Cerebellar ataxia and Purkinje cell dysfunction caused by Ca\textsuperscript{2+}-activated K\textsuperscript{+} channel deficiency

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Malfunctions of potassium channels are increasingly implicated as causes of neurological disorders. However, the functional roles of the large-conductance voltage- and Ca\textsuperscript{2+}-activated K\textsuperscript{+} channel (BK channel), a unique calcium, and voltage-activated potassium channel type have remained elusive. Here we report that mice lacking BK channels (BK\textsuperscript{−/−}) show cerebellar dysfunction in the form of abnormal conditioned eye-blink reflex, abnormal locomotion and pronounced deficiency in motor coordination, which are likely consequences of cerebellar learning deficiency. At the cellular level, the BK\textsuperscript{−/−} mice showed a dramatic reduction in spontaneous activity of the BK channel in cerebellar Purkinje neurons, which generate the sole output of the cerebellar cortex and, in addition, enhanced short-term depression at the only output synapses of the cerebellar cortex, in the deep cerebellar nuclei. The impairing cellular effects caused by the lack of postsynaptic BK channels were found to be due to depolarization-induced inactivation of the action potential mechanism. These results identify previously unknown roles of potassium channels in mammalian cerebellar function and motor control. In addition, they provide a previously undescribed animal model of cerebellar ataxia.

Potassium channels are the largest and most diverse class of ion channels underlying electrical signaling in the brain (1). By causing highly regulated, time-dependent, and localized polarization of the cell membrane, the opening of K\textsuperscript{+} channels mediates feedback control of excitability in a variety of cell types and conditions (1). Consequently, K\textsuperscript{+} channel dysfunctions can cause a range of neurological disorders (2–6), and drugs that target K\textsuperscript{+} channels hold promise for a variety of clinical applications (7).

Among the wide range of voltage- and calcium-gated K\textsuperscript{+} channel types, one stands out as unique: the large-conductance voltage- and Ca\textsuperscript{2+}-activated K\textsuperscript{+} channel (BK channel, also termed M\textsubscript{o} or Maxi-K) differs from all other K\textsuperscript{+} channels in that it can be activated by both intracellular Ca\textsuperscript{2+} ions and membrane depolarization (8). These channels are widely expressed in central and peripheral neurons, as well as in other tissues (9), and are regarded as a promising drug target (10). However, the functions of the BK channels in vivo have not previously been directly tested in any vertebrate species. We therefore decided to examine the functions of these channels by inactivating the gene encoding the pore-forming channel protein.

Methods
A complete description of the methods is given in Supporting Methods, which is published as supporting information on the PNAS web site.

Generation of BK Channel α Subunit-Deficient Mice. In the targeting vector (Fig. 5, which is published as supporting information on the PNAS web site), the pore exon was flanked by a single loxP site and a floxed neo/ko cassette. Correctly targeted embryonic stem cells were injected into C57BL/6 blastocysts and resulting chimeric mice were mated with C57BL/6. Homozygous BK-deficient mice (F\textsuperscript{2} generation) were produced. Either litter- or age-matched WT and BK\textsuperscript{−/−} mice on a hybrid SV129/C57BL6 background (always F\textsuperscript{2} generation) were randomly assigned to the experimental procedures, in keeping with German legislation on the protection of animals.

Brain in Situ Hybridization and Immunohistochemistry. BK channel α subunit mRNA transcript antisense probes complementary to the pore exon were labeled with [α-\textsuperscript{35}S]dATP to a specific activity of 10\textsuperscript{8} cpm/μg. Sections (15 μm) were cut on a cryostat and fixed in 4% paraformaldehyde. Sections were prehybridized with hybridization buffer containing yeast tRNA and salmon sperm DNA and exposed to labeled probe (5,000 cpm/μl) in hybridization buffer. Sections were washed and exposed for 28 days.

Immunohistochemistry was performed by using cryostat slices from perfused and postfixed brains. Coronal cerebellar were incubated with anti-BKα\textsubscript{674–1115} tagged with a peroxidase-conjugated goat anti-rabbit IgG.

Motor Function, Motor Learning, and Footprint Patterns. For analyzing balance and motor coordination, 12 male and 12 female mice of each genotype (4–6 months) were tested on the accelerating rotarod (11); the rotational speed increased from 4 to 40 rpm over 5 min. Mice were trained for 3 days with five trials per day before latency to fall was recorded.

