Conditional deletion of \( L1CAM \) in human neurons impairs both axonal and dendritic arborization and action potential generation

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Hundreds of \( L1CAM \) gene mutations have been shown to be associated with congenital hydrocephalus, severe intellectual disability, aphasia, and motor symptoms. How such mutations impair neuronal function, however, remains unclear. Here, we generated human embryonic stem (ES) cells carrying a conditional \( L1CAM \) loss-of-function mutation and produced precisely matching control and \( L1CAM \)-deficient neurons from these ES cells. In analyzing two independent conditionally mutant ES cell clones, we found that deletion of \( L1CAM \) dramatically impaired axonal elongation and, to a lesser extent, dendritic arborization. Unexpectedly, we also detected an \( \sim 20-50\% \) and \( \sim 20-30\% \) decrease, respectively, in the levels of ankyrinG and ankyrinB protein, and observed that the size and intensity of ankyrinG staining in the axon initial segment was significantly reduced. Overexpression of wild-type \( L1CAM \), but not of the \( L1CAM \) point mutants R1166X and S1224L, rescued the decrease in ankyrin levels. Importantly, we found that the \( L1CAM \) mutation selectively decreased activity-dependent Na\(^+\)-currents, altered neuronal excitability, and caused impairments in action potential (AP) generation. Thus, our results suggest that the clinical presentations of \( L1CAM \) mutations in human patients could be accounted for, at least in part, by cell-autonomous changes in the functional development of neurons, such that neurons are unable to develop normal axons and dendrites and to generate normal APs.

X-linked neurodevelopmental disorders that produce intellectual disability are relatively common diseases resulting from mutations in X-chromosomal genes, with \( \sim 1/600-1/1,000 \) males affected (Gécz et al., 2009). One particular gene associated with X-linked intellectual disability is \( L1CAM \), with an estimated incidence of 1/25,000–1/60,000 males (Halliday et al., 1986; Weller and Gärtner, 2001). \( L1CAM \) is a type 1 transmembrane protein of the immunoglobulin superfamily that is conserved in vertebrates and invertebrates (Neuroglian/Sax-7 and that was initially described as a neuronal cell adhesion molecule involved in axonal growth (Rathjen and Schachner, 1984; Chang et al., 1987). \( L1CAM \) contains six extracellular N-terminal Ig domains, followed by five fibronectin type 3 (FN3) domains, a single transmembrane segment, and a short C-terminal intracellular sequence. \( L1CAM \) is the founding member of the \( L1CAM \) family that includes neurofascin, NrCAM, and CHL1, which exhibit similar domain structures and are composed of homologous sequences. \( L1CAM \) is highly expressed in the developing nervous system and has also been implicated in cancer progression (Schäfer and Altevogt, 2010; Kiefer et al., 2012; Schäfer and Frotscher, 2012).

Approximately 350 pathogenic mutations in the \( L1CAM \) gene have been described in patients with a broad spectrum of neurological abnormalities and mental retardation, summarized by the term L1 syndrome. This spectrum includes the MASA syndrome (mental retardation, aphasia, shuffling gait, and adducted thumbs), hydrocephalus due to stenosis of the aqueduct of Sylvius, agenesis of the corpus callosum, and SPG1 (X-linked hereditary spastic paraplegia type 1), which are referred to collectively as CRASH syndrome (Rosenthal et al., 1992; Stumpel and Vos, 1993; Jouet et al., 1994, 1995; Fransen et al., 1997; Weller and Gärtner, 2001; Vos et al., 2010). Besides a reported whole gene deletion (Chidsey et al., 2014), these mutations include frameshift, nonsense, and missense mutations, resulting in the production of truncated proteins or proteins with mutations in structurally defined key residues (Stumpel and Vos, 1993). Missense mutations most likely lead to alterations of intracellular trafficking and impaired function and mobility caused by additional cysteines on the surface of the molecule or aberrant ligand binding (De Angelis et al., 1999, 2002; Kenrick et al., 2000; Schäfer et al., 2010). Pathological mutations are known to affect binding of \( L1CAM \) to itself, Neurexin-1, Tax-1/Axonin-1, ankyrins, and integrins, or to impair triggering of epidermal growth factor receptor and Erk1/2 signaling (De Angelis et al., 1999; Schäfer and Altevogt, 2010). Overall, most
of the disease-causing mutations in L1CAM appear to be loss-of-function mutations.

