An Epilepsy-Associated KCNT1 Mutation Enhances Excitability of Human iPSC-Derived Neurons by Increasing Slack K_{Na} Currents

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Mutations in the KCNT1 (Slack, K_{Na} 1.1) sodium-activated potassium channel produce severe epileptic encephalopathies. Expression in heterologous systems has shown that the disease-causing mutations give rise to channels that have increased current amplitude. It is not known, however, whether such gain of function occurs in human neurons, nor whether such increased K_{Na} current is expected to suppress or increase the excitability of cortical neurons. Using genetically engineered human induced pluripotent stem cell (iPSC)-derived neurons, we have now found that sodium-dependent potassium currents are increased several-fold in neurons bearing a homozygous P924L mutation. In current-clamp recordings, the increased K_{Na} current in neurons with the P924L mutation acts to shorten the duration of action potentials and to increase the amplitude of the afterhyperpolarization that follows each action potential. Strikingly, the number of action potentials that were evoked by depolarizing currents as well as maximal firing rates were increased in neurons expressing the mutant channel. In networks of spontaneously active neurons, the mean firing rate, the occurrence of rapid bursts of action potentials, and the intensity of firing during the burst were all increased in neurons with the P924L Slack mutation. The feasibility of an increased K_{Na} current to increase firing rates independent of any compensatory changes was validated by numerical simulations. Our findings indicate that gain-of-function in Slack K_{Na} channels causes hyperexcitability in both isolated neurons and in neural networks and occurs by a cell-autonomous mechanism that does not require network interactions.

Key words: action potential; EIMFS; epileptic encephalopathy; MMPSI; potassium channels; seizures

Significance Statement

KCNT1 mutations lead to severe epileptic encephalopathies for which there are no effective treatments. This study is the first demonstration that a KCNT1 mutation increases the Slack current in neurons. It also provides the first explanation for how this increased potassium current induces hyperexcitability, which could be the underlining factor causing seizures.

Introduction

Mutations of the Slack K^+ (K_{Na} 1.1) channel lead to a highly intractable childhood epilepsy syndrome called malignant mi-

grating partial seizures of infancy (MMPSI), also called epilepsy of infancy with migrating focal seizures (EIMFS) (Barcia et al., 2012; Kim et al., 2014; Shimada et al., 2014; Vanderver et al., 2014; Møller et al., 2015; Mikati et al., 2015; Rizzo et al., 2016; Tang et al., 2016). This is a devastating epileptic encephalopathy with no effective treatments. Children with MMPSI experience scores of multifocal seizures per day (Coppola et al., 1995). They are profoundly intellectually disabled, and brain development halts at an infantile stage with many children remaining unable to speak or walk. Mortality is high, and sudden unexpected death in

Received June 28, 2018; revised July 8, 2019; accepted July 17, 2019.

Author contributions: I.H.Q., K.P.M., F.H.G., and L.K.K. designed research; I.H.Q., S.S., K.P.M., Y.Z., S.R.A., M.R.M., M.C.M., and L.K.K. performed research; I.H.Q., S.S., K.P.M., Y.Z., S.R.A., M.R.M., M.C.M., and L.K.K. analyzed data; I.H.Q. wrote the first draft of the paper; I.H.Q., S.S., K.P.M., Y.Z., S.R.A., M.J.M., and L.K.K. edited the paper; K.P.M., M.J.M., and E.M.J. contributed unpublished reagents/analytic tools.

Acknowledgements: This work was supported by National Institutes of Health NS102239 to L.K.K. and a Swebilius Foundation Grant to I.H.Q.

K.P.M., M.J.M., and E.M.J. are employed by FUJIFILM Cellular Dynamics, Inc. The remaining authors declare no competing financial interests.

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<https://doi.org/10.1523/JNEUROSCI.1628-18.2019>
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epilepsy (SUDEP) has been reported (Møller et al., 2015). Slack mutations have now also been associated with a spectrum of focal epilepsies including other infantile epileptic encephalopathies and later onset autosomal dominant nocturnal frontal lobe epilepsy (ADNFLE) (Heron et al., 2012; Martin et al., 2014; Møller et al., 2015; Ohba et al., 2015).

Slack, most recently termed $K_{Na}1.1$ (Kaczmarek et al., 2017), is a sodium-activated potassium channel encoded by *KCNT1* (Yuan et al., 2003; Kaczmarek, 2013). Functions attributed to the channel include regulation of neuronal bursting in sensory neurons (Tamsett et al., 2009; Nuwer et al., 2010; Biton et al., 2012; Lu et al., 2015) and maintaining temporal accuracy in auditory brainstem neurons (Yang et al., 2007). Slack has also been proposed to regulate neuronal development through interactions with the Fragile X Mental Retardation Protein (FMRP) and translational machinery (Brown et al., 2010; Zhang et al., 2012; Kim and Kaczmarek, 2014; Bausch et al., 2015).

In heterologous expression systems, epilepsy-causing mutations cause substantially increased Slack channel activity over that of WT channels with no change in the level of channel protein (Barcia et al., 2012; Kim et al., 2014). This gain-of-function is seen with a large variety of different mutations (Kim et al., 2014; Martin et al., 2014; Milligan et al., 2014). Although altered voltage and/or sodium dependence contribute in part to the increased current for a subset of mutations (Kim et al., 2014; Tang et al., 2016), for most mutations the level of increased current is too large to be explained by changes in these parameters (Kim et al., 2014). Evidence indicates that enhanced positive cooperativity between channels is a major factor that increases channel activity (Kim et al., 2014). However, it is not yet known whether similar gain-of-function occurs in neurons, or if high Slack currents directly enhance neuronal excitability. Slack-mediated K_{Na} currents activate at relatively negative potentials (Santi et al., 2006; Yang et al., 2007), and increases in current near the resting potential would be expected to suppress rather than increase excitability.

