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Published in:
Journal of Biomedical Optics

DOI:
10.1117/1.JBO.19.2.025003

2014

Link to publication

Citation for published version (APA):

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Evaluation of composition and mineral structure of callus tissue in rat femoral fracture

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Abstract. Callus formation is a critical step for successful fracture healing. Little is known about the molecular composition and mineral structure of the newly formed tissue in the callus. The aim was to evaluate the feasibility of small angle x-ray scattering (SAXS) to assess mineral structure of callus and cortical bone and if it could provide complementary information with the compositional analyses from Fourier transform infrared (FTIR) microspectroscopy. Femurs of 12 male Sprague–Dawley rats at 9 weeks of age were fractured and fixed with an intramedullary 1.1 mm K-wire. Fractures were treated with the combinations of bone morphogenetic protein-7 and/or zoledronate. Rats were sacrificed after 6 weeks and both femurs were prepared for FTIR and SAXS analysis. Significant differences were found in the molecular composition and mineral structure between the fracture callus, fracture cortex, and control cortex. The degree of mineralization, collagen maturity, and degree of orientation of the mineral plates were lower in the callus tissue than in the cortices. The results indicate the feasibility of SAXS in the investigation of mineral structure of bone fracture callus and provide complementary information with the composition analyzed with FTIR. Moreover, this study contributes to the limited FTIR and SAXS data in the field. © 2014 Society of Photo-Optical Instrumentation Engineers (SPIE) [DOI: 10.1117/1.JBO.19.2.025003]

Keywords: cortical bone; fracture healing; composition; mineral structure; Fourier transform infrared microspectroscopy; small angle x-ray scattering.

1 Introduction

Callus formation is a critical step for successful fracture healing.1 About 5% to 10% of all fractures suffer from delayed healing or lead to a nonunion.2 Bone morphogenetic proteins (BMPs) increase callus formation3 and have been shown to be valuable in the treatment of fracture nonunions. BMPs, however, increase the speed of remodeling and thereby also induce resorption via the RANK pathway. On the other hand, bisphosphonates can be used to reduce the resorption of bone.4 These manipulations of the molecular composition and mineral structure, respectively, of trabecular5 and cortical bone.5,10–13 However, the composition and mineral structure of the callus tissue during fracture healing has received little attention.14–16

Each molecular structure has its own specific IR absorption spectrum. When coupled to a microscope, FTIR imaging microspectroscopy provides a tool for fast measurements of the spatial composition of bone.10,17,18 Several compositional parameters can be calculated from the bone IR spectra, e.g., degree of mineralization, carbonate substitution, collagen maturity, crystallinity, and acid phosphate substitution (APS).19–22

In SAXS, a sample is irradiated by a narrow well-collimated x-ray beam and the intensity of x-rays scattered by the sample at small angles is measured. In bone, the scattering is related to bone mineral structure, i.e., the spatial arrangement of mineral crystals and collagen fibrils. When coupled with an automated scanning stage, spatial maps can be recorded where each pixel contains a two-dimensional (2-D) scattering pattern.13 Several parameters can be determined from the scattering pattern of bone, e.g., the mineral plate thickness, predominant orientation, and degree of orientation.8,13,23,24 Spatial distributions of these parameters provide a good insight to the variation in mineral structure in cortical bone and newly formed callus tissue. Different methods have been used to determine the mineral plate thickness. Fratzl et al. proposed a method in which a mineral phase fraction of 50% is assumed.23,25 This assumption might not be valid for, e.g., newly formed bone where the degree of mineralization may vary substantially. Another approach introduced by Bünker et al. evaluates the mineral plate thickness through an iterative curve-fitting method where no assumptions of the mineral phase fraction are needed.13 The latter was used in this study.

This study used FTIR imaging microspectroscopy and scanning SAXS to assess the molecular composition and mineral structure of newly formed callus tissue, the adjacent cortical
bone, and the nonfractured contralateral cortical bone, during long bone fracture healing in a rat nonunion model. The first aim was to evaluate the feasibility of SAXS to assess the mineral structure of callus tissue and if it could provide complementary information to the compositional analyses from FTIR. The subsequent aim was to study the differences in molecular composition and mineral structure among fracture callus, fracture cortex, and control cortex.

