for a new user to become familiar with the API will be reduced.

6.4 Virtual fields

A virtual field is a field that is calculated *on demand* or *on the fly*. This can remove redundancy, which in turn will lower the risk for programming errors. It also makes the code more memory efficient. By providing a method, say, `getM()`, the virtual field `M` is automatically provided. Analogously, `setM(obj,value)` will add the option to assign values by `obj$M <- value`. Virtual fields are implemented by the `Object` class. With virtual fields, details can be hidden from the user, fields can be made read-only, and the correctness of assignment to fields can be asserted at the time of the assignment using specially written set methods. Moreover, with virtual fields it is possible to change the internal representation of the microarray objects and at the same time remain compatible with old packages, but also with new packages such as those developed under the Bioconductor project [17]. For instance, assume that a new normalization method that normalizes the fields `ch1` and `ch2` in a list structure is available in another package. For an aroma class to utilize this new method all that is needed is to provide the `get` and `set` methods `getCh1()` and `setCh1()`, respectively, and the same for `ch2`.

7 Obtaining the package

7.1 Requirements

In addition to a recent installation of R [13], the aroma package requires that the `R.classes` bundle is installed. The package can be used on a typical Windows, Mac OS X, various Linux systems, Sun Solaris and any operating system that supports R. Memory requirements are constrained by the size of the microarray data,

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7. Obtaining the package

but typically at least 256 Mb is required although 512-1024 Mb is recommended.

7.2 Installation and updates

To download and install aroma and its required packages, write

```
install.packages(c("R.classes", "aroma"),

contriburl="http://www.maths.lth.se/help/R")
```

For source code and manual installation go to http://www.maths.lth.se/help/R/. The R software is available from http://www.r-project.org/.

7.3 License

The R system is licensed under GNU General Public License (GPL). R.classes and aroma are licensed under GNU Lesser General Public License (LGPL). The GPL and LGPL can be obtained from http://www.gnu.org/copyleft/ or by writing to the Free Software Foundation, Inc., 59 Temple Place – Suite 330, Boston, MA 02111-1307, USA.

Please cite this document whenever using aroma. When using R.classes, please cite [4].
References


Affine calibration for microarrays with dilution series or spike-ins

Henrik Bengtsson¹ and Ola Hössjer²

¹ Mathematical Statistics at the Centre for Mathematical Sciences, Lund University, Box 118, SE-221 00 Lund, Sweden. Email: hb@maths.lth.se.
² Mathematical Statistics at the Mathematics Department, Stockholm University.

ABSTRACT

Background: In this theoretical study we follow up our previous theoretical and applied work on affine calibration and normalization methods by suggesting a stochastic affine model for two- or multi-channel microarrays containing dilution series. With replicated probes (spots) at various, not necessarily known, concentration levels it is possible to identify the channel biases uniquely. For this to be true it is necessary that some genes are differentially expressed, but which they are or what their relative gene-expression levels are do not have to be known. The method presented may also be applied to so called spike-in data. From the maximum likelihood estimator, estimates of the relative gene-expression levels for all dilution series follow directly. Given the estimated channel biases and scale factors, back transformation gives calibrated probe signals that are proportional to the amount of fluorophore in each channel. With the assumption that most genes are non-differentially expressed, the proposed calibration method makes the signals in each channel proportional to the corresponding gene-expression levels.

Results: The model suggested is a heteroscedastic affine model where the standard deviations of the error terms are equal for all replicates that belong to the same gene, but may vary freely between genes. For a specific probe, the standard deviation in one channel is assumed to be proportional to the other channels. These constraints were chosen as a compromise between model flexibility and feasible parameter estimates. We investigate the properties of the parameter estimates from simulated data by comparing bootstrap and asymptotic confidence intervals based on normal approximation for different setups of number of genes and replicates. We simulate two-channel gene-expression data with technical dilution series from the suggested model and, in order to investigate the robustness of
the estimator against model misspecification, a more general model for which the standard deviations are not only proportional to the gene-expression levels, but also the probe concentrations. We find that the standard deviation of the bias-parameter estimates is inversely proportional to the square root of the product of the number genes and replicates, provided the latter is not too small. For modest number of replicates, the standard deviation based on the Fisher information is a good approximation to the bootstrap ditto. When too few replicates are available, normality can not be assumed and the standard errors are under-estimated. For similar reasons, for different setups of number of genes and replicates such that the total number of probes are the same, we found that it is better to use more replicates and fewer genes, if possible. Comparing dilution series with uniformly and logarithmic-uniformly distributed concentrations better results are obtained using the latter.

**Availability:** All methods have been implemented in the cross-platform R package called aroma, which is available for free from [http://www.braju.com/R/](http://www.braju.com/R/).

**Keywords:** spotted microarrays; systematic effects; bias; calibration; normalization; dilution series; technical replicates; profile likelihood; Fisher information; stochastic affine model; heteroscedastic noise.

1 Introduction

On a microarray there is a large number of disjoint probes, which each consists of a homogeneous set of immobilized DNA with a known or at least an identifiable sequence. For a typical microarray experiment, for each cell sample for which a gene-expression profile is to be obtained, RNA is extracted and reversely transcribed into cDNA. In the same or a following process, the cDNA is labeled with a unique dye, commonly a fluorescent dye. The set of cDNA is referred to as the target. In single-channel technique identically labeled cDNA constitutes the target whereas in two-channel techniques two differently labeled cDNA sets of equal amount constitute the target. The target is then let to hybridize to the microarray so that, ideally, sequences of target cDNA that match the DNA in a specific probe will hybridize to that probe and nowhere else. The effect is a massive parallel and spatial sort of target sequences to the corresponding probes. Next, using a microarray scanner the microarray is scanned in different
wavelength channels, one for each dye used, resulting in one or several high-resolution spatial images. Using image analysis techniques, the pixels in the images corresponding to the probe regions are identified and the amount of labeled cDNA in each probe and channel is quantified based on the pixel intensities.

Thus, the main objective when scanning a microarray is to quantify the amount of fluorescent molecules in each probe and each channel. Then, by assuming that the amount of hybridized DNA is proportional to the amount of fluorophores and so on, estimates of the gene-expressions can be obtained. However, in this paper we will focus solely on the problem of correctly quantifying the amount of different fluorophores in the probes.

As an extension to our previous work [3, 2], we suggest an affine calibration (normalization) method based on a stochastic model for two- or multi-channel microarrays containing dilution series. With minor additional assumptions, the same calibration method can be used for so-called spike-in calibration data where genes with known amounts of cDNA have been added. Without being further discussed, with additional assumptions it may also be applied to gene-expression signals obtained from multiple single-channel microarrays.