There were no significant differences in the ability of the mice to traverse a graded beam with smooth or irregular surface was assessed (12). Before testing, the mice were trained for 3 days with four trials per day.

Hindpaws and forepaws of six WT and seven BK\textsuperscript{−/−} mice (litter- or age-matched, 3–6 months) were dipped in red and blue watercolor, respectively, before walking on paper (12). Mice were trained on 3 consecutive days with 3 trials per day. Footprint patterns were analyzed for stride basis, stride length, and paw abduction.

Conditioned Eye Blink. Eye-blink conditioning (13) from 8 WT and 8 BK\textsuperscript{−/−} mice (8–12 weeks) was performed with a tone (5 kHz) as conditioned stimulus (CS) and an air puff (10 psi) as uncond-

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Abbreviations: BK channel, large-conductance voltage- and Ca\textsuperscript{2+}-activated K\textsuperscript{+} channel; AHP, afterhyperpolarization; PC, Purkinje cell; AP, action potential; DCN, deep cerebellar nuclei; IBTX, iberiotoxin.

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ditioned stimulus (US). The US was coterminated with the CS. Conditioning (each session consisting of 90 paired CS-US trials and 10 CS trials alone) was performed successively for 5 days and extinction (each session consisting of 20 CS alone) for 3 days.

**Electrophysiological Analysis of Cerbellar Purkinje Cells (PCs).** Sagittal slices (350–400 µm thick) from the cerebellar vermis of 4- to 5-week-old WT and BK−/− mice were prepared and kept at 34–36°C in artificial cerebrospinal fluid with 10 µM bicuculline free base [a concentration that has been shown to not measurably affect currents of Ca2+–activated K+ channels with small conductance (SK) or afterhyperpolarizations (AHPs) in rat brain slices (14, 15) and to cause only a ∼20% inhibition of BK channels expressed in frog oocytes (16)] and 10 µM 6,7-dinitroquinoxaline-2,3-dione (D600) to block synaptic transmission. Whole-cell (Axoclamp 2A, Axon Instruments, Union City, CA) and extracellular (Multiclamp 700A, Axon Instruments) recordings were obtained from PC somata under visual control. To compare the first action potential (AP) and AHP evoked by a depolarizing current pulse, a weak hyperpolarizing DC was injected to silence the cell before testing.

**Recording of Inhibitory Synaptic Potentials in Deep Cerebellar Nuclei (DCN).** Cerbellar slices from 13- to 17-day-old mice were prepared and superfused with artificial cerebral spinal fluid at room temperature. Whole-cell recordings (Axopatch 1-D) from large DCN neurons (above 15 µm, hence presumably glutamatergic projecting neurons) in the lateral or medial nuclei, with 4 mM kynurenic acid blocking excitatory amino acid neurotransmission. PC axons were stimulated with a pair of tungsten microelectrodes, with different interstimulus intervals. Paired-pulse depression was quantified as the ratio between the average peak amplitudes of the responses to second and first stimuli.

**Results**

**Generation of BK−/− Mice.** The BK channel consists of four α subunits and four optional auxiliary β subunits (17). The pore-forming α subunit is encoded by only a single gene, KCNMA1 (also called Slo), from which multiple splice isoforms are generated (18, 19), whereas there are four different β subunit genes with tissue-specific expression (20–23). To determine BK channel functions, we generated mice lacking functional BK channels. The pore exon of the α subunit, which also encodes part of the S6 segment, was deleted through homologous recombination (Fig. 5). We obtained BK (mSlo) channel-deficient mice (BK−/−) that completely lack the BKα mRNA and protein (Fig. 1a and b). In the cerebellum, in situ hybridization experiments (Fig. 1a) indicate that mRNA encoding the BKα subunit is predominantly observed in PC somata, whereas only very low message levels are detected in the molecular layer, the granule cell layer, and the DCN, respectively. In clear contrast, BK channel protein (Fig. 1 b and c) is observed in PC somata, the molecular layer (where PC dendrites arborize), and the DCN (where PC axons project to). This finding suggests that the expression of BK channels in the cerebellum is predominantly, but not exclusively, in PCs where they are likely to be targeted to both the somato-dendritic and axonal compartments. Other principal neurons of the cerebellum express BK channels at levels considerably below that of PCs (24).

