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Published in:
Journal of Neuroscience

DOI:
10.1523/JNEUROSCI.3226-10.2010

2010

Link to publication

Citation for published version (APA):

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

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Synopsis of gain control and information storage in neural networks are mediated by alterations in synaptic transmission, such as in long-term potentiation (LTP). Here, we show using both in vitro and in vivo recordings from the rat cerebellum that tetanization protocols for the induction of LTP at parallel fiber (PF)-to-Purkinje cell synapses can also evoke increases in intrinsic excitability. This form of intrinsic plasticity shares with LTP a requirement for the activation of protein phosphatases 1, 2A, and 2B for induction. Purkinje cell intrinsic plasticity resembles CA1 hippocampal pyramidal cell intrinsic plasticity in that it requires activity of protein kinase A (PKA) and casein kinase 2 (CK2) and is mediated by a downregulation of SK-type calcium-sensitive K conductances. In addition, Purkinje cell intrinsic plasticity similarly results in enhanced spine calcium signaling. However, there are fundamental differences: first, while in the hippocampus increases in excitability result in a higher probability for subsequent potentiation and lowers the Purkinje cell output. Second, intrinsic plasticity raises the spontaneous spike frequency of Purkinje cells. The latter effect does not impair tonic spike firing in the target neurons of inhibitory Purkinje cell projections in the deep cerebellar nuclei, but lowers the Purkinje cell signal-to-noise ratio, thus reducing the PF readout. These observations suggest that intrinsic plasticity accompanies LTP of active PF synapses, while it reduces at weaker, nonpotentiated synapses the probability for subsequent potentiation and lowers the impact on the Purkinje cell output.

Introduction

A classic view in neuroscience holds that information storage and learning in neural circuits are made possible by alterations in synaptic transmission such as 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). More recently, it has been shown that activity-dependent alterations in hyperpolarization-activated I\textsubscript{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 (Sausbier 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 Na2HPO4, 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. 1A, B, D, E; picrotoxin: Fig. 1F).

Whole-cell patch-clamp recordings were performed under visual control using a 40× 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-glucurate, 3.48 MgCl2, 10 HEPES, 4 NaCl, 4 NaATP, 0.4 NaGTP, 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 Microlmaging). 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 (2–8 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 calbindin (P2B) 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 (C57BL6/OlaHsd background). Littermates of the following genotypes were used for the experiments: P2B-lox/P2B-lox/L7Cre (L7-PP2B) and P2B-lox/P2B-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 C57BL6/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 stereotaxic 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-amplifier (Molecular Devices). The intracellular electrode solution contained the following (in mM): 134 K-glutamate, 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 *in vivo* might be different than in slices (Alviña et al., 2008). However, the tonic spike frequency ranges reported here are well within the ranges found in awake rat DCN recordings (LeDoux et al., 1998); thus, it is unlikely that tonic PF stimulation affects spontaneous firing in DCN neurons differently *in vivo*. Purkinje cell axons were stimulated using a pair of tungsten microelectrodes located in the white matter around the cerebellar nuclei. The stimulus intensity was adjusted at 0.1 Hz to induce a pause in the spontaneous firing of DCN neurons. Kynurenic acid (3 μM) was bath applied to block excitatory neurotransmission. Repetitive stimulation was applied at 33.3 and 50 Hz. The switch from 33.3 to 50 Hz was applied after 2–4 min of stimulation at 33.3 Hz. Recordings were digitized and stored using programmable software (Spike 2, Cambridge Electronic Design). 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 dissected, 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-calbindin (Sigma) 1:1000. Subsequently, sections were rinsed in TBS, incubated with FITC- or Cy3- (Jackson ImmunoResearch) conjugated goat-anti-mouse or goat-anti-rabbit secondary antibodies at 1:200 for 1.5–2 h at room temperature. Sections were rinsed again in TBS and mounted directly on coverslips. Fluorescence images were taken with a Zeiss LSM 510 confocal laser-scanning microscope. The rabbit anti-SK2 channel antibodies obtained from both Alomone and Sigma were directed against a sequence in the C-terminal domain of SK2 channel subunits, which corresponds to amino acids 542–559. The immunostainings obtained with both antibodies were very similar, which is why we only show stainings obtained with the anti-SK2 antibody purchased from Alomone Labs (see Fig. 4). The specificity of this antibody was confirmed using slices from SK2+/− mice and littermate controls (supplemental Fig. 6, available at www.jneurosci.org as supplemental material) (for protocol details, see Pichon et al., 2007).