The outline of the paper is as follows. In the "Model" section, we present the stochastic affine model describing scanned microarray data with within-array replicates. We outline the necessary assumptions and model constraints in order to make all parameters identifiable. In the following "Methods" section, we provide a model estimator based on maximum likelihood techniques, which forms the basis for the calibration method described thereafter. Different properties of the parameter estimates are discussed as well as the two bootstrap methods used. How the microarray data is simulated from two different models and with two types of dilutions series is described in the "Data" section. In the "Results" section, the properties of the estimates for microarrays with different number of genes, different number of replicates, and different types of replicates etc. are presented. A brief discussion then follows and conclusions are given in the last section.
2 Model

Consider a microarray experiment involving genes $i = 1, \ldots, I$ from RNA extracts $c = 1, \ldots, C$ with $C \geq 2$. From now on, we will refer to each set of signals from each RNA extract as a channel. Let $\mu_{c,i}$ be the true gene-expression (transcription) level of gene $i$ in channel $c$. Ideally, statistical analysis is done on these quantities, but here we will focus on the problem of quantifying the (relative) amount of fluorescent dyes of a certain kind in each probe.

Within-array technical replicates are probes on the microarray that have identical nucleotide sequences. For instance, in spotted microarrays, technical replicates are spots that are printed with the same probe sequence from the same clone library. Formally we say that for gene $i$ there exist replicates $j = 1, \ldots, J_i$ on the microarray so that in total there are $J_i = \sum_{i=1}^{I} J_i$ probes on the array. Let index $(i, j)$ refer to the $j^{th}$ replicated probe for gene $i$. Furthermore, let $z_{c,i,j}$ be the concentration of fluorescent molecules that are attached to DNA sequences hybridized to probe $(i,j)$ in channel $c$. Note, some of these may sit on cross-hybridized targets. However, we do not differentiate between cross-hybridized and true target DNA in this study. With the assumption that the relative amount of true and cross-hybridized target DNA is the same for all replicated probes of the same gene, we have, if assuming everything else is equal, that

$$z_{c,i,j} = z_{c,i}; \forall j. \quad (1)$$

The wavelength profiles of the fluorescent light from replicated probes are expected to be equal up to a global scale factor.

A dilution series is a set of within-array technical replicates where the replicated probes have different concentrations of DNA sequences. Let the probe concentration of probe $(i,j)$ be denoted by $b_{i,j}$ (in arbitrary units). Moreover, assume that the brightness (excitation coefficient times quantum yield) of a probe is proportional to the probe concentration and the amount of excitation light, which in turn is assumed to be constant, and can be summarized in a channel-specific quantity called $b_c$. The total amount of fluorescent light $x_{c,i,j}$ from probe $(i,j)$ with “gain” $b_c$ can then be modeled as

$$b_c x_{c,i,j} = b_c b_{i,j} z_{c,i}; \forall c. \quad (2)$$
2. Model

We assume no spectral cross-talk and no saturation. Added to the fluorescent light is also stray white and stray laser light, auto-fluorescent light, photocathode noise, electronic noise [3] etc., which we assume all can be modeled as a channel-specific but probe-independent constant $a_c$ plus zero-mean noise.

2.1 The affine stochastic model

Let $y_{c,i,j}$ be the observed expression-level for replicate $j$ of gene $i$ in channel $c$. We assume that

$$y_{c,i,j} = a_c + b_c x_{c,i,j} + \varepsilon_{c,i,j}$$

(3)

where $\varepsilon_{c,i,j}$ is independent noise with $E[\varepsilon_{c,i,j}] = 0$ and $V[\varepsilon_{c,i,j}] = \sigma_{c,i,j}^2$. Furthermore, assume that

$$\sigma_{c,i,j} = \sigma_{c,i}$$

(4)

$$\sigma_{c,i} = k_c \sigma_i$$

(5)

for $c = 1, \ldots, C$ where $k_1 = 1$. In words, this means that the relative standard deviation of the noise in the different channels does not depend on replicate. Let

$$\beta_{c,i} = \frac{b_c z_{c,i}}{b_1 z_{1,i}} = \frac{b_c x_{c,i,j}}{b_1 x_{1,i,j}}, \forall j.$$  

(6)

Thus, $\beta_{c,i}$ is up to a scale factor equal to $z_{c,i}/z_{1,i}$, which is the “expression level” of gene $i$ in channel $c$ in units of the corresponding expression level in channel 1. With the above parametrization and no further model assumptions $b_c$ and $x_{c,i,j}$ are non-identifiable. Define $\gamma_{c,i,j} = b_c x_{c,i,j}$. The model can then be written as

$$y_{c,i,j} = a_c + \beta_{c,i} \gamma_{1,i,j} + \varepsilon_{c,i,j}; \forall c, i, j$$

(7)

where all parameters are identifiable. The $2C - 1$ structural parameters, here the parameters independent of gene, are $\theta = (a, k)$ where $a = (a_1, \ldots, a_C)$ and $k = (k_2, \ldots, k_C)$. The $C + J_i$ incidental parameters for gene $i$ are $\xi_i = (\beta_i, \gamma_{1,i,j})$, where $\beta_i = (\beta_2,i, \ldots, \beta_{C,i})$ and $\gamma_{1,i,j} = (\gamma_{1,i,1}, \ldots, \gamma_{1,i,J_i})$. Let $\xi = \{\xi_i\}_i$, $\beta = \{\beta_i\}_i$, $\gamma_1 = \{\gamma_{1,i}\}_i$, and $\sigma = \{\sigma_i\}_i$. In total there are $|\theta| + |\xi| = C(I + 2) + J_{i} - 1 = p$ parameters.
3 Methods

We will next present how the above model with Gaussian noise can be estimated using maximum likelihood techniques.

3.1 Maximum likelihood estimation

Let \( y = \{y_i\}_i \) where \( y_i = (y_{1,i,1}, \ldots, y_{1,i,J_i}, \ldots, y_{C,i,1}, \ldots, y_{C,i,J_i}) \in \mathbb{R}^{CJ_i} \) is the vector of the observations for gene \( i \) in all channels and all replicates \( j = 1, \ldots, J_i \). The log-likelihood function of the parameters given the data is then

\[
l(\theta, \xi; y) = \sum_{i=1}^{I} l_i(\theta, \xi_i; y_i). \tag{8}\]

Assuming Gaussian zero-mean noise with probe specific variation \( \varepsilon_{c,i,j} \in N(0, \sigma^2_{c,i,j}) \), the log-likelihood for observation \( y_{c,i,j} \) is up to an additive constant

\[
l_{c,i,j}(\theta, \xi_i; y_{c,i,j}) = -\ln \sigma_{c,i,j} - \frac{1}{2} \left( \frac{y_{c,i,j} - (a_c + \gamma_{c,i,j})}{\sigma_{c,i,j}} \right)^2 \tag{10}\]

With assumptions (4) and (5), the log-likelihood for gene \( i \) becomes

\[
l_i(\theta, \xi_i; y_i) = \sum_{c=1}^{C} \sum_{j=1}^{J_i} l_{c,i,j}(\theta, \xi_i; y_{c,i,j}) \]

\[
= -\sum_{c=1}^{C} \sum_{j=1}^{J_i} \ln (k_c \sigma_i) - \frac{1}{2} \sum_{c=1}^{C} \sum_{j=1}^{J_i} \left( \frac{y_{c,i,j} - (a_c + \gamma_{c,i,j})}{k_c \sigma_i} \right)^2 
= -J_i \sum_{c=2}^{C} \ln k_c - CJ_i \ln \sigma_i - \frac{1}{2} \sigma_i \sum_{c=1}^{C} \frac{1}{k_c^2} \sum_{j=1}^{J_i} \left( y_{c,i,j} - a_c - \gamma_{c,i,j} \right)^2. \tag{11}\]

