Intrinsic Plasticity Complements Long-Term Potentiation in Parallel Fiber Input Gain Control in Cerebellar Purkinje Cells

Belmeguenai, Amor; Hosy, Eric; Bengtsson, Fredrik; Pedroarena, Christine M.; Piochon, Claire; Teuling, Eva; He, Qionger; Ohtsuki, Gen; De Jeu, Marcel T. G.; Elgersma, Ype; De Zeeuw, Chris I.; Jörntell, Henrik; Hansel, Christian

Published in:
Journal of Neuroscience

DOI:
10.1523/JNEUROSCI.3226-10.2010

2010

Citation for published version (APA):

General rights
Copyright and moral rights for the publications made accessible in the public portal are retained by the authors and/or other copyright owners and it is a condition of accessing publications that users recognise and abide by the legal requirements associated with these rights.

• Users may download and print one copy of any publication from the public portal for the purpose of private study or research.
• You may not further distribute the material or use it for any profit-making activity or commercial gain
• You may freely distribute the URL identifying the publication in the public portal

Take down policy
If you believe that this document breaches copyright please contact us providing details, and we will remove access to the work immediately and investigate your claim.
Intrinsic Plasticity Complements Long-Term Potentiation in Parallel Fiber Input Gain Control in Cerebellar Purkinje Cells

Amor Belmeguenai, 1,2,‡* Eric Hosy, 1* Fredrik Bengtsson, 3‡ Christine M. Pedroarena, 3‡ Claire Piochon, 6 Eva Teuling, 1,6 Qionger He, 6 Gen Ohtsuki, 1,6 Marcel T. G. De Jeu, 6 Ype Elgersma, 6 Chris I. De Zeeuw, 1,7 Henrik Jörntell, 1 and Christian Hansel 1,6

1Department of Neuroscience, Erasmus University Medical Center, 3000CA Rotterdam, The Netherlands; 2Université de Lyon, Université Lyon 1, Centre National de la Recherche Scientifique, Unité Mixte de Recherche 5123, Villeurbanne, France; 3CTRS-IDEE, Hospices Civils de Lyon, Lyon, France; 4Department of Experimental Medical Science, Section for Neuroscience, Lund University, 22184 Lund, Sweden; 5Department of Cognitive Neurology, Hertie-Institute for Clinical Brain Research, University of Tübingen, 72076 Tübingen, Germany; 6Department of Neurobiology, University of Chicago, Chicago, Illinois 60637; and 7Netherlands Institute for Neuroscience, Royal Academy of Sciences (KNAW), 1105 BA Amsterdam, The Netherlands

Received June 22, 2010; revised Aug. 6, 2010; accepted Aug. 9, 2010.

This study was supported by grants from De Nederlandsche Organisatie voor Wetenschappelijk Onderzoek (separate grants to A.B., C.I.D.Z., and C.H.), SENSOPAC and the Prinses Beatrix Fonds (C.I.D.Z.), SF (C.P.), the Swedish Medical Research Council (H.J.), and National Institute of Neurological Disorders and Stroke Grant NS-62771 (C.H.).

Correspondence should be addressed to Christian Hansel, University of Chicago, Department of Neurobiology, 947 East 56th Street, Chicago, IL 60637. E-mail: chansel@bsd.uchicago.edu.

*F.B. and C.M.P. contributed equally to this work (co-first authors).
‡ Christine M. Pedroarena, and Christian Hansel contributed equally to this work (co-second authors).

Copyright © 2010 the authors 0270-6474/10/3013630-14$15.00/0

DOI:10.1523/JNEUROSCI.3226-10.2010

Introduction

A classic view in neuroscience holds that information storage and learning in neural circuits is made possible by alterations in synaptic transmission such as in long-term potentiation (LTP) and long-term depression (LTD). In the cerebellum, for example, bidirectional plasticity at PF-to-Purkinje cell synapses may be involved in motor learning (Ito, 1984; Jörntell and Hansel, 2006). More recently, plasticity of the intrinsic excitability of neurons has received attention as a potential correlate of learning (Hansel et al., 2001; Zhang and Linden, 2003; Frick and Johnston, 2005; Mozzachiodi and Byrne, 2010), but it remains unclear how intrinsic plasticity complements synaptic plasticity in memory formation. Excitability changes result from modifications of voltage- or calcium-dependent ion channels. So far, different types of K channels have been implicated in intrinsic plasticity, including A-type K channels (Schreurs et al., 1998; Frick et al., 2004), calcium-activated BK channels (Nelson et al., 2005), and SK channels (Sourdet et al., 2003; Lin et al., 2008). Moreover, it has been shown that activity-dependent alterations in hyperpolarization-activated I_h currents can adjust the excitability of neurons (Wang et al., 2003; Fan et al., 2005; Brager and Johnston, 2007).

In cerebellar circuits, activity-dependent intrinsic plasticity has been described in granule cells (Armano et al., 2000), Purkinje cells (Schreurs et al., 1998), as well as in the target neurons of Purkinje cells in the deep cerebellar nuclei (DCN) (Aizenman and Linden, 2000) and the vestibular nuclei (Nelson et al., 2005). In Purkinje cells, BK channel downregulation results in cerebellar ataxia (Saussier et al., 2004), while an enhancement of SK channel function has been shown to improve ataxia (Walter et al., 2006). These observations show how crucial the control of Purkinje cell activity patterns and intrinsic excitability is for proper motor coordination and suggest that Purkinje cells might use activity-dependent excitability alterations to fine-adjust the electrical output of the cerebellar cortex.

During recent years, research on the cellular basis of cerebellar motor learning has largely focused on LTD at PF synapses onto...
Purkinje cells, and more recently on LTP (Lev-Ram et al., 2002; Jörntell and Hansel, 2006; Dean et al., 2010). It has to be kept in mind, however, that the total synaptic gain (the gain of neuronal responsiveness to synaptic activity) can be modified, not only by changing properties of synaptic transmission (as in LTD and LTP), but also by local modifications of intrinsic excitability. To examine whether Purkinje cell intrinsic plasticity can play a role in cerebellar information storage, we characterized the cellular mechanisms underlying intrinsic excitability alterations, and determined how intrinsic plasticity affects the spontaneous activity of Purkinje cells and PF synaptic gain.

Materials and Methods

Purkinje cell recordings in vitro. Sagittal slices of the cerebellar vermis (250 μm) were prepared from postnatal day 17 (P17)–P28 Sprague Dawley rats after decapitation and isoflurane anesthesia. This procedure is in accordance with the guidelines of the Animal Care and Use Committees of both the Erasmus University Medical Center and the University of Chicago. In some experiments, mice (P17–P28) were used, as specified below. The slices were cut on a vibratome (Leica VT1000S) using ceramic blades. Subsequently, the slices were kept in artificial CSF (ACSF) containing the following (in mM): 124 NaCl, 5 KCl, 1.25 NaH2PO4, 2 MgSO4, 2 CaCl2, 26 NaHCO3, and 10 n-glucose, bubbled with 95% O2 and 5% CO2. The slices were allowed to recover for at least 1 h, and were then transferred to a submerged recording chamber superfused with ACSF at near-physiological temperature (34–35°C). In some recordings, the ACSF was supplemented with either 20 μM bicuculline methiodide or 100 μM picrotoxin to block GABA_A receptors. Picrotoxin was used in some experiments, as bicuculline salts have been reported to interfere with SK channel function (Seutin and Johnson, 1999). In our hands, however, intrinsic plasticity was observed independent of the type of antagonist used (bicuculline: Fig. 1 A, B, D, E; picrotoxin: Fig. 1 F).