2 Materials and Methods

2.1 Experimental Overview

Twelve male Sprague-Dawley rats at 9 weeks of age were anaesthetized with ketamine HCl (75 mg/mL, Parnell Laboratories, Rosebery, Australia) and xylazine (10 mg/mL, Ilium, Smithfield, Australia).25 The right femurs of the rats were osteotomized, stripped of periosteum and muscle, and fixed with an intramedullary 1.1 mm Kirschner wire. BMP-7 was placed locally around the fracture and zoledronate (ZO) or saline (NaCl) was injected after 2 weeks. Thus, four groups were created (A) NaCl, (B) BMP-7 + NaCl, (C) BMP-7 + ZO, and (D) ZO. After the operation, the rats received subcutaneous physiologic saline and buprenorphine (Temgesic, Reckitt and Coleman, Hull, UK) at 0.05 mg/kg twice a day. Rats were sacrificed after 6 weeks and both femurs, fractured (right), and control (left) were stripped of the soft tissue, defatted, and dehydrated in ascending series of ethanol solutions and embedded in polymethylmethacrylate (PMMA). The experimental protocol was approved by the local animal ethics committee. The experimental model results in 52% nonunions when the femur is left untreated.25

2.2 Imaging of Callus Size

The femurs from the fracture side were imaged with a micro-CT system (Skyscan 1172, Aartselaar, Belgium) to visualize the fracture callus and cortex in each animal (Fig. 1). Images were acquired with an isotropic voxel size of 36 μm using 100 kV, 100 μA with a 0.5 mm aluminum filter, and 10 repeated scans. Image reconstruction was performed (NRecon, Skyscan, v. 1.5.1.4) by correcting for ring artifacts and beam hardening. Scans were created (A) NaCl, (B) BMP-7 + NaCl, (C) BMP-7 + ZO, and (D) ZO. After the operation, the rats received subcutaneous physiologic saline and buprenorphine (Temgesic, Reckitt and Coleman, Hull, UK) at 0.05 mg/kg twice a day. Rats were sacrificed after 6 weeks and both femurs, fractured (right), and control (left) were stripped of the soft tissue, defatted, and dehydrated in ascending series of ethanol solutions and embedded in polymethylmethacrylate (PMMA). The experimental protocol was approved by the local animal ethics committee. The experimental model results in 52% nonunions when the femur is left untreated.25

2.3 Small Angle x-ray Scattering

From the PMMA embedded specimens, 300 μm sections were sawed (EXAKT 400 CS, Cutting Grinding System, Hamburg, Germany). Two samples from each treatment group and one section from each sample were measured and analyzed. SAXS measurements were conducted at the 1911-SAXS beamline at the 1.5 GeV ring (MAX II) of the MAX IV Laboratory (Lund University, Lund, Sweden).27 The wavelength of the monochromatic radiation obtained from a Si(111) crystal was 0.91 Å and the size of the collimated synchroton x-ray beam at the sample position was approximately 0.2 × 0.2 mm². The used detector was a MarCCD (Rayonix, L.L.C.) with a 165 mm active area and 79 μm pixel size. It was placed at 1911 mm behind the sample and the exposure time to collect each SAXS pattern was 5 s. The q-range measured at each measurement point was 0.01 to 0.30 Å⁻¹. Bone sections were mounted in a sample holder which was placed in a motorized x–y scanning stage to map the sample with a step size of 0.2 mm in both directions. About 23 mm² areas of the sections including the fracture callus and cortex and 10 mm² areas of the control cortex were measured [Fig. 2(b)].

From the anisotropic (noncircular symmetric) 2-D SAXS intensity pattern, I(q, θ), the size, shape, and orientation of the mineral crystals were analyzed for each measurement point [Fig. 3(a)]. Analysis of SAXS data were done by averaging the scattering pattern over 360 deg azimuthal range8,13,23,24 after masking of the beamstop and the area outside the detector [Fig. 3(a)]. This results in a one-dimensional (1-D) scattering pattern I(q). From this pattern, the mineral plate thickness was evaluated following the approach suggested by Bünger et al.13 Shortly, it is based on a curve fitting where the mineral crystals are assumed to be plates with a finite thickness, T, in 1-D and infinite size in the other 2-D. The scattering from one plate is

\[
P(q) = \frac{1}{q^2} \left| \frac{\sin(qT/2)}{qT/2} \right|^2 .
\]  