Finally, the overall log-likelihood is

\[
l(\theta, \xi; y) = -J \sum_{c=2}^{C} \ln k_c - C \sum_{i=1}^{I} J_i \ln \sigma_i 
- \frac{1}{2} \sum_{c=1}^{C} \frac{1}{k_c^2} \sum_{i=1}^{I} \frac{1}{\sigma_i^2} \sum_{j=1}^{J_i} \left( y_{c,i,j} - a_c - \gamma_{c,i,j} \right)^2. \tag{12}\]
3. Methods

3.1.1 Estimation of structural parameters

The profile log-likelihood for the structural parameters is

\[
l_p(\theta; y) = \sup_{\xi} l(\theta, \xi; y) = \sum_{i=1}^{J} \sup_{\xi_i} l_i(\theta, \xi_i; y_i) = \sum_{i=1}^{J} l_i(\theta, \hat{\xi}_i(\theta); y_i)
\]  

(13)

where \(\hat{\xi}_i(\theta)\) maximizes (11) given \(\theta\). Finally,

\[
\hat{\theta} = \hat{\theta}_{\text{ML}} = \arg \max_{\theta} l_p(\theta; y)
\]

(14)

is the maximum likelihood estimate of \(\theta\). In practice, we search over an iteratively refined \((2C - 1)\)-dimensional grid box constrained for \(\theta\).

3.1.2 Estimation of incidental parameters

The parameter vector \(\xi_i = (\beta_{c,i}, \gamma_{1,i}, \sigma_i)\) can be estimated as follows. We begin by estimating the parameters \(\beta_i = (\beta_{1,i}, \beta_{2,i}, \ldots, \beta_{C,i})\) given the assumption that \(\theta\) is the true structural parameter. Let

\[
y_{c,i,j} = y_{c,i,j} - \kappa_c = \gamma_{c,i,j} + \varepsilon_{c,i,j} = \beta_{c,i} \gamma_{1,i} + \varepsilon_{c,i,j}
\]

\[
\tilde{y}_{c,i,j} = \tilde{\beta}_{c,i} \gamma_{1,i} + \tilde{\varepsilon}_{c,i,j}
\]

where \(\tilde{\beta}_{c,i} = \beta_{c,i}/\kappa_c\) and \(\tilde{\varepsilon}_{c,i,j} = \varepsilon_{c,i,j}/\kappa_c\) so that \(V[\tilde{y}_{c,i,j}] = V[\varepsilon_{c,i,j}] = \sigma_i^2\).

Thus, given the true structural parameters \(\theta\) with noise-free observations, the \(J_i\) translated and rescaled points \(\tilde{y}_{i,j} = (\tilde{y}_{1,i,j}, \ldots, \tilde{y}_{C,i,j})\), \(j = 1, \ldots, J_i\) in \(\mathbb{R}^C\) are located along the line \(\tilde{L}_i(0, \tilde{\beta}_i)\) that goes through the origin and has direction \(\beta_i = (\hat{\beta}_{1,i}, \ldots, \hat{\beta}_{C,i})\) with \(\beta_{c,i} \geq 0; \forall c\). With independent and identically distributed noise, the observations cluster along the line \(L(0, \beta_i)\) and estimates of \(\beta_{c,i}\) and \(\gamma_{1,i,j}\) given \(\theta\) are found from the line in \(\mathbb{R}^C\) that minimizes the sum of squared orthogonal distances to \(\{\tilde{y}_{i,j}\}\). A version of principal component analysis (PCA) where data is not centered in advance can be used to estimate this linear subspace, where the direction of the estimated \(\tilde{L}_i\) is defined by the first principal component of \(\tilde{y}_i = \{\tilde{y}_{i,j}\}\), which is described by the components of \(v_i = (v_{1,i}, \ldots, v_{C,i}); \|v_i\| = 1\). Each component is then, up to a scale factor, an estimate of the corresponding \(\beta_{c,i}\). Since \(\beta_{1,i} = k_1 = 1\), we obtain

\[
\hat{\beta}_{c,i}(\theta) = \frac{k_c}{\|v_{1,i}\|} \cdot v_{c,i}; \forall c.
\]

(16)
Figure 1: Example of the estimate of $\beta_i$ and $\gamma_{1,i}$ for $C = 2$ and $J_i = 10$. **Left:** $J_i$ data points in $(y_1, y_2)$. $\alpha = (20, 70)$ is marked with a cross. **Middle:** The same data points in the translated space. **Right:** The translated and rescaled $(k = (1, 5))$ space in which the PCA fit is done. The fit is displayed as a line. The same fit is illustrated in the other spaces too by back transformation.

To estimate $\gamma_{1,i}$, we observe that when projecting the $j^{th}$ replicate $\tilde{y}_{i,j}$ of gene $i$ onto the PCA estimated line $\tilde{L}_i(0, v_i)$, the first coordinate of the projection is $\hat{\gamma}_{1,i,j}(\theta)$ and its distance from the origin is $\lambda_{i,j}$. From this it follows

$$\hat{\gamma}_{1,i,j}(\theta) = v_1 \lambda_{i,j}; \forall j.$$

Finally, given $\theta$ and estimates of $\beta_{c,i}$ and $\{\gamma_{1,i,j}\}_{j}$, a bias-corrected estimate of $\sigma_i^2$ is

$$\hat{\sigma}_{corr, i}^2(\theta) = \frac{1}{J_i C - (C + J_i - 1)} \cdot \sum_{c=1}^C \frac{1}{k_c^2} \sum_{j=1}^{J_i} (y_{c,i,j} - a_c - \hat{\beta}_{c,i}(\theta)\hat{\gamma}_{1,i,j}(\theta))^2.$$  

Thus, given $\theta = (\alpha, k)$ we have obtained an estimate of $\xi_i = (\beta_i, \gamma_{1,i}, \sigma_i)$ for gene $i$.

### 3.1.3 Calibration

With the assumption that most probes are non-differentially expressed, it is straightforward to estimate $b_c; \forall c$. Hence, estimates of the channel $c$ to channel
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One relative concentration of fluorescent molecules \( \{ z_{c,i} / z_{1,i} \}_{c,i} \) follow immediately from (6) and the maximum likelihood estimates of \( \{ \beta_{c,i} \}_{c,i} \). If it is only these quantities, which each summarizes all signals from a complete calibration series, that are of interest, no further analysis or pre-processing of the signals is needed. However, if for various reasons further calibration or normalization is to be applied on the probes separately, or another method to combine the signals from the replicated probes is to be used, the calibrated probe signals can be calculated as

\[
\hat{x}_{c,i,j} = \frac{y_{c,i,j} - \hat{a}_c}{\hat{b}_c}; \quad \forall c, i, j.
\]  

(19)

3.2 Properties of the estimates

Main focus will be on the properties of the bias-parameter estimates \( \{ a_c \}_c \) as these are of major importance in order to calibrate the measurements correctly, but we will also look at the estimates of the relative gene-expression levels \( \{ \beta_{c,i} \}_{c,i} \). Especially, we study the standard deviation and confidence intervals of the estimates. Such can be obtained both by large sample normal approximations and from bootstrap simulations, as described next.