Whole-cell patch-clamp recordings were performed under visual control using a 40X water-immersion objective mounted on a Zeiss Axioskop 2FS microscope. Patch pipettes (3–4 MΩ) were filled with internal saline containing the following (in mM): 9 KCl, 10 KOH, 120 K-glutamate, 3.48 MgCl2, 10 HEPES, 4 NaCl, 4 NaATP, 0.4 Na_GTP, and 17.5 sucrose, pH adjusted to 7.25. For cell-attached recordings, the pipette saline contained the following (in mM): 125 NaCl, 10 HEPES, 3 KCl, and 2 CaCl2. All drugs were purchased from Sigma. Patch-clamp recordings were performed in current-clamp mode (capacitance cancellation switched off) using an EPC-10 amplifier (HEKA Electronics). Membrane voltage and current were filtered at 3 kHz, digitized at 8 kHz (for action potential kinetics: 33 kHz), and acquired using Pulse software (HEKA Electronics). In the majority of recordings, a hyperpolarizing bias current was applied to prevent spontaneous spike activity. In some experiments (see Fig. 8), injection of bias currents was used to adjust the background spike rate of Purkinje cells. For PF stimulation, glass pipettes filled with ACSF were placed in the molecular layer, and for CF stimulation in the granule cell layer. Test responses were evoked at a frequency of 0.05 Hz using ~3 μA pulses that were applied for 500 μs. Intrinsic plasticity was monitored during the test periods by injection of brief (550 ms) depolarizing current pulses (100–200 pA) adjusted to evoke 5–15 spikes. The spike count was taken as a measure of excitability. Input resistance (Ri) was measured by injection of hyperpolarizing test currents (200 pA; 100 ms) and was calculated from the voltage transient toward the end of current injection.

Confocal calcium imaging. Calcium transients were recorded using a Zeiss LSM 5 Exciter confocal microscope equipped with a ×63 Zeiss Apochromat objective (Carl Zeiss Microimaging). Fluorescence was excited at 488 nm using an argon laser (Lasos Lasertechnik). For the calcium imaging experiments, sagittal slices of the cerebellar vermis (190–220 μm) were prepared from postnatal day 23–31 Sprague Dawley rats. Purkinje cells were loaded with the fluorescent calcium indicator dye Oregon Green BAPTA-2 (200 μM). After 10 min of dye loading, the patch pipette was removed and the dye was allowed to diffuse into the dendrite for another ~40 min. Subsequently, the cell was repatched with a second dye-filled patch electrode. This repatching protocol was used to avoid wash-out effects that might result from too long exposure to the pipette saline. The ACSF was supplemented with picrotoxin (100 μM), and in some experiments with the anti-oxidant vitamin C (100 μM). The recordings were performed at room temperature. Intrinsic plasticity was monitored using current pulses as in the nonimaging experiments, before and after tetanic current injection. Spine calcium transients were triggered by 100 Hz PF stimulation at intervals of 0.5–2 min. The number of PF pulses was adjusted to evoke a pronounced calcium transient (~8 pulses). Before tetanization, calcium transients were recorded during a 10 min baseline period. For each sweep, the data acquisition frequency was in the range of 19–58 Hz, depending on the size of the selected region of interest. Fluorescence changes were normalized to resting levels and expressed as the ratio ΔF/F(t) = [F(t) – F]/F, where F(t) is the fluorescence value at time t, and F is the averaged fluorescence obtained during the baseline period preceding the stimulus application. Data were discarded when changes in the baseline fluorescence exceeded 20%, with the
exception of recordings (then up to 25%) in which the change in the spine calcium transient (area) was at least twice as large.

**Genetically modified mice.** Mutant mice with a Purkinje cell-specific deletion of calcineurin (PP2B) were obtained by crossing the floxed CNB1 line (Zeng et al., 2001) with a L7-Cre line (Barski et al., 2000), resulting in a Purkinje cell-specific knock-out of the regulatory subunit (CNB1) of PP2B (C57BL/6/OlaHsd background). Littermates of the following genotypes were used for the experiments: PP2B-lox/PP2B-lox/L7Cre (L7-PP2B) and PP2B-lox/PP2B-lox (littermate controls). αCaMKII knock-out mice (Elgersma et al., 2002) and littermate control mice were obtained by breeding heterozygous knock-out mice in the C57BL/6/OlaHsd background.

**Purkinje cell recordings in vivo.** To obtain single unit recordings in a nonanesthetized in vivo preparation, adult Sprague Dawley rats were decerebrated as previously described (Bengtsson and Jörntell, 2007). After decerebration, the rats were paralyzed and mounted in a stereotactic frame for increased mechanical stability. Tungsten-in-glass microelectrodes were used for unitary, extracellular Purkinje cell recordings (5–25 μm exposed tip). Tungsten-in-glass microelectrodes (50–100 μm exposed tip) for parallel fiber stimulation (intensity 10–30 μA, pulse width 0.1 ms) were placed at middle depth of the molecular layer (100–150 μm depth from the surface), at 200–600 μm away from the recording electrode. Single-pulse PF stimulation was used to verify that simple spike activity was evoked in the recorded PC. Burst PF stimulation (15 pulses at 100 Hz, repeated at 1 Hz for 5 min) was delivered at intensities just above threshold for evoking simple spikes. The experimental procedures for the in vivo recordings were approved by the local Swedish Animal Research Ethics Committee.

**DCN recordings in vitro.** Cerebellar slices (275–290 μm) from P16–P20 rats were prepared and superfused with artificial CSF containing the following (in mM): 125 NaCl, 2.5 KCl, 1.3 NaH2PO4, 1.5 MgCl2, 26 NaHCO3, 20 glucose, 2.5 CaCl2. The solution was bubbled with 95% O2 and 5% CO2. Whole-cell recordings in current-clamp mode were obtained at 31°C ± 0.5 from large DCN neurons in the lateral or interpositus nuclei using an Axoclamp2B-ampifier (Molecular Devices). The intracellular electrode solution contained the following (in mM): 134 K-glucenate, 6 KCl, 10 K-HEPES, 0.1 EGTA, 0.3 NaGTP, 2 K-ATP, 10 phosphocreatine, 2 MgCl2. All recorded DCN neurons showed spontaneous firing (7.9–38 Hz) and triphasic afterhyperpolarizations (AHPs). Here, we took advantage of an in vitro preparation to investigate the effect of enhanced tonic Purkinje cell spike firing on spontaneous spike firing in DCN neurons. It has been pointed out that DCN spike patterns of enhanced tonic Purkinje cell spike firing on spontaneous spike firing lar nuclei. The stimulus intensity was adjusted at 0.1 Hz to induce a pause tungsten microelectrodes located in the white matter around the cerebel-

**Data analysis.** Data obtained from the Purkinje cell recordings in vitro were analyzed using Pulsifit (HEKA Electronics) and Igor Pro software (WaveMetrics). For statistical analysis, we used the paired Student’s t-test and the Mann–Whitney U test, when appropriate. Peristimulus time histograms (PSTHs) and cumulative spike probabilities (see Fig. 8) were calculated using SigmaPlot software (Hearne Scientific Software). Data obtained from the in vivo recordings were analyzed using the two-tailed Student’s t test. Data obtained from the DCN neuron recordings were analyzed off-line using programmable software Spike 2 (Cambridge Electronic Design), Igor (WaveMetrics), and SigmaStat (SPSS). Statisti-

cal analysis was performed using the Wilcoxon signed-rank test on data taken from 100 s before and 100 s after the switch from 30 to 50 Hz stimulation. The data from the first 10 s after the switch were excluded from the analysis. In all figures, the values shown represent the mean ± SEM.

**Results**

**Purkinje cell intrinsic plasticity can be observed in vitro and in vivo** To monitor intrinsic excitability changes in Purkinje cells, we performed whole-cell patch-clamp recordings in cerebellar slices (250 μm thick) obtained from P17–P28 Sprague Dawley rats at near-physiological temperature (34–35°C). Intrinsic excitability was measured in current-clamp mode by injecting brief depolarizing currents (~100–200 pA) that at the beginning of the recordings were adjusted to evoke 5–15 spikes. During the test periods before and after tetanization, these current steps were delivered at 0.05 Hz. GABA_A receptors were blocked by bath application of bicuculline methiodide (20 μM) or picrotoxin (100 μM). In these recordings, the number of depolarization-evoked spikes was taken as a measure of Purkinje cell excitability. An enhanced spike count was observed after repeated injection of depolarizing currents that were delivered at 5 Hz for 3 s (255.4 ± 20.3% of baseline ± SEM; n = 9; last 5 min; p = 0.00006) (Fig. 1A). Under control conditions, the spike count remained stable (103.4 ± 10.9%; n = 10; p = 0.19) (Fig. 1A). In the same recordings, we also monitored the amplitude of the AHP following the depolarizing current steps. After tetanization, the AHP amplitude was significantly reduced (81.9 ± 7.2%; n = 9; p = 0.02985) (Fig. 1B), but remained stable under control conditions (102.2 ± 3.3%; n = 10; p = 0.47) (Fig. 1B). Intrinsic plasticity was also observed when no GABA_A receptor antagonists were present in the bath (spike count: 155.2 ± 5.4%; n = 15; p = 0.00007) (Fig. 1C). However, the excitability increase was significantly larger when inhibition was blocked (p = 0.00006), suggesting that spontaneous GABAergic transmission limits intrinsic plasticity. A pronounced increase in the number of depolarization-

**Ethics Committee.** These procedures were done according to guidelines of the University of Tübingen and the local Committee on Animal Care and Use.