**Data analysis.** Data obtained from the Purkinje cell recordings *in vivo* were analyzed using Pulsefit (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). Statistical 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 = 0.0006) (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 = 0.02895) (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 = 0.0006), suggesting that spontaneous GABAergic transmission limits intrinsic plasticity. A pronounced increase in the number of depolarization-evoked spikes was also observed following PF tetanization (1 Hz, 5 min: 645.0 ± 113.9%; n = 9; p = 0.00138) (Fig. 1D). This 1 Hz PF stimulation protocol has previously been used for PF-LTP induction (Coensmans et al., 2004), and reliably elicited LTP un-
der 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 = 0.00024) (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 GABA_A 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-
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 \textit{in vitro} (Ramakers and Storm, 2002; Frick et al., 2004). When 4-AP was bath applied (100 \muM), 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: 180.81 ± 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 IA 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: 0.35 ± 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 IA 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 (Sourdret 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 ibetoxin. 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 ibetoxin (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>Condition</th>
<th>Amplitude (mV)</th>
<th>AHP (mV)</th>
<th>Em after spike (mV)</th>
<th>10%-90% (ms)</th>
<th>FWHM (ms)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Control before IP</strong></td>
<td>91.6 ± 5.3</td>
<td>-6.9 ± 1.2</td>
<td>-49.6 ± 1.6</td>
<td>0.198 ± 0.015</td>
<td>0.288 ± 0.028</td>
</tr>
<tr>
<td><strong>Control after IP</strong></td>
<td>89.9 ± 4.5</td>
<td>-6.1 ± 2.1</td>
<td>-48.4 ± 1.6</td>
<td>0.186 ± 0.015</td>
<td>0.276 ± 0.022</td>
</tr>
<tr>
<td><strong>Before 4AP</strong></td>
<td>87.7 ± 6.9</td>
<td>-4.9 ± 1.6</td>
<td>-52.0 ± 2.3</td>
<td>0.180 ± 0.001</td>
<td>0.270 ± 0.026</td>
</tr>
<tr>
<td><strong>Under 4AP</strong></td>
<td>88.3 ± 3.2</td>
<td>9.6 ± 1.9</td>
<td>-54.1 ± 2.1</td>
<td>0.190 ± 0.009</td>
<td>0.435 ± 0.039</td>
</tr>
<tr>
<td><strong>Under 4AP after IP</strong></td>
<td>87.6 ± 8.4</td>
<td>10.6 ± 3.2</td>
<td>-54.4 ± 2</td>
<td>0.188 ± 0.007</td>
<td>0.465 ± 0.043</td>
</tr>
<tr>
<td><strong>Before HpTX-2</strong></td>
<td>76.1 ± 5.5</td>
<td>-7.1 ± 1.5</td>
<td>-59.9 ± 1.3</td>
<td>0.210 ± 0.019</td>
<td>0.320 ± 0.040</td>
</tr>
<tr>
<td><strong>Under HpTX-2</strong></td>
<td>74.4 ± 4.5</td>
<td>-6.4 ± 1.4</td>
<td>-57.4 ± 3.1</td>
<td>0.220 ± 0.017</td>
<td>0.340 ± 0.041</td>
</tr>
<tr>
<td><strong>Under HpTX-2 after IP</strong></td>
<td>75.3 ± 5.4</td>
<td>-6.8 ± 1.4</td>
<td>-58.8 ± 2.8</td>
<td>0.215 ± 0.024</td>
<td>0.335 ± 0.045</td>
</tr>
<tr>
<td><strong>Before apamin</strong></td>
<td>98.4 ± 5.4</td>
<td>-7.8 ± 2.2</td>
<td>-51.9 ± 1.3</td>
<td>0.180 ± 0.013</td>
<td>0.220 ± 0.018</td>
</tr>
<tr>
<td><strong>Under apamin</strong></td>
<td>98.8 ± 4.1</td>
<td>-7.5 ± 2.2</td>
<td>-50.2 ± 1.3</td>
<td>0.175 ± 0.014</td>
<td>0.235 ± 0.016</td>
</tr>
<tr>
<td><strong>Under apamin after IP</strong></td>
<td>100.6 ± 5.7</td>
<td>-7.1 ± 1.9</td>
<td>-49.9 ± 1.5</td>
<td>0.165 ± 0.010</td>
<td>0.230 ± 0.115</td>
</tr>
</tbody>
</table>

Parameters of the action potential waveform that were measured 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 (dV/dt, measured 4.5–6 ms after the spike peak). The table summarizes these parameters (see also Fig. 3D) 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 (Coemsans 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
There was no statistical difference between these two groups as the molecular switch mechanism involving PP1, 2A, and 2B as well involves an activation of protein phosphatases 1, 2A, and 2B.