3.2.1 Confidence interval from Fisher information

From model (3) with error structure (5), it is possible to obtain a closed form expression for the Fisher information of the bias-parameter estimates and, hence, standard deviation and confidence interval estimates. For large samples, classical theory gives that the parameter estimates are approximately normally distributed, with a standard deviation that can be estimated as the square-root of the diagonal terms of the inverse of the \( p \times p \) Fisher information matrix \( I \). In particular, for the bias parameters \( a_1, \ldots, a_C \) we obtain

\[
\hat{\sigma}_{a_c}^2 = \hat{V}[\hat{a}_c] \approx (\hat{I}^{-1})_{a_c a_c}
\]

where \( \hat{I} \) is an estimate of \( I \) obtained by replacing unknown parameter values by estimates. As shown in appendix A, for \( C = 2 \) we have

\[
1/(\hat{I}^{-1})_{a_c a_c} = \frac{1}{\hat{\beta}_c^2} \sum_{i=1}^I w_i (\hat{\beta}_{2,i} - \hat{\beta}_2)^2
\]

(20)

where \( \hat{\beta}_c = \left( \sum_{i=1}^I w_i \hat{\beta}_{c,i} \right) / \sum_{i=1}^I w_i \) is a weighted average of all “relative expression levels” in channel \( c \), and similar for \( \hat{\beta}_c^2 = \left( \sum_{i=1}^I w_i \hat{\beta}_{c,i}^2 \right) / \sum_{i=1}^I w_i \).
The weights are
\[ w_i = \frac{J_i}{\sigma_i^2 (k_i^2 + \beta_i^2, i)} \cdot \frac{\sum_{j=1}^{J_i} (b_{i,j} - \bar{b}_i)^2}{\sum_{j=1}^{J_i} b_{i,j}^2} \]  
(21)
where \( \bar{b}_i = \frac{\sum_{j=1}^{J_i} b_{i,j}}{J_i} \) is the average probe concentration for gene \( i \). It is interesting to notice from (20)-(21) that the asymptotic standard error \( \hat{\sigma}_{ac} \) decreases the more the expression levels \( \{\gamma_{1,i,j}\}_j \) vary between replicates. Equivalently, the more similar the \( \{b_{i,j}\}_j \) are, the smaller the Fisher information is. Moreover, when the relative expression levels \( \{\beta_{2,i}\}_i \) are close to each other the standard error is large.

It now follows that an approximate \((1 - \alpha)\)-level confidence interval for bias parameter \( a_c \) is
\[ R_{ac} = [\hat{a}_c - \lambda_\alpha/\sqrt{\hat{\sigma}_{ac}}, \hat{a}_c + \lambda_\alpha/\sqrt{\hat{\sigma}_{ac}}] \]  
(22)
where \( \lambda_\alpha = \Phi^{-1}(1 - \alpha) \) and \( \Phi(\cdot) \) is the distribution function for \( N(0, 1) \). Note that, with simulated data we can obtain standard deviations and confidence intervals using both the true incidental parameters \( \{k_c\}_c, \{\sigma_i\}_i, \{\beta_{2,i}\}_i, \) and \( \{\gamma_{1,i,j}\}_i,j \), as well as their estimates.

For the gene-expression levels we have \( \hat{\sigma}_{\beta_{2,i}}^2 = \hat{\upsilon}(\hat{\beta}_{c,i}) \approx (\hat{I}^{-1})_{\beta_{c,i}, \beta_{c,i}} \). It is shown in the appendix that for \( C = 2 \),
\[ 1/(\hat{I}^{-1})_{\beta_{2,i}, \beta_{2,i}} = \frac{\sum_{j=1}^{J_i} b_{i,j}^2}{\sigma_i^2 (k_i^2 + \beta_i^2, i)} + O(J_i^{-1}), \]  
(23)
where the remainder term of size \( O(J_i^{-1}) \) is present since the bias parameters \( a_1 \) and \( a_2 \) are unknown and have to be estimated. For large \( J_i \) we may drop this term and obtain confidence intervals \( R_{\beta_{2,i}} \) analogously to (22).

### 3.2.2 Confidence interval from bootstrap samples

Alternatively, standard deviation and confidence interval may be calculated from \( B \) bootstrap estimates obtained from both parametric and nonparametric bootstrap methods [5], as described next. Let \( \gamma_i = (\gamma_{1,i,1}, \ldots, \gamma_{1,i,J_i}, \ldots, \gamma_{C,i,1}, \ldots, \gamma_{C,i,J_i}) \in \mathbb{R}^{CJ_i} \). Moreover, let \( \mu_i = (a_1 1_{J_i}, \ldots, a_C 1_{J_i}) + \gamma_i \), and let
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\[
\Sigma_i = \sigma_i^2 \text{diag}(1_{J_i}, k_i^2 1_{J_i}, \ldots, k_i^2 1_{J_i}) \text{ be a diagonal } (C_i J_i) \times (C_i J_i) \text{ matrix where } 1_n \text{ is a vector of ones of length } n. \text{ Now, with independent Gaussian noise } (9), \text{ model } (7) \text{ can be written as }
\]

\[
y_i \in N (\mu_i, \Sigma_i); \forall i. \tag{24}
\]

First, we consider the parametric bootstrap method where the gene vectors \( \mu_i \) and the error terms \( \{ \varepsilon_{c,i,j} \}_{c,j} \) are resampled using the estimated parameters. Let \( \hat{\mu}_i \) and \( \hat{\Sigma}_i \) be defined in analogue to the above but with estimates \( \{ \hat{\alpha}_c \}, \hat{\gamma}_i, \hat{\sigma}_i \), and \( \{ \hat{k}_c \} \) replacing the true parameters. A parametric bootstrap method is then

\[
y_i^* \in N (\hat{\mu}_i, \hat{\Sigma}_i); \forall i. \tag{25}
\]

With this approach, it is possible to obtain confidence intervals for the structural as well as the gene-specific incidental parameters. Second, we consider the nonparametric method where we resample from genes;

\[
y_i^* = y_{u_i}; \forall i \tag{26}
\]

where \( \{ u_i \} \) are independent and uniformly distributed indices on \( i = 1 \ldots, I \). Because \( J_i \) is typically small, it is not reasonable to bootstrap within genes this way, that is to resample from replicates. For instance, with \( J_i = 2 \) half of the samples would contain genes with identical replicated signals making it impossible to estimate \( \beta_{c,i} \) and \( \{ \gamma_{1,i,j} \} \). The nonparametric method (26) is somewhat simpler to implement, especially in a distributed computational environment since it does not require parameter estimates in order to resample. However, contrary to the parametric approach, it only allows us to calculate confidence intervals for the structural parameters.