**Immunohistochemistry.** Two-week-old, four-week-old, and adult (3–4 months old) rats were anesthetized with pentobarbital and perfused transcardially with 4% paraformaldehyde. The brain was carefully dis-sected, postfixed in 4% paraformaldehyde for 1 h at 4°C, and rinsed overnight in 0.1 M phosphate buffer (PB) containing 30% sucrose. Forty micrometer sections of the cerebellum were cut on a freezing microtome and collected in 0.1 M PB. Sections were heated up in 0.25 M sodium citrate to 80°C for 30 min, rinsed in TBS, and blocked in TBS containing 10% normal horse serum (NHS) and 0.5% Triton, for 1 h at room temperature. Sections were incubated for 48–72 h at 4°C in TBS containing 2% NHS, 0.4% Triton, and primary antibodies at the following concentrations: rabbit anti-SK2 (Alomone or Sigma) 1:500; mouse anti-calbi-
nder the recording conditions described here (120.8 ± 5.0%; n = 6; p = 0.00198) (Fig. 1E). Finally, we adopted a PF burst protocol (10–15 pulses at 100 Hz, repeated at 1–3 s intervals for 5 min), which was designed to reflect granule cell activity patterns in vivo (Chadderton et al., 2004; Jörntell and Ekerot, 2006). Application of this 100 Hz PF burst protocol triggered intrinsic plasticity (179.6 ± 19.8%; n = 5; p = 0.00388) (Fig. 1F), and elicited LTP (supplemental Fig. 1, available at www.jneurosci.org as supplemental material) (see also Smith and Otis, 2005). None of these three tetanization protocols caused a significant change in the input resistance (current injection protocol: 100.3 ± 2.5%; n = 9; p = 0.91; 1 Hz PF protocol: 96.8 ± 4.7%; n = 9; p = 0.08; 100 Hz PF protocol: 100.0 ± 0.5%; n = 5; p = 0.95) (supplemental Table 1, available at www.jneurosci.org as supplemental material). However, intrinsic plasticity (current step protocol) (Fig. 1A) was associated with a decrease in the spike threshold (required current injection from −65 mV; before: 260.7 ± 54.2 pA; after tetanization: 225.0 ± 46.9 pA; n = 7; p = 0.0465).

To evaluate the physiological relevance of Purkinje cell intrinsic plasticity, we performed unitary, extracellular Purkinje cell recordings in nonanesthetized, decerebrated adult rats (P60–P90). In these in vivo recordings, the spontaneous spike rate was 42.4 ± 4.9 Hz (control recordings; n = 5). Application of the 100 Hz PF burst protocol caused an increase in the rate of spontaneous spike firing of Purkinje cells (control: 100.1 ± 1.0%; n = 5; p = 0.34; after tetanization: 115.0 ± 2.8%; n = 5; p = 0.0023) (Fig. 2A, B). Application of the 1 Hz PF stimulation protocol did not result in spike rate changes (supplemental Fig. 2, available at www.jneurosci.org as supplemental material). These in vivo recordings differ from the intrinsic plasticity experiments shown in Figure 1, in that the spontaneous spike rate (rather than the number of depolarization-evoked spikes) was measured and inhibition was left intact, thus allowing us to assess intrinsic plasticity under more physiological conditions. To examine whether intrinsic plasticity can be observed under these conditions in vitro, we applied both the current step protocol and the 100 Hz PF burst protocol, respectively, to cerebellar slices, when GABAergic inhibition was left intact, and Purkinje cells were allowed to spontaneously discharge (hyperpolarizing bias currents were omitted), resulting in an average background spike rate of 24.1 ± 2.9 Hz (control recordings; n = 8) (Fig. 2C). Under these conditions, repeated current injection induced a lasting increase in the spike rate (140.9 ± 15.3%; n = 10; p = 0.02565; control: 85.7 ± 5.9%; n = 8; p = 0.04557) (Fig. 2C). Application of the 100 Hz PF burst protocol also resulted in an increase in the background spike rate (175.2 ± 15.4%; n = 8; p = 0.0024) (Fig. 2D). As the in vivo recordings were performed using P60–P90 rats, we also examined whether intrinsic plasticity can be observed in that age group in vitro. In recordings from P60–P100 Purkinje cells in slices, we indeed observed an excitability increase when the 100 Hz PF stimulation protocol was applied (spike count: 280.6 ± 24.9%, n = 8; p = 0.00001) (supplemental Fig. 3, available at www.jneurosci.org as supplemental material). These recordings suggest that intrinsic plasticity is a robust phenomenon that can be observed both in vitro and in vivo, in young adult and adult animals, and in the absence and presence of GABAA receptor blockers. Furthermore, the change in intrinsic excitability can be monitored when counting depolarization-evoked spikes, or when measuring spontaneous spike activity. It can be argued that both whole-cell patch-clamp recordings from Purkinje cells in vitro and single-unit recordings in vivo might disrupt physiological Purkinje cell responses. We thus additionally characterized intrinsic plasticity using cell-attached recordings in vitro, which might provide a less invasive recording technique. When cell-attached recordings were performed from P20–P27 rats, an average background spike frequency of 32.5 ± 4.8 Hz (control recordings; n = 7) (Fig. 2F) was observed. Application of the 100 Hz PF burst protocol resulted in an increase in the spontaneous spike rate (140.3 ± 6.9%; n = 10; p = 0.0002; control: 94.9 ± 6.6%; n = 7; p = 0.83) (Fig. 2E,F). These cell-attached recordings confirm that intrinsic plasticity is a robust phenomenon that can be observed under a wide range of experimental conditions. In most subsequent experiments, we used the nonsynaptic cur-

![Figure 2](https://example.com/figure2.png)

**Figure 2.** Changes in the spontaneous Purkinje cell spike rate recorded in vivo and in vitro. **A,** 100 Hz PF burst stimulation caused an increase in simple spike firing in vivo (n = 5). The time graph shows the percentage change of the recorded spike frequency (Hz). The traces show single-unit extracellular recordings before (top) and after tetanization (bottom). Calibration: 0.2 mV, 50 ms. **B,** Bar graph comparing the spike rate observed after PF burst stimulation (n = 5; 0–30 min after tetanus) to that measured under control conditions (n = 5). **C,** Repeated current injection enhanced the spontaneous Purkinje cell spike rate in vitro (closed circles; n = 10). Under control conditions, the spike rate was not enhanced (open circles; n = 8). **D,** 100 Hz PF tetanization also caused an increase in the tonic spike rate in vitro (n = 8). **C,D,** Calibration: 20 mV, 200 ms. E, In cell-attached recordings in vitro, 100 Hz PF stimulation enhanced spontaneous spike firing (n = 10). F, This increase was not seen under control conditions (n = 7). Calibration: E, 100 pA, 200 ms; F, 200 pA, 200 ms. For these recordings, only cells were used that showed regular simple spike firing. The arrows indicate the time point of tetanization. All experiments shown in this figure were performed in the absence of GABAA receptor blockers in the bath. Error bars indicate SEM.
Intrinsic plasticity is mediated by a downregulation of SK channels

Intrinsic excitability changes can be mediated by alterations in voltage- or calcium-sensitive ion channels (Zhang and Linden, 2003; Frick and Johnston, 2005). In adult Purkinje cells, high-quality dendritic voltage control is difficult to achieve. Because of this so-called “space-clamp” problem, we screened for types of ion channels involved by addressing three questions after bath application of ion channel antagonists: (1) Is the spike count upregulated after drug application? (2) Is there a change in the action potential waveform? and (3) Does application of an antagonist occlude an excitability increase caused by tetanic current injection (5 Hz, 3 s)? Under control conditions, the amplitude and kinetics (rise time/spike width) of individual action potentials were unaffected after tetanization. Rather, we observed that after tetanization, the rate of depolarization toward spike threshold was enhanced following each action potential (Fig. 3A, B, Table 1) (n = 5), resulting in the elevated spike rate.