Here, \( q = \left(4\pi/\lambda\right) \sin(\theta/2) \) is the scattering vector magnitude, where \( \lambda \) is the x-ray wave length and \( \theta \) is the scattering angle. The variation of the thickness of the mineral plates is taken into account by assuming that the thickness variation follows a Schultze-Zimm distribution \( D(T, T_{av}) \). The average scattering then reads

\[
P_{av}(q) = \frac{\int_0^\infty T^2 P(q) D(T, T_{av})dT}{\int_0^\infty T^2 D(T, T_{av})dT} .
\]  

Equation (1) represents the single particle scattering function. In bone, the mineral content is high and we also need to take into account the spatial arrangement of mineral platelets, through a structure factor. Bünger et al.13 suggested a structure factor that includes both short range and long range particle correlations. To describe the short range repulsive interactions, they considered the random phase approximation (RPA), where the effective RPA structure factor can be written as

\[
S_{RPA}(q) = \frac{1}{1 + \nu P_{av}(q)} ,
\]  

where ν is an adjustable parameter depends on the strength of the interactions and that typically increases with increasing concentrations. In addition, the increasing scattering intensity

![Image](https://via.placeholder.com/150)

Fig. 1 Micro-CT images reveal the state of healing of the bone fractures after 6 weeks in (a) NaCl, (b) BMP-7 + NaCl, (c) BMP-7 + ZO, and (d) ZO treatment groups. The images were used to qualitatively assess callus size and healing state.
of the section and 1 means that all mineral crystals are aligned perfectly in the same directions. These parameters were calculated for each measurement point (Fig. 4). A threshold of the integrated intensity of the 1-D scattering pattern $I(q)$ was used to mask the background (PMMA) and each analyzed area contained the whole measured cortex or callus.

### 2.4 Fourier Transform Infrared Microspectroscopy

Three-micrometers thick longitudinal sections of bone were cut (Polycut S, Reichert-Jung, Germany) from the PMMA plugs, and placed on ZnSe windows. Three samples of each treatment group and two sections of each sample from the control cortex, fracture cortex, and callus tissue were measured and analyzed. Analysis of bone composition was conducted with a PerkinElmer Instrument (Spectrum Spotlight 300, Perkin Elmer Inc., Wellesley, Massachusetts) in transmission mode. A 25 μm spatial resolution and spectral resolution of 4 cm$^{-1}$ with eight repeated scans per pixels were used. The background spectrum was recorded on a clean area of the ZnSe window by using the same measurement parameters but with an average of 75 scans. Data were collected at the wavenumber range of 2000 to 800 cm$^{-1}$. Areal measurements (~30 mm$^2$) were performed on the fractured femurs, including both the cortex and callus [Fig. 2(c)], and on the control cortex of the intact femur. In addition, one areal measurement of only PMMA was measured and averaged. The bone spectra were normalized by using the averaged PMMA spectrum. Subsequently, the PMMA spectrum was subtracted from the bone spectra.

From the preprocessed bone spectra, the areas of linearly baseline corrected amide I (1720 to 1585 cm$^{-1}$), phosphate...
(1200 to 900 cm\(^{-1}\)), and carbonate (890 to 850 cm\(^{-1}\)) peaks were calculated.\(^{17,19-21}\) These peak areas were used to determine the mineral/matrix (phosphate/amide I) and carbonate/phosphate ratios. Mineral/matrix ratio is an indicator of the degree of mineralization, whereas carbonate/phosphate ratio describes the carbonate substitution.\(^{19}\) Collagen maturity and crystallinity of the hydroxyapatite (HA) crystals were evaluated through second derivative peak fitting. The ratio of areas of the subpeaks under the amide I peak at 1660 and 1690 cm\(^{-1}\) was used to describe the collagen maturity (collagen cross-linking ratio—XLR),\(^{21}\) whereas the ratio of areas of the subpeaks at 1030 and 1020 cm\(^{-1}\) under the phosphate peak was used as an indicator of crystallinity,\(^{20}\) i.e., the size and perfection of HA crystals. APS, which is associated with new mineral deposition, was determined as a ratio of intensities at 1127 and 1096 cm\(^{-1}\).\(^{22}\) The total absorption of the background corrected (PMMA subtracted) spectra was used for masking out the bone and each analyzed area contained the whole measured cortex or callus. All preprocessing and analyses of the FTIR and SAXS data were performed using MATLAB (MATLAB R2011b, The MathWorks, Inc., Natick, Massachusetts).

