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**Cognitive deficits and impaired hippocampal long-term potentiation in K<sub>ATP</sub>-induced DEND syndrome**

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ATP-sensitive potassium (K<sub>ATP</sub>) gain-of-function (GOF) mutations cause neonatal diabetes, with some individuals exhibiting developmental delay, epilepsy, and neonatal diabetes (DEND) syndrome. Mice expressing K<sub>ATP</sub>-GOF mutations pan-neuronally (nK<sub>ATP</sub>-GOF) demonstrated sensorimotor and cognitive deficits, whereas hippocampus-specific hK<sub>ATP</sub>-GOF mice exhibited mostly learning and memory deficiencies. Both nK<sub>ATP</sub>-GOF and hK<sub>ATP</sub>-GOF mice showed altered neuronal excitability and reduced hippocampal long-term potentiation (LTP). Sulfonylurea therapy, which inhibits K<sub>ATP</sub>, mildly improved sensorimotor but not cognitive deficits in nK<sub>ATP</sub>-GOF mice. Mice expressing K<sub>ATP</sub>-GOF mutations in pancreatic β-cells developed severe diabetes but did not show learning and memory deficits, suggesting neuronal K<sub>ATP</sub>-GOF as promoting these features. These findings suggest a possible origin of cognitive dysfunction in DEND and the need for novel drugs to treat neurological features induced by neuronal K<sub>ATP</sub>-GOF.

**Results**

Pan-Neuronal Expression of K<sub>ATP</sub>-GOF Mutations Associates with DEND-Like Sensorimotor Deficits. To study the neurological features of DEND, we generated pan-neuronal nK<sub>ATP</sub>-GOF mice (SI Appendix, Fig. S1). Because K<sub>ATP</sub> channels require both Kir6.2 and SUR1 subunits (26), only those neurons that endogenously express SUR will form functional transgenic K<sub>ATP</sub> channels (14). As expected, Kir6.2 messenger RNA (mRNA) levels were increased in nK<sub>ATP</sub>-GOF mice, while SUR1 levels were unaffected (SI Appendix, Fig. S1 H, Left), suggesting no changes in channel density in mutant mice. Enhanced green

**Significance**

Gain-of-function (GOF) mutations in the ATP-sensitive potassium (K<sub>ATP</sub>) channel cause neonatal diabetes, with some individuals exhibiting developmental delay, epilepsy, and neonatal diabetes (DEND) syndrome. In this study, we uncover the direct effects of neuronal expression of K<sub>ATP</sub>-GOF mutations, and not diabetes per se, on the neurological features of DEND. Our results show a close link between neuronal K<sub>ATP</sub>-GOF expression and cognitive dysfunction in DEND and reveal that antidiabetic sulfonylureas, which successfully treat diabetes, mitigate some sensorimotor problems but not cognitive deficits. These results have critical implications for humans, revealing the need for novel drugs to treat learning and memory deficits not only for K<sub>ATP</sub>-induced DEND but also for other pathologies arising from altered ion channels in the brain.

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The authors declare no competing interest.