Continuing, we estimate the standard deviation both by the sample standard deviation, but also by the median absolute deviation (MAD) of the bootstrap estimates. An \( \alpha \)-level percentile confidence interval \( \tilde{R}_{\sigma_c} \) for bias parameter \( \alpha_c \) in channel \( c \) is calculated from the bootstrap estimates \( \{ \hat{\alpha}_c^* \} \) as

\[
\tilde{R}_{\sigma_c} = (\hat{\alpha}^*_{c,(i_1)}, \hat{\alpha}^*_{c,(i_2)}) \tag{27}
\]

where \( \hat{\alpha}^*_{c,(i)} \) is the \( i \)th ranked \( \hat{\alpha}^* \), \( i_1 = [B\alpha/2] \) and \( i_2 = [B(1 - \alpha/2)] \)

with \( \alpha \in [0,1] \). A bias-corrected version of this is where \( i_1 = [B\rho(\alpha/2)] \)
and \( i_2 = \lfloor B \rho(1 - \alpha/2) \rfloor \) with quantiles \( \rho(\alpha) = \Phi(2\lambda + \lambda_1 - \alpha) \) and \( \lambda = \Phi^{-1}(P^*(\hat{a}_c^* \leq \hat{a}_c)) \). We estimate \( P^*(\hat{a}_c^* \leq \hat{a}_c) \) by the proportion of bootstrap estimates \( \hat{a}_c^* \) that are less or equal to the original estimate \( \hat{a}_c \).

Similar to the above, we can calculate a confidence interval \( \tilde{R}_{i\beta_c,i} \) for each (relative) gene-expression level \( \beta_{c,i} \); \( c = 2, \ldots, C; i = 1, \ldots, I \). The confidence interval for gene \( i \) in channel \( c \) from bootstrap estimates is

\[
\tilde{R}_{i\beta_c,i} = (\hat{\beta}_{c,i}^*(i_1); \hat{\beta}_{c,i}^*(i_2))
\]

with notations analogue to the above. Percentile or bias-corrected estimates can be obtain by choosing \( i_1 \) and \( i_2 \) as before.

### 3.2.3 Average gene-expression coverage

Given confidence-interval estimates \( \{\tilde{R}_{i\beta_c,i}\} \) and the true \( \{\beta_{c,i}\} \) parameters, we calculate the average coverage of the relative gene-expression levels in channel \( c \) as

\[
\hat{C}_{\beta_c} = \frac{1}{I} \sum_{i=1}^{I} \mathbb{I}(\beta_{c,i} \in \tilde{R}_{i\beta_c,i})
\]

where \( \mathbb{I}(\cdot) \) is the indicator function. Note, to calculate the degree of coverage \( \hat{C}_{\beta_c,i} \) for each individual gene, and similarly for \( a_c \), another, say, \( T - 1 \) sets of samples, which each has \( B \) bootstrap estimates, have to be generated. Due to time constraints, this was not feasible.

### 4 Data

All data in this report is based on simulated data as we wish to study the properties of the above estimates for the different model and parameter setups. Real-data experiments are planned, but not covered in this theoretical study.

### 4.1 Simulated models

We simulate microarray data from the affine model (3) with independent error terms \( \varepsilon_{c,i,j} \sim N(0, \sigma_{c,i,j}^2) \) and, in order to investigate the estimator’s robustness.

---

17Since \( \beta_{1,i} = 1 \) by definition, there is no confidence interval for \( c = 1 \).
4. Data

toward model misspecification, with two different error structures:

\[
\begin{align*}
\sigma_{c,i,j} &= \tilde{z}_i k_c \sigma \\
\sigma_{c,i,j} &= b_{i,j} b_c z_{c,i} k_c \sigma
\end{align*}
\]

(30) (31)

for all channels, genes and replicates where \(\tilde{z}_i = \sum_{c=1}^{C} z_{c,i} \), \(b_c\), \(b_{i,j}\), and \(z_{c,i}\) are as in (3) with a proportionality constant \(k_c \sigma\) that depends on the channel \(c\) and satisfies the constraint \(k_1 = 1\). Note that the former error structure (30) is a special case of (4)-(5) such that the standard deviation of the noise is proportional to the amount of DNA hybridized to the dilution series as a whole, cf. (1). The latter error structure is such the standard deviation is proportional to the probe concentration as well as the channel-specific amount of hybridization, cf. [13, 12, 6].

For both error structures we study two different types of dilution series, one with uniformly and one with log-uniformly distributed concentrations:

\[
\begin{align*}
b_{i,j} &= j/J_i \\
b_{i,j} &= 2^{(j - J_i)/J_i}
\end{align*}
\]

(32) (33)

such that \(0 < b_{i,j} \leq 1\) and \(b_{i,J} = 1\) for all genes.

Without loss of generality, we have chosen to study the above for setups with two channels, the same number of replicates for all genes, the same biases, the same scale parameters, and the same relative standard deviation for all channels. More precisely, \(C = 2, J_i = J; \forall i, a = (20, 70), b_c = 1,\) and \(k_c = 1; \forall c.\) For error structure (30) we use \(\sigma = 1/16\) and for (31) we use \(\sigma = 1/4.\)

4.2 Simulation of “expressions level”

The expression levels \(\{z_{c,i}\}_{c,i}\) are simulated as follows. First we sample from \(\log_2 \gamma_{1,i} \in \Gamma(1, 1/3)\) with shape parameter 1 and rate parameter 1/3. Samples for which \(\log_2 \gamma_{1,i} > 16\) are censored to 16. To simulate fold changes we sample from \(\log_2 \beta_{c,i} \in N(0, 2^2)\) and assign \(z_{c,i} \leftarrow \beta_{c,i} \tilde{\gamma}_{1,i} + \delta; \forall c\) where \(\delta = 100\) is a small constant in order to avoid gene expressions close to zero.
4.3 Simulated array setups

For the nonparametric bootstrap method, we have simulated data from the model setups and obtained original and \( B = 500 \) bootstrap estimates of all structural and incidental parameters for most combinations of \( I = 100, 300, 600, 1000, 1500 \) and \( J = 2, 3, 4, 5, 10, 30, 100 \). For the parametric bootstrap method, we choose slightly different sets of \( I \) and \( J \), but of the same order.

5 Results

With the data sets in hand, we have studied how the standard deviation of the bias-parameter estimates vary i) with varying number of replicates \( J \) and fixed number of genes \( I \), ii) with varying \( I \) and fixed \( J \), and iii) with different combinations of \( I \) and \( J \) such that the total number of probes, \( J_{\Sigma} = IJ \), is fixed. We also compare the estimates obtained from arrays with log\(_2\)-uniformly and uniformly distributed dilution series. Similarly, we investigate the properties of the relative expression estimates \( \{\beta_{c,i}\}_{c,i} \) and the average coverage of these. We end this section with a visual comparison of non-calibrated and calibrated probe signals, as well as a comparison between estimated and true relative gene expression ratios. All analysis was done with data simulated under the correct model (30) and the misspecified model (31).