Human induced pluripotent stem cell (hiPSC) technology offers a novel tool to advance our understanding of neurological diseases. hiPSCs surpass heterologous systems because regular, adjunct biology is conserved (Liu et al., 2013; LaMarca et al., 2018). In a wide variety of neurological disorders, hiPSCs have not only advanced understanding of disease mechanisms, but have also provided a more relevant avenue for drug screening (Tidball and Parent, 2016; Csobonyeiova et al., 2017; Gonzalez et al., 2017; Balan et al., 2018).

To understand how Slack mutations contribute to hyperexcitability, we have generated and evaluated a hiPSC-derived neuron line harboring an MMPSI-associated mutation known to have gain of function, P924L Slack (Milligan et al., 2014). The mutation was generated by nuclease-mediated engineering, allowing a true isogenic control. We characterized neurons derived from these cells using voltage-clamp recordings as well as multielectrode arrays (MEA) and current-clamp recordings to evaluate firing patterns and the properties of action potentials. Current-clamp recordings were performed at an earlier [days in vitro (DIV) after plating 7–8] and later (DIV after plating 14–16) maturation stage.

Materials and Methods

Generation of Slack P924L iPSC-derived neurons. Human neurons differentiated from induced pluripotent stem cells (iCell Neurons) were provided by FUJIFILM Cellular Dynamics. These cells were generated through a proprietary forebrain differentiation protocol developed by

Cellular Dynamics, resulting in a population of neurons without glial or oligodendrocyte cells. The cells have been characterized to demonstrate their similarity to primary neurons. Thawed neuron populations are >95% pure neurons, rapidly form neural networks via functional synapses, and display long-term viability and reproducibility (Berry et al., 2015). They have previously been described and characterized including methods for plating and maintenance (Chai et al., 2012; Haythornthwaite et al., 2012; Chatzidaki et al., 2015). Gene expression and pharmacology closely resembles neonatal prefrontal cortex (Dage et al., 2014). To generate a *KCNT1* P924L allelic variant, nuclease-mediated SNP alteration of the healthy control cell line was performed to introduce a site-specific mutation using proprietary licensed protocols, similar to prior efforts that generated disease-specific models of cardiomyocytes (Drawnel et al., 2014). The resulting cells have the P924L variant in both *KCNT1* alleles.

Measurement of Slack expression. Cellular lysates were separated on 4–15% gradient SDS gels, transferred to nitrocellulose paper, probed overnight at 4°C with primary antibody as indicated. For Slack expression we used a custom polyclonal pan-Slack antibody (1:500) or GAPDH antibody (1:500, Santa Cruz Biotechnology) with goat anti-rabbit (#7074, Cell Signaling Technology) HRP-conjugated secondary antibodies (1:1000 in 5% milk solution). The Slack antibody was designed as a custom polyclonal antibody against the C terminus region of rSlack using the peptide sequence GCDVMNRVNLGYLQDEMNH (Brown et al., 2008). Generation of this rabbit polyclonal anti-pan-Slack IgG was performed by Biosynthesis. A stable rat Slack-expressing HEK cell line was used as a positive control for the antibody. Other membrane proteins were measured including Nav1.1 (1:1000, Alomone Laboratories), Kv1.1 (1:500, NeuroMab), BK (1:500, BD Bioscience), and Na,K-ATPase antibody (1:1000, Cell Signaling Technology, for normalization of membrane proteins).

Patch-clamp electrophysiology. Cells were plated on 12 mm glass coverslips that were transferred into a recording chamber. Whole-cell patch-clamp recordings were performed in one preparation at 7–8 d after plating for current clamp and another preparation at 7–10 d after recording for voltage clamp and then repeated with a second preparation for current clamp at 14–16 d. Recordings were made from five coverslips for voltage-clamp and two coverslips for each of two preparations for current clamp. Pipette tip resistance was typically 2–4 MΩ for voltage-clamp and 10–15 MΩ for current clamp. Signals were amplified with an EPC-7 (HEKA) or a Multiclamp 700B (Molecular Devices) amplifier and recorded with Clampex 10.2 software (Molecular Devices). Data were acquired at a sampling rate of 2 kHz (for voltage clamp) or 20 kHz (for current clamp) and analyzed using Clampfit 10 (Molecular Devices) and MATLAB 2017b (The MathWorks). All measurements were conducted at room temperature.

For voltage-clamp recordings, the extracellular medium contained the following (in mM): either 139 NaCl or 139 *N*-methyl-D-glucamine (NMDG), 5.4 KCl, 1 CaCl₂, 1 MgCl₂, and 10 glucose, with pH 7.3, and patch electrodes were filled with (in mM) 97.5 K⁺ gluconate, 32.5 KCl, 10 HEPES, 1 EGTA, 2 MgCl₂. For current clamp, the extracellular recording medium contained the following (in mM): 139 NaCl, 10 HEPES, 4 KCl, 2 CaCl₂, 1 MgCl₂, 10 D-glucose (310 mOsm, pH 7.4), and patch electrodes were filled with internal solutions containing the following (in mM): 130 K⁺ gluconate, 6 KCl, 4 NaCl, 10 Na-HEPES, 0.2 K-EGTA, 0.3 GTP, 2 MgATP, 0.2 cAMP, 10 D-glucose, 0.15% biocytin and 0.06% rhodamine. The pH and osmolarity of the internal solutions were brought close to physiological conditions (pH 7.3, 290–300 mOsm).

For measurements of K_{Na} currents, whole-cell voltage-clamp measurements were performed with cells held at −70 mV and given 600 ms voltage pulses in 10 mV steps over a fixed range of −80 to +70 mV. Recordings were obtained independently for cells in extracellular medium containing either 140 mM Na⁺ or 140 mM NMDG. Current traces from NMDG were averaged and subtracted from the individual current traces obtained from the Na⁺ condition. The difference current at the end of the voltage pulse was considered to be the steady-state K_{Na} current.