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fluorescent protein (eGFP), coexpressed with the mutant channel, was only detected in nKATP-GOF mice (SI Appendix, Fig. S1 H, Right), confirming transgene expression. We verified on-target cerebellar and hippocampal recombination in nKATP-GOF mice in frozen brain slices costained with anti-GFP and anti-Cre antibodies, and we compared them to absent staining in controls (SI Appendix, Fig. S2 A–C). nKATP-GOF mice showed a significant increase in ambulatory activity \( [F(1, 24) = 24.74, P = 0.00005] \), vertical rearing frequency \( [F(1, 24) = 6.354, P = 0.0188] \), distance traveled in the periphery \( [F(1, 24) = 20.71, P = 0.0001] \) and center entries \( [F(1, 24) = 13.70, P = 0.0011] \), no differences in time spent in the center, but reduced overall resting time \( [\text{Genotype: } F(1, 24) = 17.34, P = 0.0003] \) (Fig. 1 A–F). nKATP-GOF showed a significant increase in pole climb-down time \( [F(1, 24) = 10.08, P = 0.0041] \) and decreased time on the elevated ledge \( [F(1, 24) = 5.653, P = 0.0257] \) (Fig. 1 G and H). nKATP-GOF mice took significantly less time to climb to the top of a 90° screen \( [F(1, 24) = 48.91, P = 0.00005] \), distance traveled in the periphery \( [F(1, 24) = 11.61, P = 0.003] \), and swimming speeds \( [F(1, 19) = 9.032, P = 0.0073] \) (Fig. 4 A–C). They also showed significant deficits in escape latency \( [F(1, 19) = 9.280, P = 0.0066] \) and swimming speeds \( [F(1, 19) = 10.18, P = 0.0048] \) but not in path length \( [F(1, 19) = 3.305, P = 0.0849] \) in MWM place trials (Fig. 4 D–F). Robust differences in probe trial retention variables such as platform crossings \( [F(1, 19) = 33.45, P = 0.00005] \), time in the target quadrant \( [F(1, 19) = 4.700, P = 0.0431] \), and spatial bias were demonstrated in hKATP-GOF mice (Fig. 4 G and H). Similar baseline freezing levels were found in hKATP-GOF and control mice on day 1 of fear conditioning (Fig. 4); however, hKATP-GOF mice showed significantly elevated freezing \( [F(1, 19) = 9.545, P = 0.006] \) during tone–shock training. An analysis of the contextual fear data only revealed a genotype-by-time interaction \( [F(7, 133) = 2.519, P = 0.0360] \). Pairwise comparisons indicated similar degrees of freezing up to the middle of the test session followed by a greater habituation response in hKATP-GOF mice (Fig. 4). No significant differences between groups were observed during the altered-context baseline, auditory cue test, or on shock sensitivity (Fig. 4K and SI Appendix, Table S4).