To determine the mechanism underlying this change in the firing rate, we first tested the effects of 4-aminopyridine (4-AP), an antagonist of some types of voltage-gated Kv channels, including A-type K currents (I_A). A previous study has demonstrated that delay eyeblink conditioning in rabbits is associated with an increase in Purkinje cell excitability (Schreurs et al., 1998), resembling similar excitability changes observed in CA1 pyramidal cells after trace eyeblink conditioning (Moyer et al., 1996). This type of Purkinje cell intrinsic plasticity was described as being mediated by changes in I_A (Schreurs et al., 1998). I_A has also been implicated in excitability alterations in CA1 hippocampal pyramidal cells in vitro (Ramakers and Storm, 2002; Frick et al., 2004). When 4-AP was bath applied (100 μM), the spike count was significantly enhanced (177.0 ± 8.4%; last 3 min; n = 11; p = 0.000007) (Fig. 3C). However, the width of individual action potentials was significantly increased, too (before: 0.27 ± 0.03 ms; 4-AP: 0.44 ± 0.04 ms; n = 11; p = 0.025) (Fig. 3C, Table 1). Such a prolongation was not seen under control conditions (before: 0.29 ± 0.03 ms; after tetanization: 0.28 ± 0.02 ms; n = 5; p = 0.18) (Table 1). Thus, a downregulation of 4-AP-sensitive Kv conductances does not mediate the form of intrinsic plasticity described here. Accordingly, bath application of 4-AP did not occlude further excitability increases following tetanic current injection [158.0 ± 13.9%; n = 6 (Fig. 3D) as compared to 155.2 ± 5.4%; n = 15 (Fig. 1C); p = 0.09]. At the beginning of these occlusion experiments, the spike count was reset to baseline levels (5–15 spikes) by ad-
justing the amplitude of the current steps and the amplitude of the hyperpolarizing bias currents. On average, there was no significant difference in the amplitudes of these currents between the control group and the drug application groups described here (current steps; control: 180.8 ± 17.74 pA; n = 15; 4-AP: 160.0 ± 26.64 pA; p = 0.52; n = 7; HpTX-2: 150.0 ± 32.56 pA; p = 0.51; n = 5; apamin: 139.54 ± 11.99 pA; p = 0.09; n = 11; Student’s t test/holding currents; control: -471.56 ± 14.02 pA; n = 15; 4-AP: -450.86 ± 45.65 pA; p = 0.57; n = 7; HpTX-2: -384.0 ± 36.69 pA; p = 0.06; n = 5; apamin: -527.18 ± 29.38 pA; p = 0.08; n = 11; Student’s t test) (supplemental Fig. 4, available at www.jneurosci.org as supplemental material). Moreover, there was no significant correlation between the current step amplitude or holding current amplitude, respectively, and the intrinsic plasticity amplitude (current steps; control: p = 0.35; 4-AP: p = 0.81; HpTX-2: p = 0.16; apamin: p = 0.54; holding currents: control: p = 0.51; 4-AP: p = 0.99; HpTX-2: p = 0.06; apamin: p = 0.08; Pearson test) (supplemental Fig. 4, available at www.jneurosci.org as supplemental material), excluding the possibility that differences in the occlusion experiments might be due to differences in the amplitude of either the current steps or the bias holding currents.

We next tested the effects of heteropodatoxin-2 (HpTX-2), a selective inhibitor of Kv4.2-mediated \( I_A \) currents. Bath application of HpTX-2 (100 nm) significantly enhanced the spike count (187.1 ± 8.6%; n = 7; p = 0.00004) (Fig. 3E). Like 4-AP, HpTX-2 increased the action potential width (before: 0.32 ± 0.04 ms; after tetanization: 0.34 ± 0.04 ms; n = 6; p = 0.033) (Fig. 3E, Table 1), but with HpTX-2 this effect was less pronounced. HpTX-2 application did not occlude excitability increases triggered by tetanic current injection (165.1 ± 12.9%; n = 6) (Fig. 3F) (as compared to 155.2 ± 5.4%; n = 15) (Fig. 1C) (p = 0.88). These data show that Kv4.2-mediated \( I_A \) currents are not involved in Purkinje cell intrinsic plasticity.

Next, we examined the involvement of SK-type calcium-dependent K channels in Purkinje cell intrinsic plasticity. In layer V pyramidal neurons, SK channels have been implicated in activity-dependent excitability increases (Sourdet et al., 2003). In Purkinje cells, SK channels have been shown to control spike firing (Edgerton and Reinhart, 2003; Womack and Khodakhah, 2003). Bath application of the SK-channel blocker apamin (3 nm) caused a pronounced increase in the spike count (316.1 ± 39.7%; n = 15; p = 0.00003) (Fig. 3G), without changing the action potential width (before: 0.22 ± 0.02 ms; apamin: 0.24 ± 0.02 ms; n = 15; p = 0.08) (Fig. 3G, Table 1). However, following each action potential, the rate of depolarization toward spike threshold was enhanced in the presence of apamin. This effect was also observed under control conditions after tetanization (Fig. 3A, Table 1). Finally, after bath application of apamin, subsequent tetanization caused only a moderate increase in the spike count (119.6 ± 6.4%; n = 10; p = 0.008) (Fig. 3H), which was significantly lower than that observed under control conditions (155.2 ± 5.4%; n = 15; p = 0.009). This partial occlusion effect after apamin bath application suggests that a downregulation of SK-type K currents contributes to Purkinje cell intrinsic plasticity. Finally, we tested the BK-type calcium-dependent K channel blocker ibotocin. BK channels control the spread of calcium spikes in Purkinje cell dendrites (Rancz and Häusser, 2006), and might therefore be involved in excitability control as well. While bath application of ibotocin (50 nm) enhanced the spike count (184.1 ± 30.2%; n = 7; p = 0.02) (supplemental Fig. 5A, available at www.jneurosci.org as supplemental material), it did not occlude intrinsic plasticity after current injection (159.9 ± 19.6%; n = 6; p = 0.80) (supplemental Fig. 5B, available at www.jneurosci.org as supplemental material), which rules out the possibility that BK-type K channels are involved in intrinsic plasticity.

These experiments show that the form of intrinsic plasticity described here is at least partially mediated by a downregulation of SK-type calcium-dependent K channels. Of the three types of apamin-sensitive SK channel subunits (SK1–SK3), only SK2 channels are expressed in rat Purkinje cells, but their levels have been shown to decline during the first 3 postnatal weeks (Cingolani et al., 2002). In contrast to rats, mice have been shown to express SK2 channels in the adult Purkinje cell layer (Sailer et al., 2004). As our patch-clamp recordings were obtained from up to 4-week-old rats, we examined whether SK2 channels are indeed expressed in young adult and adult rat Purkinje cells as suggested by recordings from our laboratory and others (Womack and Khodakhah, 2003), and performed immunostainings using antibodies directed against the C-terminal domain of the SK2 channel sub-

### Table 1. Changes in action potential waveform parameters evoked by tetanization and by application of K channel antagonists, respectively.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Control before IP</th>
<th>Control after IP</th>
<th>Under 4AP</th>
<th>Under HpTX-2</th>
<th>Under apamin</th>
<th>Under apamin after IP</th>
</tr>
</thead>
<tbody>
<tr>
<td>Amplitude (mV)</td>
<td>91.6 ± 5.3</td>
<td>89.9 ± 4.5</td>
<td>87.7 ± 6.9</td>
<td>76.1 ± 5.5</td>
<td>98.4 ± 5.4</td>
<td>100.6 ± 5.7</td>
</tr>
<tr>
<td>AHP amplitude (mV)</td>
<td>-6.9 ± 1.2</td>
<td>-6.1 ± 2.1</td>
<td>-4.9 ± 1.6</td>
<td>-7.1 ± 1.5</td>
<td>-7.8 ± 2.2</td>
<td>-7.1 ± 1.9</td>
</tr>
<tr>
<td>Em after spike (mV)</td>
<td>49.6 ± 1.6</td>
<td>48.4 ± 1.6</td>
<td>52.0 ± 2.3</td>
<td>59.9 ± 1.3</td>
<td>51.9 ± 1.3</td>
<td>49.9 ± 1.5</td>
</tr>
<tr>
<td>10%-90% (ms)</td>
<td>198 ± 0.015</td>
<td>186 ± 0.015</td>
<td>180 ± 0.001</td>
<td>210 ± 0.019</td>
<td>180 ± 0.013</td>
<td>165 ± 0.010</td>
</tr>
<tr>
<td>FWHM (ms)</td>
<td>288 ± 0.028</td>
<td>276 ± 0.022</td>
<td>270 ± 0.026</td>
<td>320 ± 0.040</td>
<td>220 ± 0.017</td>
<td>235 ± 0.046</td>
</tr>
</tbody>
</table>