Macroscopic and microscopic analysis (Nissl staining) did not reveal any morphological abnormalities in young or adult BK−/− brains. The mutant mice had a normal life expectancy compared to their WT littermates but showed obvious ataxia. Furthermore, at 4 and 8 weeks of age, the BK−/− mice (males and females) showed 15–20% smaller body length and weight compared to their WT littermates, but the length and weight became normal at 12 weeks of age (Fig. 6, which is published as supporting information on the PNAS web site). The mutant mice also showed moderate vascular dysfunctions (∼10% increase in arterial blood pressure and changes in its regulation). These vascular effects, which were found to be due to BK channel deficiency in vascular smooth muscle and clearly unrelated to the neurological deficits, will be described elsewhere.

**Motor Impairment in BK−/− Mice.** The BK−/− mice exhibited intention tremor and abnormal gait (12). Thus, the mutant mice showed shorter strides and irregular step pattern as compared with the WT mice (Fig. 1c). The angle between the left and right hindpaw axes was almost doubled, whereas the hindpaw width was similar between WT and mutants. These alterations were seen at all ages and in both genders.

The motor impairment precluded common tests of spatial learning such as the water maze. Thus, the mutants showed reduced swim speed and more frequent floating. To test motor coordination and balance, we used beam walking (12). The BK−/− mice were reluctant to traverse the beam (Fig. 2a), and, if trying, they made more slips and hindpaw errors and often fell off the beam or off the start platform, thus indicating severe motor impairment. In the accelerating rotarod test, which also requires good sensorimotor coordination and is sensitive to cerebellar and basal ganglia dysfunction (11), the BK−/− mice of both genders showed a strongly reduced latency to fall (Fig. 2b).

Despite starting at a much lower level, the mutants improved at...
Abnormal Eye-Blink Reflex in BK<sup>−/−</sup> Mice. The motor impairment prompted us to examine cerebellar function. The cerebellum adjusts the operation of motor centers in the cortex and brainstem during movements and is needed for balance, precision timing, and sensorimotor learning, functions that appeared to be affected in the mutants. To test cerebellar function, we adopted the conditioned eye-blink response, a well established behavioral test of cerebellar learning (13). An air puff to the eye elicits the eye-blink reflex, which is abnormal in the BK<sup>−/−</sup> mice. Statistical summary of conditioned eye blinking results from eight WT and eight BK<sup>−/−</sup> mice. The conditioning phase includes 5 days followed by 3 days for extinction. All data are means ± SEM, **, P < 0.01.

Fig. 2. Motor impairment and abnormal conditioned eye blinking in BK<sup>−/−</sup> mice. (a) Beam walking performance at regular (reg; i.e., smooth) and irregular (irreg; i.e., with steps) beams: distance moved, hindpaw slips, and falls. Means of two trials per beam, n = 6 per gender. (b) Accelerating rotarod performance: time to fall off the rotarod (mean value per trial), n = 6 per gender. (c) The conditioned eye-blink reflex, a cerebellum-specific form of motor learning, was abnormal in the BK<sup>−/−</sup> mice. Statistical summary of conditioned eye blinking results from eight WT and eight BK<sup>−/−</sup> mice. The conditioning phase includes 5 days followed by 3 days for extinction. All data are means ± SEM, *, P < 0.05; **, P < 0.01.

a rate similar to the WT (Fig. 2b), indicating that they partially compensated their deficit through motor learning. The BK<sup>−/−</sup> mice showed motor impairment also in the open field test: reduced distance, path linearity, and exploration index, as well as lack of typical acceleration when leaving the center field (data not shown).

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In contrast, the BK<sup>−/−</sup> mice showed no learning of the conditioned eye blink. Instead, they showed increased eye blinking up to the highest level observed in trained WT mice, suggesting maximal disinhibition of the interpositus nucleus. These results support the hypothesis of cerebellar dysfunction in the BK<sup>−/−</sup> mice.