Impaired Hippocampal LTP Is Associated with Learning and Memory Deficits in KATP-GOF Induced DEND. To assess the effect of KATP-GOF mutations on neuronal excitability and synaptic function, we assayed long-term potential (LTP) induction in acute hippocampal slices. LTP induction was significantly reduced in nKATP-GOF mice relative to controls (Fig. 5A) with a significant difference in the simple effect of genotype \( [F(1, 15) = 11.09, P = 0.0046] \) but no genotype-by-time interaction \( [F(21, 31) = 1.185, P = 0.2620] \). The slope of the baseline field excitatory postsynaptic potentials (fEPSP) did not differ between nKATP-GOF and littermate control mice during the pretetanic baseline (SI Appendix, Fig. S4A). Similar to nKATP-GOF mice, hKATP-GOF mice showed a significant reduction in LTP, with differences in the simple effect of genotype \( [F(1, 18) = 6.518, P = 0.02] \) (Fig. 5B) and genotype-by-time interaction \( [F(21, 38) = 1.751, P = 0.2221] \). The baseline fEPSP slope did not show significant differences between hKATP-GOF and littermate control mice (SI Appendix, Fig. S4B).
Fig. 1. Locomotor and sensorimotor functions are significantly impaired in nKATP-GOF mice. Locomotor activity: (A) total ambulations, (B) vertical rears, (C) distance traveled in the periphery, (D) center entries, (E) time in the center, and (F) resting time. Sensorimotor battery tests: (G) time to climb down pole, (H) time on an elevated ledge, and (I) time to climb to the top of 90° inclined screen. (J) Stationary rod, (K) constant speed, and (L) accelerating rotarod. For rotarod, T1 and T2 refer to Trials, S1 to 3 refers to Sessions. All tests shown in this figure were acquired from the same cohort. nKATP-GOF mice (red, n = 14) and littermate control mice (gray, n = 13). *P < 0.05, **P < 0.01, ***P < 0.001, ****P < 0.0001.
Fig. 2. MWM and fear conditioning assays reveal severe deficits in spatial learning, memory, and fear conditioning in nKATP-GOF mice in MWM-cued trials: (A) escape path length, (B) escape path latency, and (C) swimming speeds. MWM place trials: (D) escape path length, (E) escape path latency, and (F) swimming speeds. MWM probe trial: (G) platform crossings and (H) target quadrant time and spatial bias for the target quadrant. Fear conditioning: (I) tone–shock baseline-training, (J) contextual fear conditioning, and (K) auditory cue tests. All tests shown in this figure were acquired from the same cohort. nKATP-GOF mice (red, n = 14) and littermate controls (gray, n = 13). *P < 0.05, **P < 0.01, ***P < 0.001, ****P < 0.0001.
Fig. 3. Locomotor, not sensorimotor, functions are impaired in hK<sub>ATP</sub>-GOF mice. Locomotor activity: (A) total ambulations, (B) vertical rearing, (C) distance in the periphery, (D) center entries, (E) time in the center, and (F) time resting. Sensorimotor battery: (G) time to climb down pole, (H) time on elevated ledge, and (I) time to climb to the top of 9° inclined screen. (J) Stationary rod, (K) constant speed, and (L) accelerating rotarod. All tests shown in this figure were acquired from the same cohort. hK<sub>ATP</sub>-GOF mice (blue, n = 12) and controls (gray, n = 12). *P < 0.05.
Fig. 4. MWM and fear conditioning test results reveal selective spatial learning, memory, and fear conditioning deficits in hK<sub>ATP</sub>-GOF mice in MWM-cued trials: (A) escape path length, (B) latency, and (C) swimming speeds. MWM place trial: (D) escape path length, (E) latency, and (F) swimming speeds. MWM probe trial: (G) platform crossings and (H) time in the target quadrant and spatial bias for the target quadrant. Fear conditioning: (I) tone–shock baseline training, (J) contextual fear conditioning, and (K) auditory cue test. All tests shown in this figure were acquired from the same cohort. hK<sub>ATP</sub>-GOF mice (blue, n = 12) and littermate controls (gray, n = 12). *P < 0.05, **P < 0.01, ***P < 0.001, ****P < 0.0001.
Hippocampal Neurons from nKATP-GOF and hKATP-GOF Mice Showed Sensitivity to K\(_{\text{ATP}}\) Channel Openers and Inhibitors. To address K\(_{\text{ATP}}\)-GOF–mediated alterations to neuronal function, whole-cell voltage ramp experiments were performed in primary cultures of dissociated hippocampal neurons. To increase the likelihood of recording K\(_{\text{ATP}}\)-specific currents and assessing K\(_{\text{ATP}}\) drug sensitivity, voltage ramps were recorded in high-potassium external solutions in the presence or absence of K\(_{\text{ATP}}\) channel openers (diazoxide) and inhibitors (tolbutamide) (Fig. 6A). The potential off-target effects of diazoxide were mitigated in these recordings by simultaneous application of the \(\alpha\)-amino-3-hydroxy-5-methyl-4-isoxazolepropionic acid receptor (AMPA receptor) inhibitor NBQX (44). A two-way ANOVA of genotype and bath condition (Na\(_1\), K\(_1\), Dz, etc.) revealed an overall reduction in current density in nKATP-GOF mice relative to controls [genotype effect: \(F(1, 24) = 7.318, P = 0.0124\); however, comparisons between similar conditions (e.g., Dz-GOF to Dz-control) revealed no significant pairwise differences (Fig. 6C). The application of 300 \(\mu\)M diazoxide increased K\(_{\text{ATP}}\) current density, whereas the addition of 500 \(\mu\)M tolbutamide decreased it (Fig. 6C). The amplitude of this effect, defined as the difference in current densities between diazoxide and tolbutamide (Dz-Tol), did not differ between nKATP-GOF and control mice (Fig. 6D), suggesting unaffected K\(_{\text{ATP}}\) drug sensitivity in mutant mice. No significant differences in current density under any conditions were found in nKATP-GOF neurons when 0.5 mM Mg-ATP was added to the internal solution (Fig. 6C). hKATP-GOF neurons demonstrated no significant differences in current densities (Fig. 6E) or K\(_{\text{ATP}}\) drug sensitivity (Fig. 6F) with respect to control neurons. As predicted, neurons from global Kir6.2-knockout (K\(_{\text{ATP}}\)-KO) mice demonstrated a marked reduction in overall currents and no sensitivity to diazoxide or tolbutamide (Fig. 6D), confirming the specific effects of these drugs on K\(_{\text{ATP}}\).