Interestingly, an ethanol-binding site disrupting the interface between Ig-domains 1 and 4 of L1CAM has been identified. This site might explain the inhibitory effects of ethanol on L1CAM-mediated cell adhesion and neurite outgrowth, and could contribute to neuropathological abnormalities observed in fetal alcohol spectrum disorders, which exhibit features that are similar to those observed in L1 syndrome patients (Ramanath et al., 1996; Bearer et al., 1999; Arevalo et al., 2008). Electron microscopy studies on L1CAM and data from a crystal structure of the N-terminal Ig domains 1–4 of the L1CAM family member neurofascin, as well as a cryo-electron tomography report on liposomes supplemented with L1CAM ectodomains, revealed a horseshoe-like structure of the Ig domains 1–4 (Schirrmann et al., 2001; He et al., 2009; Liu et al., 2011). Based on the structure of the Ig domains 1–4 of the L1CAM homologue Axonin-1, it has been suggested that two horseshoes on opposing cells interact in a zipper-like manner, mediating homophilic cell adhesion (Freigang et al., 2000). Ethanol, and disease-causing missense mutations in the ethanol-binding pocket (e.g., Leu-120-Val and Gly-121-Ser), likely disrupt the horseshoe-shaped structure and inhibit homophilic and heterophilic interactions of L1CAM (Bateman et al., 1996; De Angelis et al., 1999, 2002; Arevalo et al., 2008). However, in contrast to the notion that Ig domains 1–4 are essential for homophilic binding, neurons from a reported L1CAM mutant mouse line lacking only Ig domain 6, which contains the integrin-binding motif RGD, failed to attach to L1CAM in vitro (Itoh et al., 2004), suggesting a more complicated scenario for the homophilic activity of L1CAM on neurons.

Studies using constitutive L1CAM-deficient mice as a model system reported defects in axon guidance in the corticospinal tract, impaired growth of pyramidal layer V neuron apical dendrites, reduced size of the corpus callosum, malformations of the ventricular system and the cerebellar vermis, decreased association of axons with nonmyelinating Schwann cells, and reduced inhibitory synaptic transmission (Dahme et al., 1997; Cohen et al., 1998; Fransen et al., 1998; Demyrenko et al., 1999; Saghatelany et al., 2004). Puzzlingly, mutant mice expressing L1CAM with a truncated intracellular domain that lacks the ankyrin-binding region displayed no abnormalities observed in fetal alcohol spectrum disorders, which exhibit features that are similar to those observed in L1 syndrome patients (Ramanath et al., 1996; Bearer et al., 1999; Arevalo et al., 2008). Electro...
variation or incidental genetic changes, and were converted into cKO cells by removal of the puromycin resistance cassette using transduced flp recombinase (Fig. 3 B).

We produced iN cells from both L1CAM cKO ES cell clones by expression of Ngn2 delivered by recombinant lentiviruses. We also expressed WT active cre-recombinase (Cre) or mutant inactive cre-recombinase (ΔCre) in the iN cells during induction to generate precisely matching L1CAM-deficient mutant neurons and control neurons, respectively (Fig. 3, C and D; mutant neurons from the two clones are referred to as −/y #1 and −/y #2, and WT neurons as +/y #1 and +/y #2). As analyzed by immunoblotting, we found that, consistent with previous studies (Wolff et al., 1988; Schäfer and Altevogt, 2010), control neurons produced multiple L1CAM protein variants with apparent molecular weights between 220 and 65 kD (Fig. 3 E). Upon cre recombination, neurons derived from the two cKO clones showed a complete lack of all L1CAM variants, and exhibited no L1CAM expression as judged by immunocytochemistry (Fig. 3, E and F), demonstrating that the two cKO ES cell clones can be efficiently converted into control and L1CAM mutant human neurons with precisely matching backgrounds that only differ in the expression of L1CAM.