To quantify the total number of evoked action potentials, cells were held in current-clamp mode near −60 mV with a steady holding current,

and current injections were given starting 12 pA below the steady holding current, in a total of 20 depolarization steps, with 3 pA increments and 400 ms in duration. Neurons that needed >50 pA to be held at −60 mV were discarded from the analysis.

Numerical simulations. Simulations of the effects of K_{Na} currents on neuronal firing were performed using a simple one-compartment neuronal model comprising a Na^+ current I_{Na} , a leak current I_L , a voltage-dependent K^+ current I_{Kv} , and a Na^+ -activated K^+ current I_{KNa} , with kinetic parameters based on previous simulations of the effects of K_{Na} currents on firing patterns (Yang et al., 2007; Brown et al., 2008). Responses were simulated by integration of the equation $C dV/dt = I_{ext(t)} - I_{Na} - I_{Kv} - I_{KNa} - I_L$, where stimuli $I_{ext(t)}$ were presented as a single step current (0.01–0.04 nA, 100 ms). Equations and parameters for I_{Na} , I_{Kv} , and I_L were similar to those used in previous simulations on neuronal firing patterns (Liu and Kaczmarek, 1998; Wang et al., 1998; Richardson and Kaczmarek, 2000; Macica et al., 2003; Bhattacharjee et al., 2005; Song et al., 2005; Yang et al., 2007; Kaczmarek, 2012). Specifically, I_L and I_{Na} were given by the equations $I_{Na} = g_{Na} m^3 h (V - 50)$ and $I_L = g_L (V + 80)$ respectively. Kinetic parameters for the evolution of the variables m and h were $g_{Na} = 1.5 \mu S$, $k_{am} = 76.4 ms^{-1}$, $\eta_{am} = 0.037 mV^{-1}$, $k_{bm} = 6.93 ms^{-1}$, $\eta_{bm} = -0.043 mV^{-1}$, and $k_{ah} = 0.000135 ms^{-1}$, $\eta_{ah} = -0.1216 mV^{-1}$, $k_{bh} = 1.237 ms^{-1}$ and $\eta_{bh} = 0.0384 mV^{-1}$. The voltage-dependent K^+ current I_{Kv} was given by the equation $I_{Kv} = g_{Kv} l r (V + 80)$, with the parameters for the evolution of the gating particles l and r given by $g_{Kv} = 0.03 \mu S$, $k_{al} = 1.2 ms^{-1}$, $\eta_{al} = 0.03512 mV^{-1}$, $k_{bl} = 0.2248 ms^{-1}$, $\eta_{bl} = -0.0319 mV^{-1}$, $k_{ar} = 0.0438 ms^{-1}$, $\eta_{ar} = -0.0053 mV^{-1}$, $k_{br} = 0.0562 ms^{-1}$ and $\eta_{br} = -0.0047 mV^{-1}$. The capacitance C was 0.1 nF and the leakage conductance g_L was 0.002 μS .

To simulate Slack currents, I_{KNa} was given by the equation $I_{KNa} = g_{KNa} n^2 s^4 (V + 80)$, where the variable n describes voltage dependence and is governed by the kinetic parameters, $k_{an} = 1.38 ms^{-1}$, $\eta_{an} = -0.0105 mV^{-1}$, $k_{bn} = 0.5763 ms^{-1}$ and $\eta_{bn} = -0.0355 mV^{-1}$ (Brown et al., 2008). The conductance g_{KNa} was varied between 0 and 4.0 μS .

The variable s represents the proportion of Slack subunits activated by Na^+ ions at the cytoplasmic face of the Slack channel. As in previous simulations, changes in Na^+ concentration at the cytoplasmic face of the K_{Na} channels, $[Na^+]_i$, were modeled by the equation $d[Na^+]_i/dt = aI_{Na} + a' - b[Na^+]_i$, where the kinetic constants a and a' represent Na^+ entry through voltage-dependent Na^+ channels and a Na^+ leak, respectively, and b determines the rate of pumping of Na^+ out of the cell. In the model cell, $a = 2.5$, $a' = 0.25 mM ms^{-1}$ and $b = 0.1 ms^{-1}$. The variable s represents the proportion of Slack channels activated by $[Na^+]_i$ and evolves according to the equation $ds/dt = k_f [Na^+]_i (1-s) - k_b s$, with $k_f = 0.014 mM^{-1} ms^{-1}$ and $k_b = 0.4 ms^{-1}$.

Multielectrode array recordings. WT and Slack P924L-expressing iPSC-derived neurons were thawed and plated in individual clusters (“dots”) of 80,000 cells in a 48-well MEA plate (Axion Biosystems). Cells were cultured in BrainPhys neuronal medium (Stem Cell Technologies) and underwent 50% media change every 2–3 d. Eight-minute long recordings were made on DIV 11 post-plating ($n = 24$ per group). Raw extracellular voltage recordings were processed on a Maestro 768 channel system (Axion) with 200–4 kHz Butterworth filter. Action potentials were captured on each channel via a 6 SD detection limit. Analysis of MEA spike trains was performed via Neural Metric (Axion). Burst analysis was based on a Poisson Surprise algorithm that identified clusters with a surprise factor threshold $S \geq 10$ (Legéndy and Salcman, 1985). Synchrony between channels was assessed using a binless measure of cross-correlation (Paiva et al., 2010).

Experimental design and statistical analysis. Action potential shape analysis: The first evoked action potential was used for spike shape analysis (with the lowest injected current needed for eliciting an action potential). Spike threshold was the membrane potential at which the slope of the depolarizing membrane potential increased drastically, resulting in an action potential (the first maximum in the second derivative in dV_m/dt vs V_m phase space). The 5 ms AHP amplitude was calculated as the difference between the threshold for spiking and the value of the membrane potential 5 ms after the potential returned to cross the threshold value at the end of the action potential. The spike amplitude was calculated as the difference between the maximum membrane potential

during a spike and the threshold. Action potential width was calculated as the time it took the membrane potential to reach half the spike amplitude in the rising part of the spike to the descending part of the spike (full width at half maximum), using the threshold time as the lower limit.