### 2.5 Statistics

Micro-CT images were only evaluated qualitatively due to the small number of samples per group. Wilcoxon signed rank test was used to compare the molecular composition and mineral structure between paired samples (each rat) from the control cortex, fracture cortex, and fracture callus. For this test, all the rats were pooled into one group. Paired \(t\)-test was performed by comparing all the data points from the callus tissue and the cortices in the same animal.\(^{30}\)

#### 3 Results

Based on the micro-CT images, only small amounts of callus tissue were found in the NaCl group. Some mineralized callus tissue was forming between the fracture ends but union was not completed [Fig. 1(a)]. All fractures in the BMP-7 groups were judged to be completely healed based on the micro-CT images [Figs. 1(b) and 1(c)]. Callus size was larger in BMP + ZO group than in other groups (Fig. 1).

Based on the FTIR data with all samples pooled, the mineral/matrix ratio [Fig. 5(a)] and XLR [Fig. 5(c)] were significantly lower in fracture callus than in fracture cortex (\(p < 0.01\) and \(p < 0.05\), respectively) and control cortex (\(p < 0.05\) and \(p < 0.01\), respectively), whereas crystallinity [Fig. 5(d)] of the control cortex was significantly lower than that in the fracture cortex and callus (\(p < 0.01\) in both). APS [Fig. 5(e)] was significantly higher in the fracture callus compared to control and fracture cortices (\(p < 0.05\) and \(p < 0.01\), respectively).

When comparing all data points within one animal in each sample, the mineral/matrix ratio was lower in the callus tissue than in the cortices (\(p < 0.01\)) [Table 1 and Fig. 5(a)]. Carbonate/phosphate ratio was significantly higher in callus tissue compared to cortices in all samples in the NaCl + BMP-7 and ZO groups (\(p < 0.01\)) [Table 1 and Fig. 5(b)], but not in the other two groups. XLR was significantly lower in callus tissue compared to cortices in all the individual samples from all treatment groups [Table 1 and Fig. 5(c)]. Significantly higher crystallinity values were observed in fracture callus compared to cortices in the NaCl and ZO groups (\(p < 0.01\)), but not in the BMP-7 or BMP-7 + ZO treated samples [Table 1 and Fig. 5(d)]. APS was significantly higher in callus tissue than in fracture cortex in all samples from all treatment groups (\(p < 0.01\)) [Table 1 and Fig. 5(e)].

![Fig. 5](http://biomedicaloptics.spiedigitallibrary.org/) FTIR compositional analyses of (a) mineral/matrix ratio, (b) carbonate/phosphate ratio, (c) collagen cross-linking ratio, (d) crystallinity, and (e) acid phosphate substitution in control cortex, fracture cortex, and fracture callus for all samples together and in different treatment groups. Nonparametric Wilcoxon signed rank test for all samples: *\(p < 0.01\) and **\(p < 0.05\).
BMP-7 was significantly higher in control cortex than in callus tissue fracture cortex were observed. The mineral plate thickness samples, no systematic differences between callus tissue and fracture cortex and callus (degree of orientation was higher in control cortex than in the whereas the orientation was more random in the calluses. Crystals in the cortices was clearly along the long bone axis, whereas the orientation was more random in the calluses. Degree of orientation was higher in control cortex than in the fracture cortex and callus (p < 0.05 in both) [Fig. 6(b)]. The predominant orientation of the mineral crystals in the cortices was clearly along the long bone axis, whereas the orientation was more random in the calluses. Degree of orientation was higher in control cortex than in the fracture cortex and callus (p < 0.05 in both) [Fig. 6(c)].

When comparing all data points within one animal, the mineral plate thickness was significantly higher in callus tissue than in the fracture cortex in BMP-7 + ZO treated groups [Table 1 and Fig. 6(a)]. In NaCl and NaCl + BMP-7 treated samples, no systematic differences between callus tissue and fracture cortex were observed. The mineral plate thickness was significantly higher in control cortex than in callus tissue in samples in NaCl and ZO groups. Significantly higher predominant orientation values were evident in callus tissue compared to cortices in all samples (p < 0.01) [Table 1 and Fig. 6(b)]. Additionally, the degree of orientation was significantly higher in cortices than in callus tissue (p < 0.01) in all samples regardless of the treatment [Table 1 and Fig. 6(c)].