Learning and Memory Deficits Are Not Improved by Sulfonylurea Therapy. To test the effect of peripheral sulfonylureas on neurological features of DEND, nKATP-GOF mice were implanted with slow-release glibenclamide (Glib) pellets. Glib-nKATP-GOF mice showed significant increases in distance traveled in the periphery \([F(1, 18) = 4.644, P = 0.0450]\) (Fig. 7C) and reduced time to the top of the 90° screen \([F(1, 18) = 4.593, P = 0.046]\) (Fig. 7F) but no differences in total ambulations, rearing frequency, center entries (Fig. 7 A, B, and D), or pole climb-down time (Fig. 7E). Glib-nKATP-GOF mice showed no deficits in escape path length but had impaired latency \([F(1, 18) = 7.137, P = 0.0156]\) and swimming speeds \([F(1, 18) = 12.71, P = 0.0022]\) during MMWcued trials (Fig. 6 G–I) and impaired escape path length \([F(1, 18) = 20.01, P = 0.0003]\), latency \([F(1, 18) = 36.68, P < 0.0005]\), and swimming speeds \([F(1, 18) = 11.29, P = 0.0035]\) during place trials (Fig. 7 J–L). Glib-nKATP-GOF mice showed reduced platform crossings \([F(1, 18) = 21.94, P = 0.0002]\) and time in target quadrant \([F(1, 18) = 38.87, P = 0.00005]\) and no spatial bias during probe trials (Fig. 7 M and N). During fear conditioning, Glib-nKATP-GOF mice exhibited similar baseline freezing levels on day 1 but lower freezing levels during certain training trials \([F(1, 18) = 4.768, P = 0.0425]\, genotype-by-time interaction: \(F(2, 36) = 4.525, P = 0.0188\) (Fig. 7O). Contextual conditioning was still significantly impaired in Glib-nKATP-GOF mice \([F(1, 18) = 20.69, P = 0.0002]\) (Fig. 7P). No significant differences were observed during the altered-context baseline, but Glib-nKATP-GOF mice demonstrated significantly lower freezing levels on average across minutes during the auditory cue test \([F(1, 18) = 7.567, P = 0.0131]\) (Fig. 7Q and SI Appendix, Table S5).

Severely Diabetic βKATP-GOF Mice Did Not Demonstrate Cognitive Deficits. To test if diabetes contributes to the neurological features of DEND, pancreatic β-cell–specific K\(_{\text{ATP}}\)-GOF mice (SI Appendix, Fig. S1G) (9) were assayed on behavioral tests. βKATP-GOF mice displayed reduced ambulations \([F(1, 18) = 6.845, P = 0.0175]\) and distance traveled in the periphery \([F(1, 18) = 9.048, P = 0.0076]\) and increased resting time \([F(1, 20) = 2.770, P = 0.0188]\) (SI Appendix, Fig. S5 A, C, and E) with no differences in vertical rearing or center entries (SI Appendix, Fig. S5 B and D). Sensorimotor battery tests showed a significant increase in the time to climb to the top of 60° \([F(1, 18) = 8.274, P = 0.0100]\) and 90° screens \([F(1, 18) = 6.570, P = 0.0196]\) (SI Appendix, Fig. S6 B, C, and D).