Differences between groups for the steady-state K_{Na} current as measured by voltage clamp were compared using repeated-measures ANOVA followed by two-tailed Student's t test with Holm-Sidak correction for multiple comparisons. Morphological features of action potentials from current-clamp experiments were compared by two-tailed Student's t test. Spike counts were compared using a Mann–Whitney U test. Exact sample sizes and significance values are reported for all statistical tests. Means are reported with SEM unless noted otherwise.

Results

The P924L mutation does not alter Slack, BK, or Nav1.1 channel expression in neurons

We compared Slack expression in human iPSCs that were differentiated into neurons. A mutant cell line expressing the P924L Slack mutation was engineered from WT cells using nuclease-mediated engineering to ensure an isogenic control. Both cell lines were plated using two different methods. Conventional uniform plating of 80,000 cells on 12 mm glass coverslips was used to provide isolated cells that could be used for patch-clamp electrophysiology. The cells were also plated in high density “dots” of 60,000 cells in a single large cluster. These provided a high density of synaptic connections that were used for multielectrode array (MEA) recordings. Slack protein was expressed in both WT and Slack P924L cell lines as measured by Western blotting using a previously described “pan-Slack” antibody (Brown et al., 2008). The quantity of Slack protein measured by Western blotting was similar in the two cell types by densitometry. Moreover, the method of plating did not affect the level of channel expression (Fig. 1A).

As a screen for compensatory changes in other ion channels, we also measured the expression of BK channels, calcium-dependent K^+ channels which are structurally and functionally similar to Slack, and Nav1.1 channels, because sodium entry during action potentials is suspected to be intricately linked to the activation of K_{Na} currents. There was no obvious visual difference in either of these channels by Western blotting (Fig. 1B). By densitometry no significant differences were identified in Nav1.1 ($p = 0.45$, $n = 7$), Kv1.1 ($p = 0.19$, $n = 7$), or BK ($p = 0.73$, $n = 10$) expression between P924L and WT cell lysis products at DIV12 (Student's t test).

P924L Slack-expressing neurons have increased K_{Na} currents

Whole-cell voltage-clamp measurements were first performed using physiological internal and external media with the extracellular medium containing 139 mM Na^+ ions. Cells were held at −70 mV and stepped to potentials between −80 to +70 mV for 600 ms. Under these conditions, both isogenic control and P924L Slack-expressing neurons displayed a rapid inward current (consistent with a voltage-activated sodium current) followed by a large, slowly inactivating outward current (Fig. 2A). The outward currents measured in cells expressing P924L Slack were significantly larger than those in cells with the WT channel at all test potentials above +10 mV ($n = 17$; Fig. 2D).

The experiment was then repeated using external medium in which 139 mM *N*-methyl-D-glucamine (NMDG) was substituted for Na^+ ions. In this medium, the fast inward current was abolished, and the slower outward current was decreased. In this Na^+ -free condition, there was no statistically significant difference in the amplitude of the outward current between the P924L Slack cells and WT cells (Fig. 2B,E).

Figure 1. *A*, Slack expression measured by Western blot via a custom antibody specific for Slack and targeted to multiple epitopes (Brown et al., 2008) is similar in P924L and WT cells regardless of cell plating method. A stable rat Slack expressing cell line is used as control. *B*, No significant differences were seen in normalized Nav1.1 ($p = 0.45$, $n = 7$), Kv1.1 ($p = 0.19$, $n = 7$), or BK ($p = 0.73$, $n = 10$) expression between P924L and WT preparations (Student's *t* test of densitometry values).

To isolate the Na^+ -dependent outward current, traces from cells in NMDG were averaged and subtracted from the traces obtained in the external Na^+ medium. The difference current at the end of the voltage pulses, corresponding to the steady-state K_{Na} current, was substantially increased by the mutation, by as much as 20-fold over that in WT neurons (Fig. 2C,F).

The differences between the outward currents in WT and P924L Slack cell lines were most evident at more positive voltage steps, consistent with a voltage-activated current. At the upper limit of +70 mV, for example, the outward current in the Na^+ -free, NMDG-containing medium was not significantly different between WT (0.75 ± 0.13 nA) and the mutation (0.80 ± 0.070 nA; $p = 0.99$, $n = 18$, Student's *t* test with Holm correction). In contrast the outward current in the presence of extracellular Na^+ was 0.68 ± 0.080 nA for WT cells compared with 1.07 ± 0.16 nA for the mutation ($p = 9.5 \times 10^{-5}$, $n = 17$, Student's *t* test with Holm correction). The corresponding K_{Na} current estimates were -0.065 ± 0.080 nA for WT and 0.277 ± 0.156 nA for the mutation ($p = 1.1 \times 10^{-3}$, $n = 17$, Student's *t* test with Holm correction).

Early maturation P924L Slack-expressing neurons have briefer, more numerous evoked action potentials

To compare the firing properties of individual P924L Slack and WT neurons at DIV 7–8, the resting potential of all cells was adjusted to -60 mV in the current-clamp mode, using a small steady holding current. The cells were then hyperpolarized and depolarized with 400 ms current injections applied in 3 pA increments.

We first compared the characteristics of the earliest action potential triggered by a suprathreshold depolarizing current (Fig. 3A). The width of action potentials (full width at half maximum from the identified threshold potential) was significantly shorter in the P924L neurons (8.32 ± 0.66 ms for WT, 5.87 ± 0.61 ms for P924L, $p = 0.0128$, $n = 22$, Student's *t* test; Fig. 3A). In addition, the maximal afterhyperpolarization depth after the

first threshold action potential was increased in neurons bearing the P924L mutation (5.62 ± 1.05 mV for WT, 10.40 ± 1.27 mV for P924L, $p = 0.0105$, $n = 22$, Student's *t* test). These changes are evident in representative recordings of action potentials at the onset of the depolarizing step (Fig. 3B,C, insets).