Parameters of the action potential waveform that were monitored are the peak amplitude, the AHP amplitude, the 10–90% rise time, the full width at half magnitude (FWHM), and the rate of postspike depolarization (\( t_{200} \), measured 4.5–6 ms after the spike peak). The table summarizes these parameters (see also Fig. 3B) under control conditions (n = 5) and with 4-AP bath application (n = 11), heteropodatoxin-2 bath application (n = 7), and apamin bath application (n = 15).
unit, and the Purkinje cell-specific marker calbindin (Fig. 4; for antibody specificity, see supplemental Fig. 6, available at www.jneurosci.org as supplemental material). Cerebellar sections obtained from 2-week-old rats (top row), 4-week-old rats (middle row), and adult rats (3–4 months; bottom row) indicated presence of SK2 channel subunits at all three ages, which overlapped with the Purkinje cell-specific calbindin staining. Throughout all ages, but particularly in the adult sections, we also observed SK2 channel staining around granule cell bodies, and in the molecular layer. These immunohistochemical data show that SK2 channels are expressed in Purkinje cells during development and after maturation; therefore, SK2 channels can be involved in the type of Purkinje cell excitability alterations described here.

What molecular events are involved in intrinsic plasticity upstream of the functional downregulation of SK2 channels described above? We first examined whether intrinsic plasticity depends on dendritic calcium signaling, and the activation of protein phosphatases 1, 2A, and 2B, which have been implicated in PF-LTP (Coesmans et al., 2004; Belmeguenai and Hansel, 2005). Intrinsic plasticity was blocked when the calcium chelator BAPTA (20 mM) was added to the pipette saline (86.9 ± 13.6%; n = 11; p = 0.37) (Fig. 5A), suggesting that calcium is a prerequisite for the induction process. In the absence of tetanization, BAPTA application alone transiently enhances the spike rate, but has no lasting effect on Purkinje cell excitability (n = 10) (for wash-in control recordings for all drug experiments shown in Figs. 5 and 6, see supplemental Fig. 7, available at www.jneurosci.org as supplemental material).

When the PP1/2A inhibitor microcystin LR was added to the pipette saline (10 μM), the current injection protocol failed to enhance the spike count (105.5 ± 20.8%; n = 7; p = 0.80) (Fig. 5B). Similarly, bath application of the PP1/2A inhibitor okadaic acid (1 μM) blocked intrinsic plasticity (108.8 ± 4.2%; n = 6; p = 0.064) (Fig. 5B). When the PP2B inhibitor cyclosporin A was bath applied (100 μM), the spike count was even decreased (55.6 ± 10.1%; n = 8; p = 0.00317) (Fig. 5B). These results suggest that Purkinje cell intrinsic plasticity shares the requirement for PP1/2A and PP2B activation with PF-LTP. The PP2B dependence of intrinsic plasticity was confirmed using mice, in which PP2B was selectively knocked out in Purkinje cells (L7-PP2B mice). In wild-type mice, application of the current injection protocol enhanced the spike rate (138.9 ± 10.2%; n = 16; p = 0.009) (Fig. 5C). Intrinsic plasticity was absent in L7-PP2B knock-out mice (89.7 ± 9.4%; n = 9; p = 0.40) (Fig. 5C). Conversely, PP2B activation can enhance the spike count: when active PP2B was added to the pipette saline (5 μg/ml), the spike count increased significantly (149.1 ± 15.2%; n = 11; p = 0.00423) (supplemental Fig. 8A, available at www.jneurosci.org as supplemental material). Subsequently, intrinsic plasticity was occluded (101.5 ± 9.0%; n = 8; p = 0.87) (supplemental Fig. 8B, available at www.jneurosci.org as supplemental material). The phosphatase dependence of intrinsic plasticity is not restricted to excitability increases triggered by application of the current step protocol. Likewise, application of the 1 Hz PF stimulation protocol failed to cause excitability changes in the presence of the PP1/2A inhibitor okadaic acid (1 μM; 105.4 ± 5.8%; n = 8; p = 0.37) (Fig. 5D) or

Figure 4. SK2 channel immunostaining. Anti-SK2-channel (red) and anti-calbindin (green) antibody stainings of cerebellar sections obtained from 2 week-old (top row), 4 week-old (middle row), and adult (bottom row) rats show SK2-staining in Purkinje cell dendrites and throughout the molecular layer. Right side, Enlarged views taken from the areas indicated by white boxes on the left.
There was no statistical difference between these two groups. The reduction in the intrinsic plasticity amplitude as the tetanization share the same induction mechanism, which involves activation of protein phosphatases 1, 2A, and 2B. Intrinsic excitability changes triggered by current injection or PF tetanization (80.6 ± 5.2%; n = 12; p = 0.000179) (Fig. 5B). At both concentrations, the reduction in the intrinsic plasticity amplitude was significant (30 μM: p = 0.00026; 60 μM: p = 0.00012; Mann–Whitney U test), which shows that PKA activation is required to enhance excitability. We finally examined the involvement of casein kinase 2 (CK2) in intrinsic plasticity. CK2 is a constitutively active enzyme (Biddul et al., 2004) that phosphorylates SK2-associated calmodulin and reduces the calcium sensitivity of SK2 channels (Allen et al., 2007). Intrinsic plasticity was blocked in the presence of the CK2 inhibitor emodin (30 μM; n = 8). Intrinsic plasticity was also blocked by bath application of the CK2 inhibitor DMAT at 5 μM (open circles; n = 6) and 10 μM (closed circles; n = 6). Error bars indicate SEM.

**Figure 5.** Purkinje cell intrinsic plasticity depends on calcium signaling, and activation of protein phosphatases 1, 2A, and 2B. A, Intrinsic plasticity was blocked when BAPTA (20 mM) was added to the pipette saline (n = 11). B, Likewise, the current injection protocol was ineffective in the presence of the PP1/2A inhibitor microcystin LR (10 μM; closed circles; n = 7), the PP1/2A inhibitor okadaic acid (1 μM; open triangles; n = 6), and the PP2B inhibitor cyclosporin A (100 μM; open circles; n = 8). Cyclosporin A (100 μM; open triangles; n = 8) and cyclosporin A (100 μM; open circles; n = 8) and cyclosporin A (100 μM; open triangles; n = 8). Copyright © The Rockefeller University Press. All rights reserved.

**Figure 6.** Involvement of protein kinases in Purkinje cell intrinsic plasticity. A, Intrinsic plasticity was observed in αCaMKII knock-out mice (open circles; n = 12) and wild-type controls (closed circles; n = 7). B, Bath application of the PKA inhibitor KT5720 at 30 μM (open circles; n = 14) and 60 μM (closed circles; n = 12) affected intrinsic plasticity. C, Excitability increases were prevented by bath application of the CK2 inhibitor emodin (30 μM; n = 8). D, Intrinsic plasticity was also blocked by bath application of the CK2 inhibitor DMAT at 5 μM (open circles; n = 6) and 10 μM (closed circles; n = 6). Error bars indicate SEM.


Involvement of protein kinases in Purkinje cell intrinsic plasticity. A, Intrinsic plasticity was observed in αCaMKII knock-out mice (open circles; n = 12) and wild-type controls (closed circles; n = 7). B, Bath application of the PKA inhibitor KT5720 at 30 μM (open circles; n = 14) and 60 μM (closed circles; n = 12) affected intrinsic plasticity. C, Excitability increases were prevented by bath application of the CK2 inhibitor emodin (30 μM; n = 8). D, Intrinsic plasticity was also blocked by bath application of the CK2 inhibitor DMAT at 5 μM (open circles; n = 6) and 10 μM (closed circles; n = 6). Error bars indicate SEM.