**Discussion**

K<sub>ATP</sub>-GOF–activating mutations are the most common cause of neonatal diabetes and are associated with DEND syndrome.
Fig. 7. Glib therapy selectively improves locomotor and sensorimotor, not neurological, features of DEND. Locomotor activity: (A) total ambulations, (B) vertical rearings, (C) distance traveled in the periphery, and (D) center entries in Glib-nKATP-GOF (pink, n = 11) and littermate controls (white, n = 11). Sensorimotor battery: (E) pole climb-down time and (F) time to the top of a 90° inclined screen. MWM-cued trials: (G) escape path length, (H) escape path latency, and (I) swimming speeds. MWM place trials: (J) escape path length, (K) escape path latency, and (L) swimming speeds. MWM probe trial: (M) platform crossings and (N) time in the target quadrant and spatial bias. Glib-nKATP control mice demonstrated spatial bias for the target quadrant by spending significantly more time in it versus the other pool quadrants (all Ps < 0.0001), but Glib-nKATP-GOF did not demonstrate a similar bias. Fear conditioning: (O) tone–shock baseline training, (P) contextual fear, and (Q) auditory cue tests. *P < 0.05, **P < 0.01, ***P < 0.001, ****P < 0.0001. All tests shown in this figure were acquired from the same cohort.
Hyperactivity in nKATP-GOF mice is consistent with findings in mouse models of attention-deficit hyperactivity disorder (ADHD) (27) and correlates with ADHD diagnoses in individuals with KATP-GOF-induced DEND (4, 28). They also mirrored the opposite effects: hypoactivity and impaired motor function observed in KATP-KO mice (15). Sensorimotor deficits in nKATP-GOF mice are consistent with clinical reports showing neuromotor problems and developmental delay in motor milestones in individuals diagnosed with DEND (7, 23), deficits that had been ascribed to KATP-GOF expression in cerebellar Purkinje neurons in mice (10). The impairment in escape path length and latency in nKATP-GOF mice in MWM-ued trials suggests that nonassociative factors (visual/sensorimotor disturbances) affected their performance to locate and swim to the platform, which is supported by slower swimming speeds and poor sensorimotor performance, factors which may also have influenced their reduced performance on place and probe trials. Impairments in contextual fear and auditory cue conditioning in nKATP-GOF mice could arise from faster habituation relative to controls or from an inability to sustain freezing because of hyperactivity. These results are consistent with impairments to acquisition and early consolidation of contextual memory in C57BL/6J mice intrahippocampally injected with D2 (17). Strikingly, hKATP-GOF mice, with the expression of KATP-GOF channels in pancreatic β-cells, demonstrated severe diabetes but not broad neurological features, further suggesting that the neurological features of DEND arise from expression of KATP-GOF channels in the brain.

By contrast, hKATP-GOF mice did not show significant impairments on sensorimotor or rotorod tests, although they demonstrated increased locomotor activity. Slower swimming speeds in hKATP-GOF mice suggest that they may not be completely free of compromised motor function; however, the impaired retention performance in the probe trial was as strong as in nKATP-GOF mice. Indeed, since no significant spatial learning deficits were observed in the hKATP-GOF mice, we cannot conclude that poor performance on the probe trial is due to slower swimming speeds but may reflect a selective retention deficit. A specific contextual conditioning deficit in the absence of an auditory cue impairment in hKATP-GOF reinforces the hippocampal specificity and symptomatology of this model and suggests that additional findings in auditory fear conditioning in nKATP-GOF mice have an extrahippocampal origin. These results are in agreement with intellectual learning and memory deficits observed in human DEND (4–8, 23). Our studies not only parallel the cognitive deficits reported in humans carrying KATP-GOF mutations (29–31) but also underscore the role of hippocampal KATP channels in learning and memory processes.

We have provided substantial evidence here that the neuronal expression of KATP-GOF subunits leads to motor dysfunction and spatial learning and memory deficits in KATP-GOF mice. Surprisingly, somewhat similar effects were previously reported in global KATP-KO mice (17). However, the application of Glib directly into central neurons by intraseptal injection improved spontaneous alternation and memory deficits induced by galanin or morphine in rats (18, 19), which suggests that the loss of KATP in nonhippocampal neurons may contribute to the learning and memory deficits in the global KATP-KO animal. In our study, impaired LTP induction in hippocampal slices from KATP-GOF mice was associated with cognitive deficits. KATP channels are present pre- and postsynaptically in CA1 hippocampal neurons (20, 32). Thus, KATP-GOF mutations may provide a barrier to LTP by persistently reducing presynaptic membrane potential and decreasing Ca2+ influx while postsynaptically reducing N-methyl-D-aspartate receptor (NMDAR) function by retaining Mg2+ channel block. The KATP-GOF mutation used here is expected to increase channel open probability and reduce ATP sensitivity but maintain maximum whole-cell KATP conductances.