Intrinsic excitability changes triggered by current injection or PF sensitization (80.6 ± 5.2%; n = 12; p = 0.00179) (Fig. 6D), likewise, the current injection protocol was ineffective in the presence of okadaic acid (1 μM) which inhibited in slices prepared from L7-PP2B knock-out mice (open circles; n = 6), but was inhibited in slices prepared from L7-PP2B knock-out mice (open circles; n = 9). B. Application of the 1 Hz PF stimulation protocol failed to induce excitability changes in the presence of okadaic acid (1 μM; open circles; n = 8) and cyclosporin A (100 μM; open circles; n = 16). 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 mV) 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 = 4), and the PP2B inhibitor cyclosporin A (100 μM; open circles; n = 8). C, Intrinsic plasticity was intact in slices prepared from wild-type mice (closed circles; n = 16), but was inhibited in slices prepared from L7-PP2B knock-out mice (open circles; n = 9). D. Application of the 1 Hz PF-stimulation protocol failed to induce excitability changes in the presence of okadaic acid (1 μM; open circles; n = 8) and cyclosporin A (100 μM; open triangles; n = 16). Error bars indicate SEM.

in the presence of the PP2B inhibitor cyclosporin A (100 μM; 98.3 ± 4.7%; n = 16; p = 0.72) (Fig. 5D). These data suggest that intrinsic excitability changes triggered by current injection or PF sensitization share the same induction mechanism, which involves an activation of protein phosphatases 1, 2A, and 2B.

At PF synapses, bidirectional synaptic plasticity is governed by a molecular switch mechanism involving PP1, 2A, and 2B as well as the α isofrom of calcium/calmodulin-dependent kinase II (αCaMKII) (Jörntell and Hansel, 2006). To examine whether αCaMKII is involved in intrinsic plasticity, we applied the current injection protocol to Purkinje cells in slices prepared from αCaMKII knock-out mice (Elgersma et al., 2002), and wild-type controls. Intrinsic plasticity was observed in αCaMKII knock-out mice (119.7 ± 12.8%; n = 12; p = 0.62) (Fig. 6C). As in wash-in control experiments emodin application affected the spike rate in the absence of sensitization (supplemental Fig. 7F, available at www.jneurosci.org as supplemental material), we also used the more selective CK2 inhibitor 2-dimethylamino-4,5,6,7-tetrabromo-1-H-benzimidazole (DMAT). In the presence of DMAT, intrinsic plasticity was blocked as well (DMAT; 5 μM: 106.9 ± 12.8%; n = 8; p = 0.023; 10 μM: 113.5 ± 9.8%; n = 6; p = 0.30) (Fig. 6D), suggesting that active CK2 is required for intrinsic plasticity.