5.1 Varying number of replicates

5.1.1 Correct model

*Bias-parameter estimates:* For a constant number of genes, the standard deviation of the bias-parameter estimates decreases as shown in the left graphs of Figure 2. For not too small number of replicates (or genes), the standard deviation estimates based on the *estimated* Fisher information, which is fast to calculate, are very close to the estimates based on computationally intensive bootstrap simulations. However, for a typical microarray experiment where the number of replicates is small (here about \( J < 5 \) or so depending on \( I \)), the former tend to be too small. See also Figure 5. We believe this is because the asymptotic normality assumption does not hold for small \( J \) as illustrated by the density plot in the middle graph at the top. Moreover, for large number of genes, the standard deviation calculated from the *true* Fisher information is close the estimated ditto. For small number
5. Results

Figure 2: Standard deviation of the bias-parameter estimate $\hat{a}_1$ in model (30) for $C = 2$ and $\log_2$-uniformly distributed replicates as a function of the number of replicates $J$ and as a function of the number of genes $I$. All estimates are based on the parametric bootstrap method. Top left: The standard deviation for $I = 96$ and $J = 2, \ldots, 56$. The solid and the dashed black curves (with asterisks) are the sample standard deviation and MAD of the bootstrap estimates, respectively. The solid orange and blue curves (with circles) are based on the observed and the true Fisher information, respectively. Top middle: Kernel density estimate illustrating the distribution of the bootstrap estimates for $I = 96$ and $J = 2$. The solid and dashed vertical lines are the estimated and the true bias parameter, respectively. The horizontal boxes are four different 95% confidence intervals. The bottom two are based on the percentile and bias-corrected bootstrap estimates according to (27). The two on top are based on the true and the estimated Fisher information, where the latter is hardly visible. Top right: As in the top left graph, but for $J = 6$ and $I = 96, \ldots, 1152$. Bottom left: The standard deviation for $I = 432$ and $J = 2, \ldots, 56$. Bottom middle: Distribution of bootstrap estimates for $I = 432$ and $J = 4$. Bottom right: The standard deviation for $J = 32$ and $I = 96, \ldots, 1152$.

of genes the former tends to be underestimated. See also Figure 2.

Relative-expression estimates: We have calculated the average coverage of the true relative gene-expression levels $\{\beta_{2,i}\}$ for the percentile and bias-corrected bootstrap confidence intervals estimated for each simulation set. The average coverage increases with the number of replicates as depicted in the left graph of Figure 3. The coverage of uniformly and $\log_2$-uniformly distributed probe concentrations are very close to each other.
5.1.2 Misspecified model

Bias-parameter estimates: For the more general model (31), the standard deviation of the parameter estimates based on the estimated Fisher information are still close to those calculated based on bootstrap estimates. See left and right graphs of Figure 4, which depict estimates based on parametric bootstrap samples. However, the standard deviation calculated from the true Fisher information is under-estimated with several orders of magnitude, especially for large number of replicates. This discrepancy tends to be large the more replicates there are as in Figure 4. The graphs shown are for log₂-uniformly distributed dilution series. For uniform dilution series the corresponding discrepancy (not shown) increases less rapidly with \( J \), but it still increases (on the log-scale).

Relative-expression estimates: For the misspecified model with log₂-uniformly distributed dilution series, the average coverage decreases with the number of replicates as illustrated in the right graph of Figure 3. This is because the standard deviation decreases faster than the bias of the relative gene-expression estimates. See also Figure 7. For uniformly distributed dilution series, the average coverage is stable.
or increases slightly.

5.2 Varying number of genes

Bias-parameter estimates: The standard deviation of the bias-parameter estimates for a constant number of replicates decreases with the number of genes as depicted in the right graphs of Figure 2. Observe that the curves in the left panels are smoother than the ones in the right panels. This is because the former ones are estimated with identical sets of expressions levels \( \{(\beta_i, \gamma_{1,i})\}_i \) whereas the latter ones are not. Note that since the nonparametric bootstrap estimates resample gene by gene, the corresponding graphs in that case are much smoother (not shown). The results are similar for the misspecified model.

Relative-expression estimates: We find no trend in coverage with respect to the number of genes. Neither for the correct nor the misspecified model.

5.3 Same number of probes

From the graphs in Figure 5, which summarizes the above results, we conclude that the standard deviation of the bias parameters is roughly constant for constant \( IJ \), except when the number of replicates \( J \) is small. In that case the large sample approximation is unreliable and the true standard error is larger. We have chosen to display the standard deviation estimated based on the nonparametric bootstrap samples because, as described above, these are less dependent on the simulated set of gene expressions \( \{\beta_{c,i}\}_{c,i} \). Estimates from the parametric bootstrap show a similar but more noisy pattern (not shown). The same behavior is observed (not shown) for the misspecified model.

5.4 Type of replicates

5.4.1 Correct model

Bias-parameter estimates: When it comes to minimizing the uncertainty of the bias-parameter estimates, \( \log_2 \)-uniformly distributed probe concentrations are better than uniformly distributed ones. See left graph in Figure 6, which shows the relative standard deviations, that is, the standard deviation estimates for the \( \log_2 \)-uniform replicates divided by ditto for the uniform replicates. Note that the
second factor in (21) goes to 1/4 when \( J \to \infty \) for \( \log_2 \)-uniform replicates (33), and to 1 for uniform replicates (32). Thus, with everything else the same, the relative standard error of these two replicate types is 1/2 when \( J \) is large. By definition both types of replicates are identical for \( J = 2 \), cf. (32)-(33). No corresponding trend is observed when varying the number of genes.

5.4.2 Misspecified model
A similar comparison for model (31) shows that the relative decrease when the number of replicates grows is even faster. See right graph in Figure 6.

5.5 Estimated gene log-ratios
Define the estimated gene log-ratios and the true gene log-intensities for channel \( c \) and channel 1 as

\[
\hat{M}_i = \log_2 \hat{\beta}_{c,i}, \\
A_i = \frac{1}{2} \log_2(z_{c,i} z_{1,i}),
\]

(cf. [14]). Apart from the constant \( \log_2(b_c) \), which can be estimated separately, \( \hat{M}_i \) is an estimate of the true log-ratio \( M_i = \log_2(z_{c,i} / z_{1,i}) \) for gene \( i \). In Figure 7, \( \{(A_i, M_i)\}_i \) are plotted for model (30) with \( \log_2 \)-uniformly distributed replicates.
5. Results

Figure 5: The various standard deviations of the bias-parameter estimates for different array layouts such that $IJ$ is constant. The data was simulated under model (30) with log$_2$-uniformly distributed replicates (33). Estimates based on nonparametric (26) bootstrap estimates from channel two is shown. Left: $IJ = 3000$. Right: $IJ = 6000$. Legend: 'sd*' is the standard deviation and 'mad*' is the median absolute deviation of the bootstrap estimates. 'sdI' and 'sdIˆ' are the standard deviations calculated from the true and the estimated Fisher information, respectively.