Some other characteristics of the response to depolarizing and hyperpolarizing steps did not show a significant difference between WT and P924L Slack cells. The action potential threshold, as calculated from the second derivative of a dV_m/dt vs V_m plot, was similar between the two cell types (-19.3 ± 1.6 mV in WT vs -21.7 ± 1.1 mV for P924L, $p = 0.244$, $n = 26$, Student's *t* test). The amplitudes of action potentials as measured from the holding potential were not appreciably different (72.12 ± 2.17 for WT, 75.68 ± 2.24 for mutant, $p = 0.43$, $n = 26$, Student's *t* test). We did not identify any significant difference in input resistance (WT 0.395 ± 0.100 G Ω /pF, P924L 0.230 ± 0.038 G Ω /pF, $p = 0.14$, $n = 22$, Student's *t* test), which was calculated from the change in voltage in response to the range of subthreshold hyperpolarizing current pulses in the current-clamp mode.

The number of action potentials evoked by 400 ms depolarizing currents was greatly increased in the P924L Slack-expressing neurons. In control neurons, depolarizing currents elicited by 400 ms current steps over the range of 6 to 40 pA above the action potential threshold current usually evoked only a single action potential (Fig. 3A,B). In contrast, in Slack P924L cells, the same currents typically evoked two to three action potentials (Fig. 3A,C). For example, at 30 mV above threshold, there were 1.10 ± 0.10 evoked action potentials in WT and 2.17 ± 0.42 in P924 cells (Fig. 3A). The maximum number of action potentials that could be elicited in each cell over the entire range of current step amplitudes (Fig. 3A) was a median of 1 for WT neurons (range 1–4) and 4 for P924L Slack neurons (range 1–6), corresponding to a highly significant increase ($p = 0.0005$, $n = 26$, Mann–Whitney U).

During a train of action potentials evoked by a depolarizing current, changes in action potential height and width were evi-

Figure 2. *KCNT1* P924L increases neuronal K_{Na} currents. K_{Na} currents were measured by recording outward currents in the presence or absence of extracellular sodium. **A, D**, Voltage-clamp recordings of Slack P924L cells have a larger steady-state outward current when sodium is present, consistent with an increased K_{Na} current. **B, E**, In the condition where extracellular sodium is removed, the outward current for P924L neurons is no longer different from that in WT neurons. **C, F**, The difference current, a measure of K_{Na} , is substantially larger in P924L than WT, as would be expected for increased Slack channel activity. Significance values shown are from Student's *t* test with Holm–Sidak correction for multiple comparisons. n.s., not significant.

dent throughout the train (Fig. 3C). When current steps were able to evoke more than one action potential in a WT neuron, the differences in width and subsequent afterhyperpolarization that were quantified for the first action potential in control and P924L Slack cells were also evident for these later action potentials. Figure 3D shows examples of such later action potentials that occurred at the same time following the onset of depolarization of a WT and a P924L Slack neuron.

Later maturation Slack P924L neurons continue to display increased AHP and firing rates

The initial recordings suggested an early stage of neuronal maturation based on observations of short, infrequent action potentials (Sun et al., 2016). Recordings were repeated (Fig. 4) at a later maturation stage (DIV 14–16 after plating), going as late as possible before cell attachment began to be compromised. Intracellular and extracellular solutions as well as the protocol for current clamp were unchanged. In these cells, there was still no significant difference in the voltage threshold for activation (-27.1 ± 2.3 mV for WT, -31.6 ± 1.5 mV for P924L, $p = 0.10$, $n = 30$). Narrowing of the action potential displayed enough variance that

there was no longer a significant difference in width (7.57 ± 1.1 mV for WT, 5.38 ± 0.66 mV for P924L, $p = 0.09$, $n = 30$). There was a borderline difference in the height of the first AP for the more mature cells (72.4 ± 2.0 mV for WT, 80.5 ± 3.2 mV for P924L, $p = 0.047$). This did not appear to be highly significant as evident from the degree of scatter. The most prominent difference in shape was that the AHP depth remained substantially increased with the mutation (4.7 ± 1.1 mV for WT, 11.0 ± 1.4 mV for P924L, $p = 0.001$, $n = 30$). Spike counts elicited by a voltage stimulus remained more frequent with the mutation. The maximum number of spikes elicited per pulse in an individual cell was a median of 1, mean of 1.9 in WT and median of 5, mean of 4.1 for P924L ($p = 0.009$, $n = 30$, Mann–Whitney *U* test).

Increased AHP from P924L Slack neurons persists with block of BK or Kv1 channels

The simplest interpretation of the increase in AHPs in P924L Slack neurons is that the increase in AHP is caused directly by the increase in Slack current. Because there is no specific blocker of Slack channels, this could not be tested directly. It is possible, however, that the increased AHPs could result from adaptive

Figure 3. Changes in action potential caused by Slack P924L support neuronal hyperexcitability at DIV 7–8. **A**, Current-clamp recordings of Slack P924L cells have similar action potential thresholds and heights as WT cells but shorter spike width and deeper afterhyperpolarizations. P924L cells generate more action potentials over a range of currents. The maximum number of action potentials generated by each cell is higher for P924L. **B**, **C**, Sample current-clamp recordings in response to an inward current step stimulus demonstrate that WT neurons have a single action potential followed by minimal afterhyperpolarization (**B**), whereas P924L neurons have prominent afterhyperpolarization allowing the cell to fire multiple action potentials by bringing the voltage back below threshold (**C**). Significance values shown are from Student's *t* test. Whisker plots represent SDs and error bars for stimulus-dependent spike counts (**A**, lower left) represent SEM. n.s., non significant.