No change in survival, but decreased MAP2a/b and ankyrin expression in L1CAM mutant human neurons

L1CAM-deficient human neurons exhibited similar survival as control neurons, with no evidence of cell death or degeneration as a result of the L1CAM deletion (Fig. 3, G and H). Furthermore, we observed no increase in phosphorylation of JNK/SAPK, which is activated in stressed cells (Fig. 4, C and D). Quantitative immunoblotting analyses revealed that the L1CAM deletion did not alter the majority of tested neuronal marker proteins; in particular, no changes in the levels of the majority of synaptic proteins were detected (Fig. 4, A and B). In contrast, we observed a decrease in the levels of the adaptor proteins ankyrinG and ankyrinB, which are axonal proteins that bind to the intracellular sequences of L1CAM (Davis and Bennett, 1994). We found a reduction of ~50% and ~20% of ankyrinG and ankyrinB, respectively (Fig. 4, A and B). In addition, we detected a ~30% decrease in the dendritic marker MAP2a/b and complexins. Because it was reported that L1CAM-activated signaling stimulates the mitogen-activated protein kinase (MAPK) pathway (Schaefer et al., 1999; Schmid et al., 2000; Maness and Schachner, 2007), we examined iN cells derived from both cKO clones for changes in MAPK activation, but did not detect significant changes in phosphorylation levels of Erk 1/2 or related components in the mutant neurons (Fig. 4, C and D). Together, our data suggest that the L1CAM deletion in human neurons does not significantly impair neuronal survival or activate MAPK signaling, but decreases the levels of the axonal proteins ankyrinB and ankyrinG, the dendritic protein MAP2a/b, and the presynaptic protein complexin.

Axonal and dendritic arborizations are reduced in human L1CAM mutant neurons

To test for changes in neuronal morphology, we next sparsely transfected (by the CaPO4 method) conditionally hemizygous mutant iN cells with a plasmid expressing TetO-EGFP and stained the neurons for MAP2a/b. Assuming that cytoplasmic EGFP visualizes the complete neuronal morphology and that MAP2a/b is selectively expressed in dendrites, this experiment allowed us to systematically distinguish between dendrites and axons (Fig. 5 A). By comparing mutant with WT cells, we observed in neurons derived from both independent cKO clones

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Figure 1. Axonal localization of L1CAM in human WT neurons. (A–C) Representative confocal micrographs (from three independent experiments) of L1CAM on human neurons (in cells) obtained by forced expression of Ngn2 in ES cells and analyzed 5 wk after Ngn2 induction. Neurons were double labeled by immunofluorescence for L1CAM and MAP2 (A), TuJ1 (B), or Neurofilament-L (C). Each panel shows lower (top) and higher magnifications (bottom; boxed area in lower magnification images).
Figure 2. L1CAM partially colocalizes with MAP2 in the dendrites of human neurons at early developmental stages. (A) Representative confocal micrographs of L1CAM on human neurons taken 4 and 21 d after Ngn2 induction. Neurons were triple-labeled by immunofluorescence for L1CAM, MAP2, and TuJ1. (B and D) Intensity profile of immunolabeling for the indicated proteins derived from line scans along the dashed line in the bottom left and second from the right images in A. (C and E) Summary graphs of Pearson's correlation coefficients for co-localization of L1CAM versus MAP2 (C) and L1CAM versus TuJ1 (E) determined in multiple experiments at different times after Ngn2 induction as indicated (mature = 35 d after Ngn2 induction). Data are means ± SEM; statistical comparisons were performed by Student's t test comparing various times after Ngn2 induction to mature neurons (*, P < 0.05; **, P < 0.01; ***, P < 0.001; numbers of images/independent experiments analyzed are shown in the bars).
Figure 3. Generation of human ES cells with a conditional L1CAM gene loss-of-function mutation and effect of L1CAM mutation on neuronal survival. (A) L1CAM contains six Ig-domains and five FN-type 3 domains followed by a single transmembrane sequence, and a highly conserved short intracellular cytoplasmic domain. (B) Targeting strategy. The L1CAM gene was mutated by homologous recombination in male H1 ES cells using a recombinant AAV, and correctly recombined mutant ES cells were verified by PCR. The introduced mutations flank an out-of-frame exon (exon 3) with loxP sites, and include a puromycin (Purom.) resistance cassette for selection; this cassette was removed using Frt-recombinase transduction to generate L1CAM cKO cells. For generation of precisely matched neurons containing or lacking L1CAM, mutant Cre-recombinase (ΔCre) or active Cre-recombinase (Cre) were expressed in iN cells during induction of neuronal differentiation as indicated. (C) Design of the lentiviruses used for conversion of ES cells into iN cells and for expression of inactive (ΔCre) or active Cre-recombinase (Cre). (D) Flow diagram of all experiments. cKO cells were infected at day –1 with lentiviruses shown in B, and differentiation was started on day 0 by activating Ngn2 expression with doxycycline. Analyses were conducted at least 21 d after induction of Ngn2. (E) Ponceau-stained blotting membranes (left) and immunoblots (right) of iN cells derived from two independent L1CAM cKO ES cell clones (#1 and #2). Immunoblots were stained with antibodies to human L1CAM (green) and GDI (red; loading control); signals were visualized with fluorescently labeled secondary antibodies (right). (F) Representative immunofluorescently labeled control and mutant L1CAM-deficient neurons. L1CAM, MAP2, and DAPI staining is shown. (G and H) Plot of the fraction of surviving neurons as a function of time in culture (G; dashed line indicates control neurons). As in all experiments, neurons from two independent L1CAM cKO ES cell clones (blue and orange) were analyzed. Data are means ± SEM (n = 3 independent biological experiments).
that deletion of L1CAM caused a massive decrease in the total length and number of branching points of axons, but produced no changes in cell soma size (Fig. 5, A, C, D, and G). Moreover, deletion of L1CAM decreased the total length and branch numbers of dendrites, although the effect on dendrites was smaller than that on the length and branches of axons (Fig. 5, E and F), suggesting that the partial codistribution of L1CAM and MAP2 in very young neurons is important for dendritic tree development (Fig. 2). Interestingly, ∼15–25% of L1CAM mutant neurons developed two axons, compared with only 0–3% of WT neurons (Fig. 5, B and H). To test whether this morphological phenotype persists at later developmental stages, we analyzed iN cells at 180 d after induction. Because of the immense axonal growth constraining single-cell axon analyses (Fig. 6, A and B), we measured the dendritic arborization of these “old” human neurons and detected a defect in the mutant neurons (Fig. 6, C–F). Together with the quantitative immunoblotting results and the localization of L1CAM on developing and mature human neurons, these observations indicate that loss-of-function mutations of L1CAM impair dendritic and axonal arborization of human neurons.