Purkinje cell intrinsic plasticity does not affect DCN spike rates

Under physiological conditions, Purkinje cells are spontaneously active and fire simple spikes at 30–80 Hz (Simpson et al., 1996). In our recordings from spontaneously active Purkinje cells summarized in Figure 2, we observed background spike frequencies of ~40 Hz in vivo and ~30 Hz in vitro, respectively. Under various recording conditions both in vivo and in vitro, the spontaneous background spike frequency was enhanced when intrinsic plasticity protocols were applied (Fig. 2). These results show that intrinsic plasticity can increase the rate of Purkinje cell pacemaker activity, which is largely of intrinsic origin and persists when glutamatergic transmission is blocked (Hausser and Clark, 2004).
Intrinsic plasticity-associated changes in the spontaneous Purkinje cell spike rate do not affect the tonic spike rate of DCN neurons. A. Bottom, the spike frequency of DCN neurons (F) remained stable when the increase in Purkinje cell spike rates was mimicked by a switch from 30 to 50 Hz stimulation of Purkinje cell axons (n = 7; values for the onset of 30 Hz activation are not depicted). Top, Two example recordings. The insets show the effect of a switch from 0 to 30 Hz stimulation of the inhibitory synapses. B. Top, Example recordings using a higher stimulus intensity. Bottom, Increasing the stimulus intensity resulted in larger amplitudes of control IPSPs (0.1 Hz, left), but the effect of tonic stimulation at 30 and 50 Hz on the firing rate of DCN neurons was similar using high (HI; n = 8) or low intensities (LI; n = 7). C. Bottom, At both high and low stimulus intensities, IPSP amplitudes were reduced to a larger degree at 50 Hz than at 30 Hz stimulation (averaged over 1 min before and after the frequency switch). Top, Example traces illustrating averaged IPSPs. The traces were taken from the same recording shown in B. Error bars indicate SEM. *p < 0.05.

Figure 7.

Intrinsic excitability changes affect the impact of PF signaling. What then are the functional consequences of alterations in Purkinje cell excitability? Does intrinsic plasticity affect the input–output function of Purkinje cells? It has previously been shown for weak PF inputs that the number of evoked spikes linearly reflects synaptic input strength (Walter and Khodakhah, 2006; Mittmann and Häusser, 2007). To examine whether intrinsic
were not significantly changed (0.3; enhanced this frequency ratio (the frequency recorded before. Increasing stimulus strength enhanced the frequency resulting from PF stimulation as the ratio obtained similar results when analyzing the spike frequency, significance (was slightly enhanced, but this trend did not reach statistical put of Purkinje cells by LTD and LTP, respectively. After repeated p/H11005/LTD and LTP conditions by adjusting the stimulus strength to

Varying the stimulus strength altered the number of evoked spikes ranging from one to four spikes (p/H11005/25%: 2.9, Cumulative spike probabilities at 30 Hz (p/H11005/E) and 60 Hz (p/H11005/F). The dotted line represents a fit to the baseline. H, Corrected cumulative spike probabilities after subtraction of the baseline fit at 30 Hz and 60 Hz. I, Inverse relationship between the spike rate and the net increase in spike firing, which was calculated from 100 ms time windows before and after PF stimulation (n = 7). The analysis shown in D–H is based on 477 spikes (30 Hz) and 247 spikes (60 Hz), respectively (n = 7). Error bars indicate SEM.

plasticity affects the input–output curve of Purkinje cells, we allowed the cells to spontaneously discharge and counted the number of spikes evoked by PF stimulation. We then mimicked LTD and LTP conditions by adjusting the stimulus strength to ±25% of the control level (Fig. 8A). We observed a linear relationship between stimulus strength and the number of evoked spikes, but the dynamic range was very limited: varying the stimulus strength was more pronounced than that observed at a background spike frequency of 30 Hz is more pronounced than that observed at a background spike frequency of 60 Hz (Fig. 8H), suggesting that the impact of synaptic signaling is higher at a lower tonic background spike frequency. Over several background frequencies tested, the increase in firing caused by PF stimulation (as a percentage; calculated from 100 ms time periods before and after PF stimulation) decreased with increasing spike frequencies (Fig. 8C,F).

Intrinsic plasticity enhances dendritic calcium signaling, but lowers the probability for subsequent LTP induction

Synaptic gain control is typically associated with LTP and LTD. To examine whether intrinsic plasticity interferes with synaptic plasticity, we tested whether the LTP induction probability was altered after previous enhancement of intrinsic excitability. In these experiments, PF-EPSPs and the spike count were monitored within the same sweeps (Fig. 9A). Application of the current injection protocol (5 Hz, 3 s) enhanced the spike count control: 2.3 ± 0.5; p = 0.33; n = 9; +25%: 2.8 ± 0.6; p = 0.83; n = 8; paired Student’s t test). In these experiments, changes in background spiking were prevented by injection of bias currents.

To examine whether the intrinsic plasticity-associated increase in the background spike rate itself affects PF signaling, we performed the opposite experiment and injected bias currents to vary the background spike rate (mimicking intrinsic plasticity), while the PF stimulus strength was kept constant (Fig. 8C). Alterations in the spike rate ranging from 30 to 60 Hz did not result in changes in the immediate PF response (Fig. 8C), but the enhanced background spike rate lowered the signal-to-noise ratio and thus lowered the transient PF-evoked net increase in spike firing (n = 7) (Fig. 8C–F). These data are in line with a previous report demonstrating an inverse relationship between Purkinje cell background spike rates and PF readout (McKay et al., 2007), and suggest that as a consequence of the activity-dependent plasticity described here, the impact of PF signaling is actually reduced despite of an increase in excitability.

To directly quantify the number of additional spikes that is attributable to PF stimulation at background spike frequencies of 30 and 60 Hz, respectively, we adopted an analysis method that provides an estimate of the impact of a given synaptic input on the spike output of neurons (Mittmann and Häusser, 2007). We first calculated the PSTH of the PF response patterns (Fig. 8D,E), and plotted the corrected cumulative spike probability (Fig. 8F,G). Subsequently, we fitted a line to the baseline period, which was then subtracted from the cumulative probability trace. The corrected cumulative spike probability traces (Fig. 8F) indicate the number of spikes that were caused by PF stimulation. The increase in spike firing attributable to PF stimulation at a background spike frequency of 30 Hz is more pronounced than that observed at a background spike frequency of 60 Hz (Fig. 8H), suggesting that the impact of synaptic signaling is higher at a lower tonic background spike frequency. Over several background frequencies tested, the increase in firing caused by PF stimulation (as a percentage; calculated from 100 ms time periods before and after PF stimulation) decreased with increasing spike frequencies (Fig. 8C,F).

Intrinsic plasticity enhances dendritic calcium signaling, but lowers the probability for subsequent LTP induction

Synaptic gain control is typically associated with LTP and LTD. To examine whether intrinsic plasticity interferes with synaptic plasticity, we tested whether the LTP induction probability was altered after previous enhancement of intrinsic excitability. In these experiments, PF-EPSPs and the spike count were monitored within the same sweeps (Fig. 9A). Application of the current injection protocol (5 Hz, 3 s) enhanced the spike count

Figure 8. Purkinje cell intrinsic plasticity lowers the impact of PF signaling. A, Intrinsic plasticity does not affect the synaptic input–output curve of Purkinje cells. PF stimulus strength was varied by ±25%, which approximately corresponds to amplitude changes seen after LTD and LTP induction. B, The linear relationship between stimulus strength and the number of evoked spikes was not significantly affected by intrinsic plasticity (n = 8; 9; paired Student’s t test; p > 0.05; the number of spikes evoked at all three stimulus strengths was compared before and after inducing intrinsic plasticity). The spike count includes all spikes that occurred at elevated frequency after PF stimulus onset during a 100 ms time window. The number of recordings is indicated in the brackets. C, When the background spike rate was enhanced from 30 Hz (top) to 60 Hz (bottom), the net increase in spike numbers evoked by constant PF stimulation was lowered. D, E, PSTHs calculated from PF responses at background spike frequencies of 30 Hz (D) and 60 Hz (E). F, G, Cumulative spike probabilities at 30 Hz (F) and 60 Hz (G). The dotted line represents a fit to the baseline. H, Corrected cumulative spike probabilities after subtraction of the baseline fit at 30 Hz and 60 Hz. I, Inverse relationship between the spike rate and the net increase in spike firing, which was calculated from 100 ms time windows before and after PF stimulation (n = 7). The analysis shown in D–H is based on 477 spikes (30 Hz) and 247 spikes (60 Hz), respectively (n = 7). Error bars indicate SEM.
Intrinsic Plasticity in Cerebellar Purkinje Cells

Belmeguenai et al.