Depolarization Block of Cerebellar PCs. To search for a possible cellular basis of cerebellar dysfunction in the BK<sup>−/−</sup> mice, we recorded the firing activity of PCs. We focused on this cell type because projections from these cells to the DCN are the only output from the cerebellar cortex. Because BK channels may be involved in AP repolarization and fast AHPs (25, 26), we compared AP waveforms and AHPs in cerebellar slices. To suppress spontaneous discharge and ensure comparable recording conditions, we maintained the pretest membrane potential at a constant level (−60 to −70 mV) by direct current injection, and evoked APs by depolarizing current pulses. In PCs from WT mice, the BK channel blocker iberiotoxin (IbTx, 1 μM) (1) strongly suppressed the single-spike AHP and slowed AP repolarization (Fig. 3a). In contrast, IbTx had no measurable effect on AHPs or APs in PCs from BK<sup>−/−</sup> mice. Furthermore, the AHP amplitude was significantly smaller in BK<sup>−/−</sup> (4.5 ± 0.8 mV, n = 14) vs. WT neurons (10.1 ± 1.8 mV, n = 13; P < 0.01). The mean AP 90–10% decay time was slightly but not significantly longer in BK<sup>−/−</sup> cells (0.27 ± 0.09 ms, n = 5, vs. 0.21 ± 0.01 ms in WT, n = 5), suggesting that there may be compensatory changes in other repolarizing mechanisms, or that the difference was too small to be detected statistically. Taken together, these observations suggest that the main role of BK
channels in PCs is to generate AHPs (26), with only a modest effect on the AP itself (Fig. 3a).

Normal PCs fire spontaneous APs under basal conditions, thereby tonically inhibiting the DCN (27). Surprisingly, >50% of the PCs from BK−/− mice lacked spontaneous discharge (8 of 15 cells, 53.3%), in sharp contrast to the WT cells, which all generated spontaneous APs during whole-cell recording (15 of 15 cells; Fig. 3c Left). Thus, the overall discharge activity was abnormally low in mutant PCs. The average spike frequency for all cells was ~6-fold higher in WT cells (55.2 ± 8.9 Hz) than in BK−/− cells (9.5 ± 5.6 Hz; P < 0.01) (Fig. 3c Right). PCs from both WT and BK−/− mice showed two distinct spontaneous firing patterns (28, 29): tonic-firing PCs vs. repetitive bursting PCs (Fig. 3b) and the proportions of these were not statistically different between BK−/− (20.0% tonic, 26.7% bursting) and WT mice (53.3% tonic, 46.7% bursting) (Fig. 3c Left).

To avoid possible artifacts due to dialysis via the recording pipette, we also performed somatic loose-patch extracellular recordings from PCs (30). Again, the overall spontaneous firing rate was far lower in the BK−/− (8.7 ± 3.6 Hz) than in the WT cells (65.7 ± 10.1 Hz), mainly due to more silent cells (Fig. 3c Right). In addition, among the tonic-firing PCs, the BK−/− cells showed longer interspike intervals (Fig. 3a Top) and hence lower discharge frequency (37.1 ± 7.2 Hz, n = 4) than WT cells (91.6 ± 15.8 Hz, n = 8; P < 0.01; not shown). The frequency distributions and cumulative frequency plots of the spontaneous firing rates also showed highly significant differences between BK−/− and WT cells, for both whole-cell and loose-patch recordings. In both cases, the BK−/− PC firing was skewed toward frequencies <20 Hz (Fig. 7, which is published as supporting information on the PNAS web site). Taken together, PCs from mice lacking BK channels showed an abnormally reduced spontaneous discharge activity.

Why were these mutant PCs silent? The whole-cell data showed that the resting membrane potential of the silent cells stayed at a depolarized level (~45 to ~50 mV; Fig. 4b Left). In most cases it was constant, but some (two of eight) cells showed slow shallow oscillations (0.2–0.5 Hz, 15- to 20-mV amplitude; not shown). Depolarizing current injections in silent mutant cells failed to evoke APs. However, after being hyperpolarized to ~70 or ~80 mV by current injection, each of these silent BK−/− cells could fire normal fast APs (Fig. 4b Right), indicating that their silence was caused by depolarization block, i.e., inactivation of their Na+ channels (31). Current-voltage (I/V) plots (V measured near the end of 500-ms-long injected current pulses) were nearly identical (<1-mV difference) for BK−/− (n = 11) and WT (n = 9) PCs at potentials <~70 mV, whereas at >~65 mV, the mutant PCs showed on average 2- to 5-mV larger subthreshold depolarizations than WT cells. Although this difference was not statistically significant, it may possibly reflect a change in subthreshold currents (Fig. 8, which is published as supporting information on the PNAS web site).

As expected, IbTx had no effects in mutant PCs. In the tonic WT cells, IbTx increased the discharge frequency (26) and, in some PCs, eventually precipitated a transition into bursting mode. In bursting WT cells, IbTx induced complex changes in the burst patterns (not shown). However, neither tonic nor bursting WT cells became steadily depolarized or silent during the recordings in IbTx (lasting 15–45 min). Whatever the mechanism, the lack of BK channels evidently reduced the PCs basal discharge activity to only a fraction of the normal level (Fig. 3c).