L1CAM deletion selectively decreases ankyrinG levels in the AIS

The marked decrease of ankyrinG levels in hemizygous mutant neurons (Fig. 4, A and B) suggests a potential defect in the AIS because ankyrinG is a key adaptor molecule that clusters Na+ and K+ channels at the AIS and partially co-localizes with L1CAM (Fig. 7 A; Zhou et al., 1998; Garrido et al., 2003; Kole and Stuart, 2012). To assess this aspect, we measured the area and intensity of ankyrinG staining in the AIS and found that both parameters decreased in L1CAM mutant human neurons (Fig. 7, C, E, and F). Investigating ankyrinB, which also co-localizes with L1CAM in axons...
we additionally detected a ∼30% decrease in ankyrinB staining intensity in mutant axons (Fig. 7, D and G). Together with the changes in axonal arborization (Fig. 5), these results suggest a fundamental role of L1CAM in the morphological development of axons in human neurons.

No decrease in synapse density in human mutant neurons

To test the hypothesis that L1 syndrome symptoms and its associated severe intellectual disability are caused by morphological changes at the synaptic level in human neurons, we quantified presynaptic synapsin-positive puncta (which partially overlap with L1CAM on axons from WT neurons; Fig. 8 A) on post-synaptic MAP2-positive dendrites. Both the size and the number of synapses per dendritic segment in WT and mutant neurons were compared (Fig. 8 B), and no significant differences could be detected (Fig. 8, C and D). No alterations of the intrinsic electrical properties were detected (Fig. 8, E–H), suggesting no alterations in the membrane properties.
composition. In summary, our results suggest that L1CAM is not required for synapse development as such, although we cannot exclude any functional synaptic changes at this point.

**L1CAM deletion impairs AP generation and decreases neuronal excitability in mutant human neurons**

To investigate if L1CAM deficiency, and thus a decrease of ankyrinG at the AIS, leads to functional consequences in AP firing, we performed patch-clamp recordings and measured the whole-cell density of Na\(^+\) and K\(^+\) currents in 5-wk-old iN cells in voltage-clamp configuration (Fig. 9, A–D). Cells were held at −80 mV, and then the membrane potential was sequentially depolarized in 10-mV steps for 2 s (Fig. 9, A and B). We observed a significant decrease in the peak amplitude of Na\(^+\) currents (inward), whereas K\(^+\) currents (outward) measured in the same experiments remained unchanged (Fig. 9, C and D). Thus, the removal of L1CAM seems to selectively reduce the total density of functional Na\(^+\) channels in human neurons.