Figure 6: Relative standard deviation between log-uniform and uniform replicates of the estimate of the bias parameter $a_1$ for $I = 432$ and $J = 2, \ldots, 32$. Left: Data simulated from model (30). Right: Data simulated from model (31). All estimates are from the nonparametric bootstrap samples.
and $c = 2$, $I = 96$, and for $J = 2$ (right graphs) and $J = 16$ (right graphs). The vertical lines with whiskers depict the 99% bias-corrected confidence intervals obtained from the parameteric bootstrap samples. From these graphs, it is clear that with the two noise structures chosen, the highly differentially expressed genes have a larger standard deviation than the non-differentially expressed genes. Indeed, ignoring the remainder term of (23) and applying Gauss approximation, it follows that

$$
V[\hat{M}_i] \approx \frac{V[\hat{\beta}_{2,i}]}{(\ln 2)^2/2} \approx \frac{k_2 \sigma^2}{(\ln 2)^2} \left( \frac{\beta_{2,i}}{k_2} + \frac{k_2}{\beta_{2,i}} \right) \left( \sum_{j=1}^{J_i} b_{ij}^2 \right)^{-1}
$$

with $\ln$ the base $e$ logarithm. The right-hand side is a symmetric function of $M_i = \log_2(\beta_{2,i})$ around $\log_2(k_2)$, where it has its minimum. An estimate of $\beta_{c,i}$ may be negative. This is especially true for highly differentially-expressed genes for which the probe signals in one of the channels are weak, cf. Figure 1. The same is true for the upper and lower limits of the estimated confidence intervals. Since non-positive values are not defined on the logarithm scale, we censor these to equal $\infty$ or $-\infty$ in the graphs.

5.6 Calibrated probe signals

To illustrate what non-linear effects affine transformations have on the observed relative expression levels $\{y_{c,i,j}/y_{1,i,j}\}_{i,j}$, and how these artifacts are removed by calibration method (19), we calculate the probe log-ratios and log-intensities

$$
M_{i,j} = \log_2 \frac{y_{c,i,j}}{y_{1,i,j}}
$$

$$
A_{i,j} = \frac{1}{2} \log_2(y_{c,i,j}y_{1,i,j})
$$

for all probes $(i, j)$ and similarly for the calibrated signals $\{\hat{x}_{c,i,j}\}_{c,i,j}$. As depicted by graphs in Figure 8, the intensity-dependent effects in the raw signals are to a large extent removed by the calibration method. This is true for data both from (30) and (31).

6 Discussion

Calibrated probe signals $\{\hat{x}_{c,i,j}\}_{c,i,j}$ may become non-positive due to uncertainty in the bias-parameter estimates, but also due to the existence of additive noise.
Figure 7: Example of true and estimated gene log-ratio versus true log-intensities for $C = 2$, $I = 96$, and varying $J$. Left graphs: $J = 2$. Right graphs: $J = 16$. The crosses are the true log-ratios and the thick horizontal lines are the estimated log-ratios with 99% bias-corrected confidence intervals. Top row: Data simulated from (30) with log$_2$-uniformly distributed replicates. Bottom row: Same as above, but simulated from (31).

Our model does not try to correct for this. To deal with such problems, see for instance [8]. However, note that the relative expression ratios $\{\beta_{c,i}\}_{c,i}$ are estimated in a way which makes the problem of non-positive signals somewhat smaller, although $\hat{M}_i$ still might be undefined for some genes without further restrictions on the $\beta_{c,i}$-parameters.

We have used the calibrated signals $\hat{x}_{c,i,j}$ for estimating probe-specific log-ratios. As an alternative to (34), they may be used for estimating gene-specific log-ratios too, for instance through

$$\hat{M}_i = \log_2 \frac{\sum_{j=1}^{J_i} w_{i,j} \hat{x}_{c,i,j}}{\sum_{j=1}^{J_i} w_{i,j} \hat{x}_{1,i,j}},$$

(38)

where $w_{i,j}$ are weights chosen by the user, for instance $w_{i,j} = 1$ or
Figure 8: Example of probe log-ratio versus log-intensities before (left graphs) and after (right graphs) calibration. Scale parameter $b_2$ is considered to be known, if not, the left scatter plots are shifted downward. A random set of dilutions series has been highlighted in order to illustrate the fact that the non-linear intensity-dependent artifacts are removed when applying the affine calibration method. Top row: Data show simulated from model (30) with $I = 432$ genes, $J = 4$ and log$_2$-uniform replicates. Bottom row: Data show simulated from model (31) for the same array setup.

$$w_{i,j} = w(b_{i,j}),$$ in case the latter are known.

Investigating (35), an interesting alternative error model to simulate from would be the one where, instead of the arithmetical, the geometrical mean of the hybridized dilution concentrations is used, that is, $\tilde{z}_i = \sqrt{z_{1,i} z_{2,i}}$, cf. (30).

It may even be the case that it is possible that our proposed model can be used for spotted microarrays that contain technical replicates that are not diluted. The reason why this may work is that the deposition of spots onto the array surface is imperfect and that different spots may dry differently [4]. This could result in different probe concentrations $\{b_{i,j}\}_j$. Even donut effects may be useful.
7 Conclusions

We have studied a stochastic affine model for scanned microarray data with dilution series and/or spike-ins. Using maximum likelihood techniques estimates of the relative expression levels for each dilution series (gene) follow immediately as well as calibrated signals for all probes. It is important to control the spot concentrations such that no quenching [11] takes place. The model assumes noise that is constant for all probe replicates, but may vary freely between genes. One might generalize the ML estimates to models like (31), although these will be computationally more intensive. However, our investigations show that model (30) is fairly robust against model misspecification. Even though the average coverage (29) of the relative gene-expression levels is too low for the misspecified model, we still see, from the right subplots of Figure 7, that the variability of the estimated log-ratios decreases with increasing \( J \) and is satisfactory for large \( J \). It is just the length of confidence intervals for \( \beta_{c,i} \) and \( \hat{M}_{i} \) that is too small under model misspecification.

We find that dilution series where the probe concentrations are \( \log_2 \)-uniformly distributed are favorable to the ones that are uniformly distributed, although the average coverage of the log-ratios under the misspecified model is worse for the former; the standard deviation of the estimates is smaller for the \( \log_2 \)-uniform than uniform replicates. Note that the former design is also much easier to implement in practice.

To conclude, we suggest that microarray experiments are designed with multiple dilution series for different genes that are likely to be differentially expressed. Although only series with just two replicates are required, we recommend to use more replicates in order to improve the accuracy of the estimates and the calibration as a whole. The method can easily be applied to subgroups of probes, especially spatially disjoint groups such as print-tip groups, which brings up the question on how to optimally position the replicates relative to each other on the microarray. It is beyond this article to give a thorough answer, but it will be a trade-off between accuracy of parameter estimates and size of the regions the estimates should be based on, which in turn depends on the spatial structure of the artifacts.