changes in other K^+ currents known to contribute to the fast AHP in neurons, specifically Ca^{2+} -activated BK currents or Kv1.1/Kv1.2 voltage-dependent currents. To test this possibility, we repeated the current-clamp experiments on cells at DIV 8–9 in the presence of blockers of these channels (Fig. 5). In the presence of 2 mM paxilline, a BK antagonist (Zhou and Lingle, 2011, 2014) that has no effect on Slack currents (de Los Angeles Tejada et al., 2012), WT cells had an AHP amplitude 5.5 ± 1.6 mV ($n = 11$) vs 13.6 ± 2.0 mV ($n = 15$) for P924L, again significantly different ($p = 0.007$). α -dendrotoxin is a blocker of Kv1.1 and Kv1.2, major voltage-dependent K^+ channels in the CNS, as well as of Kv1.6 (Wang et al., 1998; Harvey and Robertson, 2004). In the presence of 100 μ M α -dendrotoxin, WT cells had an AHP amplitude of 4.0 ± 0.9 mV ($n = 11$) and mutant cells had an AHP of 9.9 ± 1.0 mV ($n = 14$), which was again significantly different from WT ($p = 0.0004$) (Fig. 5). These findings indicate that compensatory increases in BK or Kv1 channels in response to the P924L Slack mutations are unlikely to be a major factor in the increased AHP amplitude.

Numerical simulations of increased K_{Na} current predict more numerous evoked action potentials.

Because activation of K^+ channels causes hyperpolarization of the plasma membrane, an increase in K^+ currents is often associated with reduced excitability. A clear exception to this is K^+ channels that activate only at positive potentials, such as Kv3-family of channels, for which increases in current promote high-frequency firing (Kaczmarek and Zhang, 2017). Although Slack channels are not strongly voltage-dependent, their dependence

on Na^+ ions implies that maximal activation occurs during action potentials when Na^+ influx is maximal. To test the potential effects of an increase in Slack currents alone on neuronal firing, we performed numerical simulations on a simple neuron model that incorporated only a Na^+ current I_{Na} , a leak current I_L , a voltage-dependent K^+ current I_{Kv} and a Na^+ -activated K^+ current I_{KNa} . The kinetic parameters for the latter were based on previous recordings and simulations of K_{Na} currents (Wang et al., 1998; Brown et al., 2008). The Na^+ concentration at the cytoplasmic face of the K_{Na} channels was determined both by Na^+ entry through voltage-dependent Na^+ channels and by a steady-state leak of Na^+ ions into the cell, and these were balanced by a first order process of pumping intracellular Na^+ from the cell. The ratios of the conductances for I_{Na} , I_L , and I_{Kv} were adjusted to generate a single action potential in response to a step depolarization, similar to the responses for neurons in the 7–8 d after plating cortical cultures.

The effects of progressive increases in I_{KNa} are shown in Figure 6A, which shows both the firing response and the accompanying changes in intracellular Na^+ levels under the plasma membrane. With the introduction of low K_{Na} currents, model neurons began to display a small fast afterhyperpolarization response. The most evident change in firing patterns in response to small increases in I_{KNa} was a progressive increase in the amplitude of the afterhyperpolarization following each action potential. As I_{KNa} was increased, an after-depolarization emerged following the initial spike. Finally, as I_{KNa} was further increased, repetitive firing was evoked throughout the duration of the depolarizing pulse. Thus, the changes in firing patterns observed in the iPSC-derived neu-

Figure 4. Current-clamp recordings of later-stage P924L-expressing IPS-derived neurons continue to display a hyperexcitable phenotype. **A**, Changes in the shape of the first threshold action potential caused by Slack P924L at DIV 14–16. **A**, Current-clamp recordings of Slack P924L cells have similar AP thresholds but deeper after hyperpolarization. P924L cells generate more action potentials in response to a stimulus over a range of current amplitudes and have a larger number of maximum elicitable spikes. Data shown as mean $\pm$ SD except for stimulus dependent spike count, which is SEM. **B**, Typical WT cells elicit a mean of two spikes. **C**, Typical P924L cells, in this case with the same current of 15 pA above baseline as the example in **B**, demonstrate a mean of more than four spikes.

rons are compatible with those expected from elevated K_{Na} currents.

Figure 6B shows the time course of activation of I_{Na} , intracellular Na^+ , the Na^+ -activated I_{KNa} current, the voltage-dependent current I_{Kv} and the leak current I_L for a simulation with a low and high value of K_{Na} conductance. Resting membrane potential was the same in both conditions. The emergence of repetitive firing is seen to be associated with the progressive accumulation of Na^+ and K_{Na} current. Figure 6, C and D, shows changes in spike threshold, height, width, and depth of afterhyperpolarizations for the simulations of Figure 6A. As with the P924L Slack cells, the emergence of sustained firing was associated with a marked increase in spike afterhyperpolarizations.

Slack P924L increases aggregate spontaneous firing rates, burst behaviors, and synchrony

To record the simultaneous activity of multiple neurons in a dense synaptically connected network, cells were plated as high-density clusters of ~60,000 cells on MEAs. Control ($n = 24$) and P924L ($n = 24$) cells exhibited spontaneous firing and, in both types of clusters, repeated bursts of action potentials were evident (Fig. 7). Overall, Slack P924L-expressing cells had an increased mean firing rate (0.360 ± 0.037 Hz for WT, 0.554 ± 0.064 Hz for P924L, $p = 0.0116$). The characteristics of the bursts of firing were also substantially different between the two cell types. Bursts were detected using a Poisson surprise analysis (Legéndy and Salcman, 1985). The frequency of these bursts was more than doubled by the mutation (4.09 ± 0.48 bursts/min for WT, 10.44 ± 1.50 bursts/min for P924L, $p = 0.0002$). Moreover, the

spike rate within each burst (burst intensity) was increased for P924L-Slack cells (6.42 ± 0.58 Hz for WT, 8.66 ± 0.69 Hz for P924L Slack, $p = 0.0174$), but burst durations were shorter (3.26 ± 0.39 s for WT, 1.70 ± 0.20 s for P924L, $p = 0.0009$). In addition, we measured the synchrony of activity between electrodes in each well and found that synchrony index values [0–1] were increased by >60% in recordings of P924L activity relative to WT ($1.037 \times 10^{-2} \pm 0.096 \times 10^{-2}$ for WT vs $1.690 \times 10^{-2} \pm 0.178 \times 10^{-2}$ for P924L ($p = 0.0023$)).