### Table 1 Comparison of composition and mineral structure parameters in callus tissue and fracture cortex when comparing all data points within one animal. Significantly decreased (↓), increased (↑), or no systematic change (—) in values from all samples are indicated, when callus tissue is compared to fracture cortex tissue. Alterations in mineral/matrix ratio (MM), carbonate/phosphate ratio (C/P), collagen maturity (XLR), crystallinity (Cryst), acid phosphate substitution (APS), mineral plate thickness (T), orientation, and degree of orientation (DoO) are indicated.

<table>
<thead>
<tr>
<th>Group</th>
<th>M/M</th>
<th>C/P</th>
<th>XLR</th>
<th>Cryst</th>
<th>APS</th>
<th>T</th>
<th>Orientation</th>
<th>DoO</th>
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<tbody>
<tr>
<td>NaCl</td>
<td>↓</td>
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<tr>
<td>BMP7 + NaCl</td>
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<td>↑</td>
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<tr>
<td>BMP7 + ZO</td>
<td>↓</td>
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<td>ZO</td>
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</table>

Based on the SAXS data with all samples pooled, the fracture cortex had a lower mineral plate thickness than the control cortex and the callus tissue [Fig. 6(a)]. The difference between the fracture cortex and callus tissue was significant (p < 0.05). The predominant orientation was significantly different in the callus tissue compared to the fracture and control cortices (p < 0.05, in both) [Fig. 6(b)]. The predominant orientation of the mineral crystals in the cortices was clearly along the long bone axis, whereas the orientation was more random in the calluses. Degree of orientation was higher in control cortex than in the fracture cortex and callus (p < 0.05 in both) [Fig. 6(c)].

When comparing all data points within one animal, the mineral plate thickness was significantly higher in callus tissue than in the fracture cortex in BMP-7 + ZO and ZO treated groups. The authors did not report other FTIR parameters. Their finding regarding the mineral/matrix ratio is consistent with our study. However, in contrast to their results, we found a higher crystallinity in the callus and fracture cortex compared to control cortex. The difference might originate from the different analysis methods, as they calculated the crystallinity from the intensity ratio in the spectra whereas we used the peak fitting method. Ouyang et al. studied the effects of estrogen and estrogen deficiency in fracture callus in rat femurs. They found a lower mineral/matrix ratio, higher carbonate/phosphate ratio, lower XLR, and slightly reduced crystallinity in the fracture callus when compared to the fracture cortex at a distance from the fracture site in the estrogen deficient rats. After 8 weeks from the start of the treatment, the estrogen sufficient rats showed similar mineral/matrix ratio and crystallinity as the estrogen deficient rats, and a slightly higher XLR and lower carbonate/phosphate ratio near the fracture. These results are generally consistent with the data presented in this study. However, they did not report statistical significance. APS is associated with new mineral deposition and high amount of APS has been shown.

### 4 Discussion

In this study, we demonstrate that FTIR and SAXS combined can be used to identify the differences in composition and mineral structure between the newly formed callus tissue and cortical bone tissue. Moreover, we initiated an investigation of the effect of treatment with BMPs and bisphosphonates on the composition and mineral structure of newly formed callus tissue.

FTIR has been widely used to evaluate the composition of bone. However, studies on composition of newly formed fracture callus bone are limited. Yang et al. used FTIR to study the composition of callus and cortex during fracture healing in wild type and interleukin-6 knockout mice. They found a higher mineral/matrix ratio in the cortex compared to callus whereas crystallinity did not differ between the callus and cortex. The authors did not report other FTIR parameters. Their finding regarding the mineral/matrix ratio is consistent with our study. However, in contrast to their results, we found a higher crystallinity in the callus and fracture cortex compared to control cortex. The difference might originate from the different analysis methods, as they calculated the crystallinity from the intensity ratio in the spectra whereas we used the peak fitting method. Ouyang et al. studied the effects of estrogen and estrogen deficiency in fracture callus in rat femurs. They found a lower mineral/matrix ratio, higher carbonate/phosphate ratio, lower XLR, and slightly reduced crystallinity in the fracture callus when compared to the fracture cortex at a distance from the fracture site in the estrogen deficient rats. After 8 weeks from the start of the treatment, the estrogen sufficient rats showed similar mineral/matrix ratio and crystallinity as the estrogen deficient rats, and a slightly higher XLR and lower carbonate/phosphate ratio near the fracture. These results are generally consistent with the data presented in this study. However, they did not report statistical significance. APS is associated with new mineral deposition and high amount of APS has been shown.