While dissociated hKATP-GOF neurons demonstrated no overall changes in whole-cell KATP currents, nKATP-GOF neurons showed an overall small reduction relative to control littersmates. Given the very small conductances and variability in the sampled measurements, this could potentially reflect compensatory changes in channel availability, but it is consistent with the lack of any gross increase in KATP channel density driving the phenotype.

In dissociated hippocampal neurons from both nKATP-GOF and hKATP-GOF mice, sensitivity to Tol is still observed, indicating inhibition of KATP by direct pharmacological action. Homomeric expression of the same Kir6.2 N-terminal truncation in recombinant cell lines resulted in reduced sensitivity to Tol (33); however, in the present study, Kir6.2[K185Q,AN30] mutant subunits are expressed with endogenous wild-type Kir6.2 subunits, approximating the human heterozygous condition and forming functional channels with varying Tol sensitivity as a result of variable wild-type and mutant subunit ratios. This may help to explain why neurons in a dissociated culture did not demonstrate any obvious difference in Tol sensitivity. Intraseptal injections of Glib attenuate galanin- or morphine-induced behavioral impairments in rats, suggesting a direct effect of this drug on KATP channels to modulate neural function and behavior (18, 19). Given this, the failure of Glib-treated nKATP-GOF mice to demonstrate broad improvements in sensorimotor and cognitive features is surprising but consistent with previous findings showing that systemic administration of Glib fails to achieve therapeutic levels in the brain and cerebrospinal fluid of normal rodents, even with doses 10 times higher than those used in our study (34). Improvements in some sensorimotor features in Glib-treated nKATP-GOF mice may thus reflect sulfonylurea action on KATP channels in the peripheral nervous system. These results are consistent with findings in humans treated with sulfonylureas in which the restoration of muscular tone and other sensorimotor functions is far more common than improvements in cognitive deficits (4, 7, 8, 23–25, 30, 35, 36).

These findings indicate that KATP-GOF channels in hippocampal neurons play an important role in learning and memory deficits and that the cognitive deficits in DEND result from neuronal KATP-GOF expression rather than from diabetes per se. The inefficacy of sulfonylureas in improving cognitive deficits in mice is in agreement with the limited neurological improvements observed in human DEND treatment. These studies have important clinical implications, pointing to potential mechanisms underlying cognitive deficits of KATP-induced DEND and indicating the need for novel drugs to treat neurological features arising from KATP-GOF mutations while also providing a platform to study other brain abnormalities induced by ion channel dysfunction.

Methods

Animals. Cre-inducible Kir6.2[K185Q,AN30]GOF transgenic mice were previously generated (9) (SI Appendix).Kir6.2[K185Q,AN30] mice were crossed to Syn-Cre mice (37), dopamine receptor D3-Cre (Drd3-Cre) mice (rederived “in house” (38)), or tamoxifen-inducible Pdx-CreERTx mice to generate pan-neuronal nKATP-GOF (SI Appendix, Fig. S1B), hippocampus-specific neuronal KATP-GOF (SI Appendix, Fig. S1C), or pancreatic β-cell-specific jKATP-GOF mice, respectively (SI Appendix, Fig. S1D), induced by tamoxifen at 6 wk old (9). Age-matched single transgenic Syn-Cre, Drd3-Cre, Pdx1-Cre, or Kir6.2[K185Q,AN30] and wild-type littermates were used as controls since no significant differences among genotypes were found (9). Because the mice were generated under different promoter-Cre backgrounds across studies, we have compared mutant KATP-GOF mice to their respective control littersates within a cohort. KATP-KO mice were generated by the targeted disruption of the gene encoding for the Kir6.2 subunit of the KATP channel (39). Mice were maintained on a normal chow diet and housed in a 12-h light–dark cycle. All experiments were performed in compliance with relevant laws and institutional guidelines and were approved by the Washington University Animal Studies Committee’s Institutional Animal Care and Use Committee (IACUC) Protocol Approval Animal Welfare Assurance No. A-3381-01.
One-Hour Locomotor Activity and Sensorimotor Battery. Locomotor activity was evaluated as previously described (40). General activity variables (total ambulations, the number of vertical rearings) were collected along with emotionality indices (time spent, distance traveled, and entries made in central zone and distance traveled in peripheral contiguous area). The following day, mice were run on a battery of sensorimotor tests (ledge and platform; inverted screen; pole, inclined screens, and rotorod) to assay balance, strength, coordination, and movement initiation (for details, see SI Appendix). The averaged "after-trial" chamber were used in all analyses. The rotorod-evaluated mice during three test sessions were separated by 4 to minimize motor learning. The rod was stationary for trial 1 and continuously rotated at constant speed (2.5 rpm) for trials 2 and 3. For trials 4 and 5, the rod rotated at an accelerating speed (0.13 rpm/s). The time an animal remained on the rod was used as the dependent variable.