All analysis was carried out using R [10, 7]. The estimation and calibration meth-
ods presented in this paper is made available for free in the aroma package [1].
References


A Fisher information

A.1 Fisher information for bias parameters

For simplicity we assume \( C = 2 \) channels. For any two subvectors \( x \) and \( y \) of \( \theta \) and \( \xi_i \) we let \( I_{x,y} = (I_{x,y})_{i,j} \) be the \( |x| \times |y| \) submatrix of \( I \). If \( I_{x,y} = 0 \), \( x \) and \( y \) are orthogonal parameter vectors. It is easy to show that \((a, \beta, \gamma)\) is orthogonal to \((k, \sigma^2)\). From properties of the inverse Fisher information of structural parameters (see for instance [9], Theorem 9.8) it follows that

\[
\frac{1}{(I^{-1})_{a_1,a_1}} = Q_{1,1} - \frac{Q_{1,2}^2}{Q_{2,2}} \tag{39}
\]

\[
\frac{1}{(I^{-1})_{a_2,a_2}} = Q_{2,2} - \frac{Q_{2,1}^2}{Q_{1,1}} \tag{40}
\]

where \( Q = (Q_{i,j}) \) is the \( 2 \times 2 \) matrix

\[
Q = I_{a,a} - I_{a,\beta_1}I_{\beta_1,\beta_1}^{-1}I_{\beta_1,a} = \sum_{i=1}^{l} J_i \rho_i (I_{\beta_i,\beta_i})^{-1}I_{\beta_1,a} \tag{41}
\]

where we define \( \beta_i = (\beta_{2i}, ..., \beta_{Ci}) \), cf. the definition in the “Model” section. One finds that

\[
I_{a,a} = \sum_{i=1}^{l} \frac{J_i}{\sigma_i} \begin{pmatrix} 1 & 0 \\ 0 & k_i^{-2} \end{pmatrix} \tag{42}
\]

and the \( i \)-th term of the right-hand side of (41) equals

\[
(I_{a,\beta_i}I_{a,\gamma_i}) \begin{pmatrix} I_{\beta_i,\beta_i} & I_{\beta_i,\gamma_i} \\ I_{\gamma_i,\beta_i} & I_{\gamma_i,\gamma_i} \end{pmatrix}^{-1} \begin{pmatrix} I_{a,\beta_i}I_{a,\gamma_i} \end{pmatrix}' \tag{43}
\]

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where $'$ denotes matrix transposition. Further,

\[
I_{\alpha\beta_i} = \frac{J_i \gamma_{1,i}}{\sigma_i^2 k_2^2} \begin{pmatrix} 0 \\ 1 \end{pmatrix}
\]  
(44)

\[
I_{\alpha\gamma_{1,i}} = \frac{1}{\sigma_i^2} \left( \begin{array}{c} 1 \\ \beta_{2,i}/k_2^2 \end{array} \right) \mathbf{1}_I
\]  
(45)

\[
I_{\beta_i\beta_i} = \frac{J_i \gamma_{2,1,i}}{\sigma_i^2 k_2^2}
\]  
(46)

\[
I_{\beta_i\gamma_{1,i}} = \frac{\beta_{2,i}}{\sigma_i^2 k_2^2} \gamma_{1,i}
\]  
(47)

\[
I_{\gamma_{1,i}\gamma_{1,i}} = \frac{k_2^2 + \beta_{2,i}^2}{\sigma_i^2 k_2^2} \text{diag}(1, \ldots, 1)
\]  
(48)

with $\bar{\gamma}_{1,i} = \sum_{j=1}^{J_i} \gamma_{1,i,j}/J_i$ and $\gamma_{2,1,i} = \sum_{j=1}^{J_i} \gamma_{2,1,i,j}/J_i$. Inserting (44)-(48) into (43), we find, after some calculations based on inversion formulas for block matrices, that the $i^{th}$ term of the sum in (41) equals

\[
\frac{J_i}{\sigma_i^2} \left( \begin{array}{ccc} 1 - \frac{\beta_{2,i}^2 S_i}{k_2^2 + \beta_{2,i}^2} & \frac{\beta_{2,i} S_i}{k_2^2 + \beta_{2,i}^2} & \frac{\beta_{2,i} S_i}{k_2^2 + \beta_{2,i}^2} \\ \frac{\beta_{2,i} S_i}{k_2^2 + \beta_{2,i}^2} & k_2^{-2} - \frac{S_i^2}{k_2^2 + \beta_{2,i}^2} \end{array} \right)
\]  
(49)

where

\[
S_i = \sum_{j=1}^{J_i} (\gamma_{1,i,j} - \bar{\gamma}_{1,i})^2 / \sum_{j=1}^{J_i} \gamma_{1,i,j}^2 = \sum_{j=1}^{J_i} (b_{i,j} - \bar{b}_i)^2 / \sum_{j=1}^{J_i} b_{i,j}^2
\]  
(50)

Inserting (42) and (49) into (41) we obtain

\[
Q_{1,1} = \sum_{i=1}^{I} \frac{\beta_{2,i}^2}{k_2^2 + \beta_{2,i}^2 \sigma_i^2} S_i = \sum_{i=1}^{I} w_i \beta_{2,i}^2
\]  
(51)

\[
Q_{2,2} = \sum_{i=1}^{I} \frac{1}{k_2^2 + \beta_{2,i}^2 \sigma_i^2} S_i = \sum_{i=1}^{I} w_i
\]  
(52)

\[
Q_{1,2} = \sum_{i=1}^{I} \frac{\beta_{2,i}}{k_2^2 + \beta_{2,i}^2 \sigma_i^2} S_i = \sum_{i=1}^{I} w_i \beta_{2,i}
\]  
(53)

where $w_i$ are gene-specific weights as in (21). Insertion of (51)-(53) into (40) gives (20) after some manipulations.
A.2 Fisher information for relative expression levels

To begin with we assume known $a$ and $C = 2$. Then $\beta_{2,i} = \beta_i$ and $\gamma_{1,i}$ are orthogonal to all other parameters and

$$1/(I^{-1}_{\beta_{2,i},\beta_{2,i}}) = I_{\beta_i,\beta_i} - I_{\beta_i,\gamma_{1,i}} I_{\gamma_{1,i},\gamma_{1,i}} I_{\beta_i,\gamma_{1,i}}.$$  \hspace{1cm} (54)

Inserting (46)-(48) into (54) it follows after some calculations that

$$1/(I^{-1}_{\beta_{2,i},\beta_{2,i}}) = \frac{\sum_{j=1}^{J_i} \gamma_{1,i,j}^2}{\sigma_i^2 (k_2^2 + \beta_{2,i}^2)} = \frac{z_{1,i}^2 \sum_{j=1}^{J_i} b_{1,i,j}^2}{\sigma_i^2 (k_2^2 + \beta_{2,i}^2)}.$$  \hspace{1cm} (55)

A more detailed analysis (not shown here) reveals that additional terms of total size $O(J_\Delta^{-1})$ are added to the right-hand side of (55) when $a$ is unknown, and this establishes (23).
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