Figure 9. Intrinsic plasticity enhances spine calcium signaling, but blocks subsequent LTP induction. A, PF-EPSPs (right) and the spike count (left) were monitored after tetanic current injection (n = 8) and after subsequent application of the 1 Hz PF tetanization protocol (n = 5). LTP induction was blocked after previous application of the intrinsic plasticity protocol (black circles; n = 3). In contrast, LTP was induced by PF stimulation, when intrinsic plasticity was not previously triggered (white circles; n = 6). Top, Traces show EPSPs and depolarization-evoked spikes under baseline conditions (left), after application of the intrinsic plasticity protocol (middle), and after application of the PF-LTP protocol (right). In all recordings shown in A, inhibition was left intact. Calibration: 20 mV, 200 ms. B–H, Confocal calcium imaging experiments reveal an increase in spine calcium transients. B, Top, Purkinje cell filled with the fluorescent calcium indicator Oregon Green BAPTA-2 (200 μM). Scale bar, 20 μm. Bottom, Enhanced view of the area marked by the red box in the top image. The red circle indicates the region of interest. Scale bar, 2 μm. C, The spike count was monitored before (left) and after tetanization (right). D, Calcium transients were evoked by 100 Hz PF stimulation (4 pulses). PF responses are shown before (left) and after tetanization (right). E, Calcium transients evoked by the PF responses shown in D. The traces represent averages of 3 calcium transients. F, An overlay of the calcium transients reveals enhanced calcium signaling. G, Bar graph showing averaged changes in the area under the curve (left) and peak of calcium transients (middle), as well as the spike count (right). These values represent averages taken during a 12 min period following tetanization (n = 7). H, Comparison of calcium transient changes (area under the curve) in spines and associated shaft regions (n = 7). Error bars indicate SEM.

(187.5 ± 18.1%, t = 15–20 min; n = 8; p = 0.00188) (Fig. 9A), but did not affect the EPSP amplitude (106.1 ± 3.4%; n = 8; t = 15–20 min; p = 0.12) (Fig. 9A). Subsequently, we applied the PF-LTP protocol (1 Hz, 5 min) to test whether intrinsic plasticity affects LTP induction. Application of the 1 Hz PF stimulation protocol further enhanced the spike count (302.4 ± 25.9%; n = 5; t = 40–45 min; p = 0.00133) (Fig. 9A). However, when intrinsic plasticity was triggered first, LTP could not be induced anymore by subsequent 1 Hz PF stimulation (104.5 ± 3.4%; n = 5; t = 40–45 min; p = 0.26) (Fig. 9A). This observation likely reveals a metaplastic interaction rather than a run-down of the LTP probability (known from LTD induction), because in control experiments, LTP could be observed after an equally long (20 min) baseline period (121.3 ± 6.4%; n = 11; p = 0.0079) (supplemental Fig. 9, available at www.jneurosci.org as supplemental material). The difference in EPSP amplitude changes monitored after (1) the application of the LTP protocol alone and (2) application of the LTP protocol following previous application of the current injection protocol was statistically significant (Mann–Whitney U test; p = 0.0351). For comparison, LTP induced after a shorter baseline period of 5 min amounted to 124.5 ± 6.4% (n = 6; p = 0.01; last 5 min) (Fig. 9A; note that an extended version of this LTP graph is shown in Fig. 1E). There was no statistical difference between the LTP amplitudes reached after baseline periods of 5 min and 20 min, respectively (Mann–Whitney U test; p = 0.45). In these experiments, intrinsic plasticity was first triggered in the absence of synaptic alterations (nonsynaptic induction protocol) to be able to examine how intrinsic plasticity as an isolated phenomenon affects the LTP induction probability (1 Hz PF protocol). The results suggest that intrinsic plasticity subsequently lowers the LTP probability. An implication of these findings is that activated PF synapses can undergo both LTP and intrinsic plasticity, with the latter reducing the probability of subsequent LTP induction at these and potentially also at neighboring, nonpotentiated synapses.

To examine whether the failure to induce LTP was associated with altered spine calcium signaling, we monitored calcium transients using confocal microscopy. Purkinje cells were loaded with the fluorescent calcium indicator Oregon Green BAPTA-2 (200 μM; excitation wavelength: 488 nm) (Fig. 9B). Intrinsic plasticity was triggered by application of the current injection protocol, and was assessed as described above (spike count) (Fig. 9C). Dendritic calcium transients were elicited by 100 Hz PF stimulation (2–8 pulses) (Fig. 9D) at intervals of typically 2 min (in some experiments 0.5 min). After application of the current injection protocol, spine calcium signaling was enhanced (Fig. 9E,F). The parameter of calcium transients that was most affected was the area under the curve, whereas the peak value was only moderately enhanced (area from 0 to 800 ms: 158.9 ± 25.8%; p = 0.025; peak: 113.6 ± 6.6%; p = 0.02; t = 0–12 min after tetanization; n = 7) (Fig. 9F,G). In the absence of tetanization, calcium transients stayed stable (area: 93.4 ± 12.2%; p = 0.30; peak: 96.0 ± 7.8%; p = 0.32; n = 7) (supplemental Fig. 10, available at www.jneurosci.org as supplemental material). There was a significant difference between changes in the area of spine calcium transients after tetanization and those recorded during the same time period under control conditions (Mann–Whitney U test; area: p = 0.04; peak: p = 0.16). The increase in calcium transients was more pronounced in spines than associated shaft regions, in which an increase was observed as well, but did not reach statistical significance (area: 129.7 ± 25.7%; p = 0.35; n = 7) (Fig. 9H). The change in spine calcium signaling was associated with an increase in intrinsic excitability (167.6 ± 5.4%; p = 0.002; t = 0–12 min after tetanization; n = 7) (Fig. 9C). A caveat of these experiments is that we can only provide a snapshot average of changes in calcium signaling and intrinsic excitability during a relatively short period (12 min) after tetanization. This limitation results from two technical factors: (1) data acquisition was adjusted to low frequencies to minimize light exposure (in most recordings images were taken all 2 min), and the calcium transients require time averaging for analysis, and (2) after 12 min post-tetanization, changes in baseline fluorescence prevented monitoring of comparable calcium transients for longer periods of time. Within this limited time period, the changes in the spike rate and the...
spine calcium transients, respectively, did not show the same kinetics, with the increase in calcium signaling reaching saturation earlier (supplemental Fig. 11, available at www.jneurosci.org as supplemental material). This difference could be due to differences in the underlying cellular mechanisms, or simply differences in the location of these alterations within Purkinje cells. Together, these observations show that intrinsic plasticity is associated with an increase in spine calcium signaling that plateaus quickly after tetanization, and is most prominent for the area under the curve of calcium transients.

Discussion

Our study shows that the intrinsic excitability of Purkinje cells can be upregulated in an activity-dependent way both in vitro and in vivo. Intrinsic plasticity was observed when PF stimulation protocols were applied that induce PF-LTP. However, we could also trigger excitability increases using a current injection protocol that does not elicit LTP. The intrinsic plasticity amplitude varied significantly with the stimulation and recording conditions used. For example, the 1 Hz protocol was very efficient in vitro, but failed to trigger intrinsic plasticity in vivo. The 100 Hz protocol induced intrinsic plasticity in vivo, but the plasticity amplitude observed was relatively small. Two factors might have contributed to this observation: (1) intact inhibition reduces the plasticity amplitude, and (2) in our in vivo recordings the background spike rate was higher (~40 Hz) than in the in vitro recordings (~25–30 Hz; intact inhibition), which might indicate that in the in vitro recordings the available plasticity range was higher to start with. Despite this amplitude variability, it should be stressed that intrinsic plasticity was observed both in vivo and in vitro, with inhibition blocked or intact, at different ages, in rats and in mice, and using different recording conditions, demonstrating that intrinsic plasticity is a robust and physiological phenomenon.