![Fig. 6](https://example.com/fig6.png) SAXS analyses of (a) mineral plate thickness, (b) predominant orientation, and (c) degree of orientation in control cortex, fracture cortex, and fracture callus for all samples together and in different treatment groups. Nonparametric Wilcoxon signed rank test for all samples: *p < 0.05.
to indicate areas of new bone formation. However, it has not been reported earlier in fracture callus tissue. Our results showed a higher APS in the fracture callus compared to the cortices, which seem highly reasonable.

The mineral plate thickness has traditionally been evaluated using a method by Fratzl et al. It is based on a two-phase assumption, in which the mineral plate thickness is determined from the ratio of the integrated intensity and the Porod constant. Further, it is assumed that the ratio of the mineral phase is 50%. This approach has been successful when a relatively constant mineral phase fraction could have been assumed in bone. The calculated mineral plate thickness increases with increasing mineral phase fraction. Thus, the method may underestimate the mineral plate thickness if the real mineral phase fraction >0.5, and in new formed bone with a lower mineral phase fraction, it might overestimate it. Moreover, the mineral fraction has been found to vary within dentin. Similar variation could be expected also in bone. Bünger et al. introduced a curve fitting method in which assumption of the fraction of the mineral phase is not needed, which may be more accurate especially when determining the mineral plate thickness in bones with clearly different degrees of mineralization.

SAXS has been used to determine the mineral structure of trabecular and cortical bone. However, fracture callus tissue has received only minor attention in SAXS studies. Liu et al. evaluated the mineral plate thicknesses in fracture callus and fracture cortex in sheep undergoing fracture healing. Using the method proposed by Fratzl et al., they found a lower mineral plate thickness in fracture callus than in fracture cortex after 2 to 6 weeks, but no difference at 9 weeks after the fracture. Our study evaluated the healing after 6 weeks in a rat model. That is roughly comparable to 9 weeks healing time in a sheep model. In the present study, the mineral plate thickness tended to be higher in control cortex compared to the fracture cortex and callus, however, no significant differences were observed. Hypothetically, with a higher number of samples this may reach a significantly higher mineral plate thickness in control cortex than in fracture cortex and callus even at this stage of fracture healing.

The number of studies combining measurements of composition and mineral structure of bone in the same samples is highly limited. Pleshko-Camacho et al. used FTIR and SAXS to study the composition and mineral structure of cortical and trabecular bone of one L-4 vertebra of a 14-month-old girl. They mapped the trabeculae with both techniques and compared the results between trabecular and cortical bone. They also compared the parameter outcome from the two techniques. They found that the total mineral content and mineral/matrix ratio were lower in trabecular bone than in cortical bone, whereas mineral plate thickness and degree of orientation were lower in cortical bone than in trabecular bone. Trabecular bone has a more irregular orientation than in cortical bone, and in some stages of fracture healing, callus tissue can be described as to be of trabecular-like. Therefore, we compare these findings to those in the present study. We also found a higher mineral/matrix ratio in cortical bone than in callus. Degree of orientation was lower in callus tissue than in cortex. In the study by Pleshko-Camacho et al., they also found a significant correlation between crystallinity, determined from IR spectra, and mineral plate thickness, determined by SAXS. In our study, no correlation was found. However, this is not surprising, since crystallinity is suggested to be related to the HA crystal length. Thickness of the crystals is different from their length and may thus diverge from crystallinity without contradiction.