MWM Navigation. Spatial learning and memory were assessed using the MWM 3 d after completion of the sensorimotor battery. A computerized tracking system (ANY-maze; Stoelting) (40, 41) recorded the swimming pathway of the mouse to the escape platform and quantified path length, latency to the escape platform, and swimming speeds during cued, place, and probe trials conducted in a pool of opaque water. Cued trials were conducted for four time/day for 2 consecutive days for a total of eight trials with an intertrial interval (ITI) of 30 min and a 60 s maximum trial, and performance was analyzed as four blocks of two trials. After 3 d, spatial learning was assessed on place condition, two blocks of two consecutive trials over 5 d with 60 s maximum trial and an ITI of 30 s spent on platform. Blocks were separated by ~2 h and analyzed over five blocks of four trials, each block representing 1 d of training. A 60-s probe trial was administered ~1 h after the last trial on day 5 when the platform was removed, and the mouse was released into the maze from the quadrant opposite where the platform was located. The amount of time the mouse spent in each quadrant of the pool and the number of times it crossed the exact location where the platform had been recorded.

Conditioned Fear. Fear conditioning was evaluated as previously described (41) after examining spatial learning and memory. Mice were placed into one of two Plexiglas conditioning chambers (26 cm x 18 cm x 18 cm; Med-Associates), each of which contained distinct visual, tactile, and olfactory cues. Freezing behavior was assessed for a 2-min baseline period prior to tone–shock conditioning. After spending 3 min in the conditioning chamber, and every 60 s thereafter, the mice were exposed to three tone–shock pairings. Each pairing consisted of 20 s broadband white noise presented at 80 dB (conditioned stimulus; CS), with 1.0 mA continuous foot shock (unconditioned stimulus) presented during the last second of the tone. Mice were placed back into the same conditioning chamber the following day, and freezing behavior was measured over an 8-min period. After 1 d, mice were placed into a different "context" chamber to assay context-independent fear conditioning. Freezing behavior was recorded for a 2-min baseline period, after which mice were assessed on the auditory cue test, which involved the presentation of the tone (CS) over an 8-min period. Freezing behavior was quantified using Freezeframe image analysis software (Actimetrics), during which freezing was defined as no movement beyond that associated with breathing. The data are presented as a percentage of time spent freezing relative to the total duration of the trial. Shock sensitivity was evaluated after fear conditioning as previously described (42).

Chronic Sulfonylurea Treatment. Glib (Glyburide) pellets at the concentration of 2.5 mg per 60-d release were obtained from Innovative Research of America. Mice that were 5-wk-old were anesthetized with Avertin (0.25 mg/kg body weight), the skin on the lateral side of the neck of the animal was lifted, and pellets were implanted under the skin of the neck using a stainless-steel precision trochar (43). A total of 3 wk after pellet implantation, all mice were tested in the same behavioral suites as above (One-Hour Locomotor Activity and Sensorimotor Battery, MWM Navigation, Conditioned Fear).