Discussion

Human iPSC-derived neurons have a Na^+ -dependent K^+ current that is substantially enhanced by the MMPSI-associated *KCNT1* P924L mutation. Previous work demonstrated gain-of-function of the Slack current using heterologous expression of this and other epilepsy-associated mutations in *Xenopus* oocytes (Barcia et al., 2012; Kim et al., 2014; Milligan et al., 2014). Our findings provide direct evidence that neurons carrying an epilepsy associated Slack mutation also have an increased K_{Na} current. The demonstration of a hyperexcitable phenotype for this mutation underscores the impact iPSCs can provide when investigating neurological disorders within their native milieu. Action potentials in these neurons are shorter and are followed by a larger fast afterhyperpolarization. The effects on this afterhyperpolarization are preserved in the presence of BK and Kv1 channel antagonists. In response to a maintained inward current, cells with the mutation can sustain more action potentials. It is likely that the increased afterhyperpolarizations accelerate recovery from the refractory period after each action potential, allowing

Figure 5. Changes in afterhyperpolarization associated with the Slack P924L mutation are maintained in the presence of antagonists to other potassium channels. **A**, Representative current-clamp recordings in response to current steps from Slack WT and P924L neurons in the presence of 100 μ M α -dendrotoxin, a Kv1.1/Kv1.2/Kv1.6 antagonist, or 2 mM paxilline, a BK antagonist. The threshold of the first action potential is marked with a dashed line. Scale bars, 20 mV and 20 ms. **B**, Group data from multiple cells in a control solution and with either drug added to the extracellular solution. Significance values are derived from two-tailed Student's *t* tests.

generation of additional spikes. The findings of the multielectrode array experiments, including increased burst frequency and intensity, also support the observation that neurons that express the mutant Slack channel have increased intrinsic excitability. The cells in these experiments appeared to be at an early maturational stage as indicated by the height of the action potentials and limited firing. Because MMPSI is an early onset neurodevelopmental disorder, these early excitable properties of neuronal maturation may more accurately represent a model of the disease than more mature neuronal membrane properties.

There are two general possible explanations for the increase in excitability in cells expressing the gain-of-function P924L Slack mutation. One explanation is that the increased K_{Na} current is directly responsible for increased firing rate. At first sight, this might appear counterintuitive because an increased potassium conductance is usually associated with lower excitability, the reverse of what is observed in epilepsy. Our numerical simulations of a very simple neuronal model, however, clearly demonstrate that an increase in the amplitude of afterhyperpolarizations, coupled with an increase in the numbers of action potentials evoked by a depolarizing stimulus is indeed produced by an increase in Slack conductance with no changes in any other parameter. The major assumption made in this simple model is that Slack channels are activated rapidly by Na^+ entry through voltage-dependent Na^+ channels. This is supported by experimental evidence demonstrating that K_{Na} currents are activated within milliseconds of onset of an action potential or of AMPA-mediated synaptic currents and that K_{Na} channels cluster with Na^+ channels at nodes of myelinated axons (Koh et al., 1994;

Hess et al., 2007; Nanou et al., 2008). This rapid transient activation of K_{Na} currents is kinetically distinct from slower components that have also been detected during neuronal firing and that are activated by persistent Na^+ channels (Budelli et al., 2009; Hage and Salkoff, 2012; Kaczmarek, 2013).

The simulations we have described should not be considered to represent a biophysically realistic model, nor an attempt to faithfully represent the detailed behavior of the cells. They incorporate only Na^+ , Kv and channels with no other active conductances. They have been highly simplified to illustrate the concept that high K_{Na} currents can lead to hyperexcitability. With any multiparametric model, and particularly a nonlinear one, the results are sure to depend greatly on the parameter set and initial conditions. This has been shown for example with BK channels (Montgomery and Meredith, 2012; Whitt et al., 2016), in which gain of function of a K_{Ca} current could either increase or decrease neuronal firing based on other parameters (in that case, associated with time of day in suprachiasmatic nucleus neurons). Despite the limitations, our simple model yielded the results shown without any parameter tuning and shows that it is in theory feasible for high K_{Na} currents (associated with increased fast AHP) to increase neuronal firing frequency, similar to our experimental observations. The principal constraint of the model is that Na^+ entry during the action potential must activate the K_{Na} channels sufficiently rapidly to contribute to the fast AHP.