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 (Häsuer and Clark,
1997), thus conveying little information on the activity state of synaptic inputs. Purkinje cells form inhibitory synapses with their DCN target cells (Ito, 1984), which on their own are spontaneously active (Jahnsen, 1986). We thus examined whether the spontaneous discharge rate of DCN neurons was affected by the enhanced background spike rate of Purkinje cells. Whole-cell patch-clamp recordings in current-clamp mode were obtained from DCN neurons in vitro, and Purkinje cell axons were stimulated in the white matter (Pedroarena and Schwarz, 2003). Two different sets of experiments were performed: In a first set of experiments, the resulting IPSPs (16.0 ± 1.9 mV; \( n = 7 \)) consistently evoked a pause in spontaneous DCN firing (Fig. 7A). In the second set of experiments, higher stimulus intensities were used to examine whether an enhanced inhibitory input (reflecting a higher number of simultaneously active Purkinje cells, 24.6 ± 6.2 mV; \( n = 8 \); \( p = 0.017 \); Mann–Whitney \( U \) test) affects DCN neurons more efficiently (Fig. 7B). Purkinje cell spontaneous activity was mimicked by protracted (2–4 min) 30 Hz activation of Purkinje cell axons, allowing the synapses to reach stationary levels of short-term depression (Telgkamp and Raman, 2002; Pedroarena and Schwarz, 2003). Intrinsic plasticity was simulated by a switch from 30 to 50 Hz activation. These frequencies were selected to mimic the increase in the spike rate observed following application of intrinsic plasticity protocols in vitro (Fig. 2C–F). Activation at 30 Hz resulted first in a transient cessation of DCN firing [Fig. 7A (insets), B (top)], followed by discharge at a reduced rate. However, there was no further reduction upon switch to 50 Hz, using either low or high stimulus intensities (low 30 Hz: 95.2 ± 2.8% low 50 Hz: 93.6 ± 4.6%; \( n = 7 \); \( p = 0.68 \); high 30 Hz: 77.4 ± 3.8%; high 50 Hz: 79.5 ± 5.0%; \( n = 8 \); \( p = 0.46 \); Wilcoxon signed-rank test) (Fig. 7). The observation that the DCN spike rate was not affected by an enhanced frequency of inhibitory Purkinje cell output is possibly explained by the filtering properties of synapses depressed to stationary levels (Pedroarena and Schwarz, 2003), since DCN neurons are affected by the frequency of inhibitory inputs if these are activated for shorter periods (Telgkamp and Raman, 2002), or if depression is lacking (Gauck and Jäger, 2000). To assess the degree of depression, IPSPs were averaged over a 1 min period before and after the switch from 30 to 50 Hz stimulation. At 30 Hz stimulation, IPSPs were reduced to 8.8 ± 1.9% at high stimulus intensity (\( n = 8 \)), and to 9.7 ± 1.4% at low stimulus intensity (\( n = 7 \)) (Fig. 7C). The strong depression at 30 Hz stimulation correlates with the moderate effect of tonic Purkinje cell activation on the firing rate of DCN neurons. At 50 Hz stimulation, IPSPs were further reduced to almost half of the levels found at 30 Hz: 5.6 ± 1.5% at high stimulus intensity (\( n = 8 \)), and to 5.1 ± 1.1% at low stimulus intensity (\( n = 7 \)) (Fig. 7C). At both intensities, this further reduction in IPSP amplitudes upon switch from 30 to 50 Hz stimulation was significant (high intensity: \( p = 0.022 \); low intensity: \( p = 0.016 \); Wilcoxon signed-rank test) and present in all neurons examined. Depression of IPSPs during a train has previously been described for shorter stimulus trains in the DCN (Telgkamp and Raman, 2002, Pedroarena and Schwarz, 2003). Together, our DCN recordings suggest that the enhanced frequency of Purkinje cell pacemaker activity, which is associated with intrinsic plasticity, does not have a direct effect on the spontaneous spike rate of DCN neurons. The results also point to the stronger depression of inhibitory Purkinje cell–DCN neuron synapses as a mechanism to counterbalance the inhibitory effect of enhanced IPSP frequencies.

**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
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 over a range of 50% led to variations in the number of evoked spikes ranging from one to four spikes (n = 11) (Fig. 8B). This observation suggests a limited control of the spike output of Purkinje cells by LTD and LTP, respectively. After repeated current injection, the spike output in all three intensity groups was slightly enhanced, but this trend did not reach statistical significance (n = 8–9; 25%: p = 0.90; control: p = 0.53; +25%: p = 0.60; paired Student’s t test) (Fig. 8B), suggesting that intrinsic plasticity has at most a weak effect on the spike output. We obtained similar results when analyzing the spike frequency, rather than the number of evoked spikes. We calculated increases in the spike frequency resulting from PF stimulation as the ratio of the frequency recorded from 3 spikes right after stimulation to the frequency recorded before. Increasing stimulus strength enhanced this frequency ratio (−25%: 1.5 ± 0.1; n = 11; control: 2.2 ± 0.3; n = 11; +25%: 2.9 ± 0.4; n = 10). After tetanization, these ratios were not significantly changed (−25%: 1.7 ± 0.4; p = 0.48; n = 9; +25%: 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–I). 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 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 LTP and LTD 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 stimulation 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 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 = 5). 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,G). 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 increase 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 vitro*, 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 vitro* and *in vivo*, 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 PP1, 2A, and 2B, whereas LTD is evoked by high calcium signals and the activation of PKC and CaMKII (Hansel et al., 2006; Jörntell and Hansel, 2006). Using phosphatase inhibitor drugs and L7-PP2B knock-out mice, we observed that PP1, 2A, and 2B 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 CaMKII knock-out mice (Hansel et al., 2006) and wild-type littermates, 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 PP1/CaMKII 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 

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, nonsignificant 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 (Lin 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 costimulation) (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 P2P2 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