As intrinsic plasticity can be triggered by PF-LTP protocols, we examined whether signaling cascades involved in LTP are shared by intrinsic plasticity. In Purkinje cells, LTP is triggered by low calcium signals and the activation of PKA and CAMKII (Hansel et al., 2006; Jörnfell and Hansel, 2006). Using phosphatase inhibitor drugs and L7-PP2B knockout mice, we observed that PKA and CAMKII are not only involved in LTP induction, but also in intrinsic plasticity. In vestibular nucleus neurons, a similar form of intrinsic plasticity has been described, which results from a downregulation of CAMKII and a reduction of BK-type K channels (Nelson et al., 2005). We did not find a difference in intrinsic plasticity between αCaMKII knock-out mice and wild-type littersmates, suggesting that intrinsic plasticity is governed by a more complex interaction between phosphatases and kinases, such as PKA and CK2, rather than a simple molecular PKA/αCaMKII switch. Accordingly, it is very possible that the various phosphatases and kinases involved have several molecular targets that complement each other in mediating intrinsic plasticity. For example, it has been shown in the hippocampus that PKA regulates the surface expression of SK2 channels (Ren et al., 2006), while CK2 reduces their calcium sensitivity (Allen et al., 2007). It remains to be determined what precise role protein phosphatases play in intrinsic plasticity and how they complement PKA and CK2 in regulating SK2 function.

Using the SK-type K channel antagonist apamin, we could show that a downregulation of SK channels partially mediates the enhancement of Purkinje cell excitability. These data are in line with the observation that intrinsic plasticity was associated with a reduction in the amplitude of the AHP (Fig. 1), which is partially mediated by apamin-sensitive SK conductances (Edgerton and Reinhart, 2003; Womack and Khodakhah, 2003). A caveat of our pharmacological approach is that by changing the membrane excitability, blockade of one type of channel might change the activation probability of other channels as well. It also needs to be pointed out that apamin application did not completely block intrinsic plasticity, suggesting that other conductances might additionally be altered. We therefore examined the involvement of other K conductances that have been implicated in excitability alterations. We were able to show that A-type K conductances and BK-type K conductances do not mediate this form of intrinsic plasticity. A remaining candidate is the Ih current, which can adjust Purkinje cell dendritic integration properties (Nolan et al., 2003; Angelo et al., 2007), but Ih has not been examined in this study.

What are the functional consequences of Purkinje cell intrinsic plasticity? In pyramidal cells, excitability increases and LTP complement each other: a downregulation of A-type K currents enhances the excitability of pyramidal cells and increases the LTP induction probability (Ramakers and Storm, 2002; Watanabe et al., 2002; Chen et al., 2006). LTP, in turn, is associated with enhanced local excitability that decreases the spike threshold (Daoudal et al., 2002; Cudmore and Turrigiano, 2004) and facilitates dendritic spike back-propagation (Frick et al., 2004). Recent studies show that SK channels play a key role in hippocampal LTP: not only does a downregulation of SK channels boost calcium signaling and promote LTP (Stackman et al., 2002; Ngo-Anh et al., 2005; Hammond et al., 2006), but LTP is associated with SK2 channel internalization, contributing to the overall EPSP increase (Lin et al., 2008). It seems that in pyramidal cells, changes in K conductances can locally act as positive regulators of synaptic gain.

A key difference between Purkinje cell intrinsic plasticity and hippocampal plasticity is the effect on subsequent LTP induction. While the enhanced excitability was associated with a small, non-significant increase in PF-EPSPs (Fig. 9A), Purkinje cell intrinsic plasticity lowers the probability for subsequent LTP induction, thus differentially affecting immediate synaptic responses and the ability to subsequently modify synaptic efficacy, respectively. A possible explanation for the reduced LTP induction probability is that intrinsic plasticity is accompanied by enhanced spine calcium signaling, as we demonstrated using confocal microscopy. In Purkinje cells, an amplification of calcium transients would promote LTD rather than LTP induction (Coemans et al., 2004). Thus, the observed failure to induce LTP, or to obtain LTD instead, might result from two factors: (1) the calcium transients were too large for LTP, but too low for LTD induction, or (2) the LTD induction threshold was reached, but LTD was not induced, because of wash-out effects. The calcium signal parameter that was most dramatically altered was the area under the curve, whereas the peak was enhanced to a lower degree. It remains technically possible that changes in the calcium peak amplitude were underestimated because of dye saturation. Nevertheless, the confocal imaging data allow us to conclude that intrinsic plasticity is associated with a significant increase in calcium signaling recruited by PF activity.

Another key difference between consequences of intrinsic plasticity in Purkinje cells and pyramidal cells, respectively, is the observed alteration in the background spike rate. In contrast to pyramidal cells, which show low levels of spontaneous spike activity (Margrie et al., 2002), Purkinje cells are characterized by
high spontaneous discharge rates of 30–80 Hz (Simpson et al., 1996). Increases in this spontaneous spike frequency lower the signal-to-noise ratio, and thus the PF readout (McKay et al., 2007). Here, we observed that Purkinje cell intrinsic plasticity is associated with a lasting increase in the spontaneous spike rate, which did not affect the tonic spike rate of DCN neurons, but lowered the net impact of PF signaling. Since pyramidal cells operate at far lower spontaneous spike frequencies, they cannot similarly control the impact of synaptic signaling by adjusting the background spike rate.

Purkinje cell intrinsic plasticity can be triggered by PF stimulation protocols that also elicit LTP. Strong PF activity will therefore induce LTP at the activated synapses, and the strengthened inputs, as much as neighboring nonpotentiated inputs, will be exposed to the enhanced excitability that accompanies LTP induction. Hippocampal recordings show that enhanced excitability can amplify EPSPs by removing the "brakes" imposed by SK channel activity (Llinas et al., 2008). In our study, we also observed that intrinsic plasticity (current step protocol) slightly enhances EPSPs (Fig. 9A) and the spike output (Fig. 8A,B), but this trend did not reach significance. It seems that, if anything, the enhanced excitability might weakly amplify EPSPs, which would particularly benefit strong, potentiated PF synapses that are likely to reach the spike threshold. In contrast, the weaker, nonpotentiated synapses are more strongly affected by the Purkinje cell-specific intrinsic plasticity features that rather have a negative impact on synaptic gain: (1) intrinsic plasticity lowers the probability for subsequent LTP induction, thus lowering the chance for weaker synapses to be potentiated at a later time point, and (2) the enhanced spontaneous Purkinje cell spike rate lowers the signal-to-noise ratio and reduces the PF readout, which will affect weaker PF synapses more dramatically than stronger ones. The reduced probability for subsequent LTP induction also ensures that the overall excitatory drive stays within limits after a set of synapses was potentiated. In this scenario, intrinsic plasticity closely interacts with LTP to optimally adjust the impact of PF signaling. Available data on cerebellar learning collected in vivo do not allow us to draw conclusions on the relative contribution of synaptic and intrinsic plasticity mechanisms, but provide an outlook on the physiological relevance of a potentiation of synaptic/intrinsic response properties. When applied in vivo, the same 100 Hz PF burst protocol used here causes a dramatic increase in cutaneous PF receptive fields in Purkinje cells, which can be reversed by application of an LTD protocol (CF stimulation) (Jörntell and Ekerot, 2002). It is conceivable that this increase in receptive field size results from PF-LTP (the LTD protocol only triggers a reversal when applied to the same set of PF synapses), but that the strengthened synapses are further amplified by an associated increase in excitability (Jörntell and Hansel, 2006). Moreover, mice with a Purkinje cell-specific knock-out of PP2B show impaired LTP and intrinsic plasticity, but LTD is unaffected. Surprisingly, these mice have severe motor learning deficits, suggesting that both synaptic and intrinsic potentiation mechanisms contribute to cerebellar motor learning (our unpublished data). These observations are in line with previous notions that multiple plasticity mechanisms beyond LTD might be involved in cerebellar learning (Raymond and Lisberger, 1998; Hansel et al., 2001).

References