An alternative explanation for increased excitability is that the presence of mutant channels induces compensatory changes in other conductances. Although we found no changes to Na_v1 , Kv1.1 or BK channel levels in neurons expressing the mutation, and the changes in AHP were preserved with blocking Kv1 family and BK conductances, we cannot eliminate the possibility of changes in other channel pore-forming subunits, their auxiliary subunits, or in enzymes that could influence these proteins. Such compensatory changes could explain the minor differences in the effects of the mutation at the two developmental stages we studied. Moreover, Slack channels interact with components of cell signaling pathways such as Phacx-1 and FMRP (Brown et al., 2010; Zhang et al., 2012; Fleming et al., 2016) and these interactions could be disrupted by the mutation. Understanding the potential consequences of such cellular alterations, which may result in changes beyond altered excitability, will likely require animal models rather than iPSC-derived neurons. Indeed there is evidence that compensatory changes in neurons with epilepsy-associated ion channel mutations contribute to pathology—for example loss of Nav1.1 channel activity leads to SK channel downregulation and increased thalamic bursting (Ritter-Makinson et al., 2019). Such compensatory changes might additionally influence treatment options (Isom, 2019). Nevertheless, the finding that the altered excitability in neurons with the P924L mutation closely matches that seen in simple model simulations

Figure 6. Numerical simulations of the effect of increasing Slack K_{Na} conductance in a simplified model neuron while holding other parameters constant. **A**, Progressively increasing the K_{Na} current increases afterhyperpolarizations and eventually trigger repetitive firing during maintained depolarization. **B**, Time course of membrane potential, I_{Na} , intracellular Na^+ , I_{KNa} , I_{KV} , and the leak current I_L for a simulation with a low and high value of K_{Na} conductance. **C**, **D**, Action potential activation thresholds (**C**) and height (**D**) show minimal changes. There is a progressive decrease in spike width (**E**) and increase in AHP depth (**F**) as the maximum K_{Na} conductance is increased. Thresholds were measured as the minimal stimulating current that triggered an action potential. Action potential height was measured as the membrane potential at the peak. Width was measured at 0 mV, and AHP depth was measured at the most negative membrane potential reached following the first action potential.

Figure 7. *KCNT1* P924L increases aggregate firing rates, bursting behaviors, and firing synchrony. **A**, Raster plots of action potentials from MEA recordings of WT or *KCNT1* P924L iPSC-derived neurons. Cells exhibit spontaneous firing and bursting behaviors on DIV 11. Two example wells of each condition are presented, with black tick marks denoting action potential (AP) timings across time (90 s) for all 16 electrodes within the well. Poisson burst statistics were used to capture and quantify momentary, high-frequency bursts. **B**, Firing and bursting statistics averaged across all wells ($n = 24$ per group) show that Slack P924L increases the mean firing rate ($p < 0.05$) and synchrony ($p < 0.01$), with greater burst frequency ($p < 0.01$) and higher burst intensities ($p < 0.05$). Although bursts in P924L cells are more frequent and more intense, they have shorter durations ($p < 0.01$) than control. Error bars indicate SEM.

of increased K_{Na} argues that the increase in this component of current alone is a major factor that increases excitability.

The multielectrode array results also suggest that the timing of firing in networks of neurons with the P924L Slack mutation is altered, producing an increase in measures of synchrony in the clusters, and increasing the overall activity of groups of neurons. Indeed, activation of Slack channels has previously been demonstrated to regulate the timing and accuracy of high-frequency firing of neurons in the auditory brainstem (Yang et al., 2007). Epilepsies associated with Slack mutations might, therefore, be considered disorders of hypersynchrony.

P924L is only one of many epilepsy-associated Slack mutations identified to date (Barcia et al., 2012; Heron et al., 2012; Kim et al., 2014; Martin et al., 2014; Milligan et al., 2014; Shimada et al., 2014; Møller et al., 2015; Mikati et al., 2015; Rizzo et al., 2016; Tang et al., 2016). Phenotypes cover a spectrum from severe infantile epilepsies (MMPSI or Ohtahara syndrome) to later-onset ADNFLE. Because these various mutations consistently show increased conductance when heterologously expressed, the mechanism of hyperexcitability demonstrated here is likely to be relevant regardless of the specific mutation. Similarly, specific inhibition of the Slack current would be expected to reduce hyperexcitability in any Slack-associated gain-of-function epilepsy.

Gain-of-function mutations in several other potassium channels have also been proposed to cause epilepsy, but the mechanism by which they lead to epilepsy is likely to differ. The most similar may be the BK channel (*KCMA1*), which has high structural similarity to Slack (Hite et al., 2015). Mutations that cause BK channels to open more, due to changes in either voltage dependence or calcium sensitivity, are known to cause epilepsy, often with paroxysmal dyskinesia (Du et al., 2005; Wang et al., 2009; Li et al., 2018). The mechanism is unknown but may parallel that for Slack, as both channels are cation-activated.

Another interesting parallel is Kv3.1 (*KCNC1*). This high-threshold channel is thought to speed up action potential recovery

and promote rapid firing (Rudy and McBain, 2001; Labro et al., 2015; Kaczmarek and Zhang, 2017) similar to what we propose for mutant Slack. Within the cerebral cortex, these channels are expressed in inhibitory neurons, and loss-of-function point mutations cause progressive myoclonus epilepsy (Muona et al., 2015).

A limitation of the study is that we have only evaluated the P924L homozygous condition, whereas a heterozygous mutation is sufficient to cause human epilepsy. Although the use of cells with a homozygous mutation makes analysis more straightforward, heterozygous cells may behave differently, in part because of heteromer formation and other potential interactions between WT and P924L subunits. Nevertheless, even the effects of a single copy mutation in iPSC-derived cells could be very different from those of the same mutation in a developing brain.

In addition to producing early-onset seizures, Slack mutations are associated with severe intellectual disability (Kim and Kaczmarek, 2014). It is not yet clear to what extent this results from seizures themselves or from other cellular alterations. ADNFLE can be caused either by mutations in Slack or in the neuronal nicotinic acetylcholine receptor. The fact that reported cases of Slack-associated ADNFLE have much more severe neuropsychiatric/neurodevelopmental consequences than those caused by nicotinic acetylcholine receptor mutations suggests that Slack mutations impair development beyond simply triggering seizures (Steinlein et al., 1995; Heron et al., 2012). Nonconducting functions of Slack including protein–protein interactions with Phactr-1 and FMRP, which regulate RNA translation and other aspects of cellular signaling, may provide another pathological mechanism distinct from the direct effect of the mutations on hyperexcitability. Pharmacological treatments that address both aspects of channel function may prove the most therapeutically useful.

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