Increased Short-Term Depression at the PC/DCN Synapses. The results so far suggest that dramatic reduction of PC activity leads to a disinhibition of DCN. To assess the inhibitory effect of the impulses that were still generated in PCs, we performed whole-cell recordings from DCN neurons from BK−/− and WT mice. Stimulation of PC axons (while excitatory receptors were blocked) evoked inhibitory postsynaptic currents (IPSCs) that could be blocked by GABAAR antagonists. The IPSCs in WT and BK−/− DCNs had similar rise times, half widths, and amplitudes, without any significant difference between the two genotypes (Fig. 9, which is published as supporting information on the PNAS web site).

If BK channels contribute to presynaptic APs or AHPs, differences in neurotransmitter release might become evident during high-frequency activation, corresponding to the high firing frequencies of PCs in vivo. Therefore, we used paired stimulation to test the frequency dependence of transmission at the PC/DCN synapses. As illustrated in Fig. 4c, the paired-pulse depression, which is also seen in normal PCs (32), was significantly increased in DCNs from BK−/− mice, but only for intervals shorter than 100 ms. Thus, BK channels apparently regulate high-frequency synaptic transmission. Possibly, the absence of BK channels impaired repolarization and/or AHP after the first spike, thus causing incomplete deinactivation of Na+ channels, reducing the subsequent spike amplitude and Ca2+ influx, thereby causing synaptic depression. Taken together, the attenuated transmission at PC/DCN synapses and the reduced activity of PCs are expected to synergistically diminish the GABAergic outflow from the cerebellar cortex, thus causing disinhibition of the DCN which, in turn, may explain the observed motor deficits.

Discussion

Deletion of the BK channel α subunit in mice permitted the first identification of the physiological functions of this unique channel type in a vertebrate species in vivo and revealed unexpected roles of BK channels in cerebellar function. The consequences of BK channel ablation exceed by far those caused by deletion of
the regulatory BK channel β subunit, which is not significantly expressed in neurons and produced no neural effects (33, 34). Nevertheless, the BK channels, despite their widespread expression in the CNS (24), appear to play surprisingly modest roles in normal brain function. This suggests that their wide neuronal expression has evolved partly because BK channels are more important under stress and deleterious conditions, a view supported by the observation that BK channels improve the survival of neurons exposed to ischemic conditions (35).

A principal finding of this study is that the mice lacking BK channels showed apparent loss of eye-blink conditioning. This learning behavior depends on the cerebellar circuitry and not on the basal ganglia or cerebral motor cortex (36). The naïve BK−/− mice showed already from the first training session an abnormally high blinking rate, suggesting a profound disinhibition of the DCN. This could be caused by the observed severe suppression of spontaneous PC activity, combined with the increased short-term depression at the inhibitory PC/DCN synapses. Although evaluation of eye-blink learning is not possible when the blinking rate is already close to maximal in the training phase, this impairment suggests cerebellar dysfunction. Hence the motor learning of BK−/− mice on the rotarod may involve mainly noncerebellar functions, e.g., corticostriatal plasticity (37).

The severe suppression of PC activity and synapses observed in the BK−/− mice is therefore a likely cause of their motor coordination deficits and ataxia. PCs are essential for cerebellar motor control (38–40) by providing specific timing signals for movement coordination (41). BK channels in PCs are activated by depolarization and Ca2+ influx, via opening of P/Q type Ca2+ channels during APs (26, 31) and contribute to AP repolarization and AHPs. The depolarization block caused by the BK channel deletion suggests that the net effect of P/Q channel activation is to polarize the membrane via activation of BK channels, thus maintaining normal excitability (31). The failure of IbTx applications in WT slices to mimic the depolarization block found in mutants may be due to incomplete block of toxin-resistant BK channels (21), or perhaps longer-lasting suppression of BK channels is needed to induce the silent state, possibly inducing compensatory changes in other membrane conductances. Interestingly, loss of P/Q channel function, due to various mutations, can cause cerebellar ataxia in humans (42). Our results suggest that these forms of inherited cerebellar ataxia may be partly due to lack of BK channel activation. Thus, the BK−/− mice, which represent a previously undescribed animal model of cerebellar ataxia, may prove useful for understanding normal and pathological cerebellar function, and studies of BK channel-dependent functions in vivo are likely to help clarify the therapeutic utility of BK channel activators or blockers (9).

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