LTP Hippocampal slice preparation. Mice were collected at P30-60, anesthetized, and then their brains were harvested and transferred immediately to ice-cold sucrose-based artificial cerebrospinal fluid with 95% carbogen and prepared by vibratome (Leica VT1200) into 300-μm coronal hippocampal slices. The preparation for all solutions and specific procedures are described in SI Appendix.

Electrophysiology recording. All hippocampal slice experiments were performed in extracellular current-clamp mode at 30 °C and recorded using pClamp 9.2 software (Axon Instruments/Molecular Devices), digitized at 20 kHz, and filtered at 2 kHz. The fEPSPs were recorded in response to current injections (Grass Instrument-S88-stimulator) in the stimulating electrode every 20 s. After establishing maximum fEPSP amplitude with an input–output curve, current injection levels were reduced to 40% of maximum and at least 25 min of baseline recorded (see SI Appendix for details). After baseline, a 100-Hz tetanus stimulus was applied to the slice to induce LTP, and recording continued for at least 1 h to assess LTP induction. The slope of the rising phase (0 to 50% maximum amplitude in trace) of the fEPSP was recorded as the main determinant of changes in synaptic plasticity (see SI Appendix for more details).

Whole-Cell Patch Clamp Electrophysiology in Dissociated Hippocampal Neurons. Hippocampal tissue from P0-P3 experimental/control mice was dissociated and plated as a single-cell suspension onto collagen-coated tissue-culture plates. Cells were first bathed in high-sodium (identical to low-potassium solution) solution prior to local perfusion with experimental solutions (see SI Appendix for more details). Dz (300 μM) or Tol (500 μM) was prepared in high-potassium solution. Membrane potential was initially clamped at −70 mV and then pulsed to −120 mV for 10 ms to allow for settling. Voltage was then ramped up at a rate of 40 mV/s to a maximum of 40 mV before being stepped down again to −70 mV with voltage ramps repeated three times per condition. Currents at −120 mV (after 10-ms recovery) were measured to maximize potassium channel–mediated conductances. All recordings were performed at room temperature.

Statistical Analyses. Mouse behavior. Data are presented as mean ± SEM. Statistical significance was determined using unpaired Student’s t tests (asterisks). Two-way repeated-measures ANOVA (asterisks within parentheses (*)) with one between-subjects (genotype) and one within-subjects (blocks of trials) variables were used to analyze locomotor activity, rotorod, MWM, conditioned fear, LTP, and GdT data. To protect against violations of sphericity, we used the Geisser–Greenhouse method to adjust α-levels for all within-subject effects containing more than two levels. Simple effects tests were conducted following certain significant interactions and subsequent pairwise comparisons and Bonferroni correction to control for alpha inflation because of multiple comparisons (GraphPad Prism 8.4.0).

LTP electrophysiology. The slope of the rising phase of the fEPSP was calculated as the average of the maximum slope within the first 0 to 50% of the rising phase and the five preceding and five subsequent digitized samples. Slopes from successive recordings within a 3-min window were also averaged. Slopes were expressed as a percentage of baseline (calculated as the average maximum slope of the 20-min preceding LTP induction). Data were analyzed using custom MATLAB scripts (MATLAB and Statistics-Toolbox-Release-2018b, TheMathWorks) and Microsoft Excel. Significance levels are denoted in each figure, and nonsignificant differences are not indicated.

Dissociated culture electrophysiology. Current amplitudes at −120 mV (after 10 ms settling) were measured and normalized to cell capacitance. These current densities were then analyzed with two-way ANOVAs with the genotype (two levels) and the bath condition (six levels) as the between- and within-subjects factors, respectively. The Geisser–Greenhouse method was used to correct for violations of sphericity. Post-hoc pairwise Student’s t tests comparing bath conditions (Na1 to Na5, Dz to Dz, etc.) across genotypes were performed using the Bonferroni correction to adjust for multiple comparisons. Within cells, KATP drug sensitivity was calculated as the difference in current density between Dz and Tol conditions. Differences in KATP drug sensitivities across genotypes with or without internal ATP were analyzed using unpaired two-tailed Student’s t tests.

Data Availability. All study data are included in the article and/or SI Appendix.

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