Lower degree of mineralization (mineral/matrix ratio), collagen maturity, and degree of orientation of the mineral crystals and higher APS are indicators of new, immature, and less organized bone. As expected, this was found in our study when the callus tissue was compared with the cortices. Also, the larger spread of angles in the callus tissue reflects the disordered nature of the forming bone. This would be expected, since the orientation of the collagen fibers and mineral crystals in cortical lamellar bone are oriented primarily in the direction of the long axis of the bone, i.e., the main loading direction. During bone healing, new bone formation occurs primarily through endochondral ossification resulting in quick formation of immature and less organized bone. Some calcified cartilage in the callus area could still be present at this stage of fracture healing, but the classification would require histology not available in this study. From the compositional data, the separation of woven bone and calcified cartilage is difficult, or even impossible. Immature bone has a more random orientation, which is reflected by the orientation parameters. With time, this immature and less organized bone tissue remodels into lamellar bone with a more organized structure, and finally, the cortex is restored. In our study, only one time point of 6 weeks was chosen. In our previous studies, this time point has corresponded to bridging of the fracture but not yet full remodeling. The amount of unremodeled callus depends both on the chosen time point and on the treatment given to the rats, which in this case was different combinations of BMP-7 and the bisphosphonate zoledronate. Based on the findings from our previous studies using the same open fracture healing model and time point, we assume that no or very little calcified cartilage remains in the callus areas. To confirm this, histological assessment would be necessary, which unfortunately was not available. In the NaCl group, the calluses were smaller compared to, e.g., BMP-treated calluses. This seems to be reflected in the orientation parameters, where the NaCl treated samples showed more similar orientation to the cortical bone compared to the large callus formations in the BMP-treated calluses.

When comparing all data points within one animal, significant differences were found between the callus tissue and the cortices. These differences were identified in some treatment groups, but not all. In all treatment groups, the mineral/matrix ratio, collagen maturity, and degree of orientation of the mineral plates were lower in the callus tissue than in the cortices. Additionally, the APS and orientation were higher in callus tissue than in the cortices in all samples and groups. However, when comparing the treatments in this manner, some differences between the treatments could also be detected. In BMP-7 + ZO and ZO treated samples, the mineral plate thickness was higher in callus tissue than in the fracture cortex. Also, crystallinity was higher in callus tissue compared to the fracture cortex in ZO treated samples. It might be postulated, that since bisphosphonates inhibit osteoclast activity and since the degree of bone resorption decreases, the mineral crystals are allowed to grow more freely and become larger, which could be seen in increasing mineral plate thickness and crystallinity.

We found that union was not achieved in the NaCl group, despite some sparse bone formation in the gap area. On the contrary, all samples in the BMP-7 + ZO groups were judged to be completely healed. These findings are consistent with
our previous studies using the same animal model with larger number of animals per group.2,5,16

In this study, the most noticeable limitation is the small number of samples. However, the current results are encouraging and suggest clear and significant differences in composition and mineral structure between the control cortex and fracture callus. Small number of samples is generally used in SAXS studies of bone.5,9,11,16 For example, Pleshko-Camacho et al. had one sample in their study,2 whereas Rinnerhalter et al. used two samples5 and in a more recent study Bünger et al. has 12 samples divided into four treatment groups.15 In the study by Liu et al., the callus tissue of four different aged sheep was investigated using SAXS.16 The main reason for the small number of samples is that the measurements are time consuming and that synchrotron beam time is highly limited. Another limitation is the spatial resolution of the SAXS scanning, which was 200 μm in this study. A more focused beam would allow better spatial resolution, in which turn would be beneficial when measuring the fine structured callus tissue. Additionally, the data were averaged over the sample thickness. The optimal thickness of SAXS samples ranges from below 100 up to 600 μm.24 The thickness in this study was 300 μm and although it is somewhat thicker than the beam size used (200 μm), it is still within these limits.

5 Conclusion

The current study contributes to the highly limited data on FTIR and SAXS on fracture healing and hard callus development and discusses important aspects of the use of SAXS on the newly formed bone. The methods indeed provide complementary information. Although FTIR enables reporting of the composition, the SAXS data enable assessment of mineral plate orientation and thickness, as well as the orientation of the collagen fibers. In conclusion, significant differences were found in the molecular composition and mineral structure between the fracture callus, fracture cortex, and control cortex. These techniques may also prove to be useful for the evaluation of effects of therapies on fracture healing. Taken together, the results indicate that SAXS is feasible when evaluating the mineral structure of the bone fracture callus and highlight the complementary nature of the two techniques.

Acknowledgments

This work was supported by the European Commission (FRACQUAL-293434), the Swedish Agency for Innovation Systems, the Swedish Research Council through the Linnaeus Center Organizing Molecular Matter, MAX IV Laboratory, the Foundation of Greta and Johan Kock, the strategic funding of the University of Eastern Finland, and the National Doctoral Programme of Musculoskeletal Disorders and Biomaterials (TBDP).

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