To test whether the reduction in Na\(^+\) channels impacts AP generation, we performed current-clamp recordings and sequentially depolarized the cells by injecting positive current. We found that L1CAM hemizygous mutant iN cells required larger current injections to reach AP threshold than precisely matching WT iN cells (Fig. 9, E–G, and M). We also assessed the firing threshold and the shape of the APs, but found no statistically significant differences between control and mutant cells (Fig. 9, H–L and N–R). These results indicate that deletion of L1CAM expression in human neurons leads to a partial loss of ankyrinG from the AIS, a decrease in the density of Na\(^+\) channels, and a change in a neuron’s response to somatic current injections.

**Ankyrin-binding site of L1CAM is required for normal expression levels of ankyrinG and ankyrinB in human neurons**

To evaluate the mechanistic role of the binding of ankylins to L1CAM in maintaining expression levels of ankylins, we next performed rescue experiments by lentiviral-mediated expression (infection on day −1; Fig. 3 D) in iN cells of WT L1CAM, or mutant L1CAM. The two pathological point mutations, R1166X and S1224L, either completely lack the ankyrin-binding site or contain a substitution in a key residue of the ankyrin-binding site (Fig. 10 A; Vos and Hofstra, 2010). Overexpressed WT, as well as R1166X and S1224L mutant, L1CAM could be detected by immunoblots at similar levels, and were both expressed on the surface of human neurons.
Interestingly, in contrast to WT L1CAM, R1166X, and S1224L mutant L1CAM did not rescue the decrease of ankyrin levels in mutant human neurons (which in this experiment were suppressed to ∼20% for ankyrinG and ∼30% for ankyrinB), suggesting a requirement of the L1CAM ankyrin protein interaction for the stabilization of ankyrins (Fig. 10, D–F).

**DISCUSSION**

In this study, we used the cre/lox technology in human neurons to investigate the effect of hemizygous L1CAM loss-of-function mutations on neuronal function. This experimental design eliminates potentially confounding effects induced by genetic background changes or selection of cell clones. So far, hundreds of mutations in the L1CAM gene (which is X-chromosomal) were reported in patients with severe developmental malformations, neurological abnormalities, and intellectual disability (Rosenthal et al., 1992; Stumpel and Vos, 1993; Jouet et al., 1994, 1995; Fransen et al., 1997; Vos et al., 2010; Adle-Biassette et al., 2013; Chidsey et al., 2014). Most of the disease-causing L1CAM mutations likely represent loss-of-function mutations. We found that conditionally L1CAM-deficient human neurons manifest three types of defects: first, a large reduction in the size of axons, a smaller decrease in dendritic arborizations, and a significant reduction in the size of the AIS; second, a decrease in the levels of ankyrinG and ankyrinB; and third, impairment of AP generation. It seems likely that these three types of defects are commonly produced by the loss of the L1CAM complex with ankyrins upon deletion of L1CAM, and that these defects are responsible, at least in part, for the symptoms of patients with L1 syndrome.

In mutant mice and in vitro systems, it has been well established that the intracellular segment of L1CAM binds to ankyrins (Davis and Bennett, 1994; Bennett and Chen, 2001). In general, it has been shown that ankyrinB binding promotes neuritogenesis by coupling L1CAM to the actin cytoskeleton (Nishimura et al., 2003). Binding L1CAM to ankyrinG, conversely, is thought to recruit L1CAM to the AIS, a highly specialized compartment in the proximal axon where APs are generated (Malhotra et al., 1998; Yoshimura and Rasband, 2014). Mice deficient for ankyrinB exhibit reduced levels of L1CAM on these axons and, because of their hypoplasia of the corpus callosum and dilated ventricles, share features of L1 syndrome patients (Scotland et al., 1998). Furthermore, it has been
shown that the cytoplasmic SFIGQY motif of L1CAM (aa 1224–1229, which is present in all L1CAM family members and is highly conserved from invertebrates to vertebrates) is essential for ankyrin binding (Garver et al., 1997; Needham et al., 2001). By introducing two pathological missense mutations, namely S1224L and Y1229H, reduced recruitment of ankyrin to the plasma membrane in vitro was observed (Needham et al., 2001). Knock-in mice expressing the missense mutation Y1229H exhibit mistargeting of mutant ganglion cell axons from the ventral retina to lateral sites in the contralateral superior colliculus (Buhusi et al., 2008). Additionally, observations of neonatal L1CAM-deficient mice suggest a decrease of ankyrinB expression in thalamocortical and corticothalamic axons, together with axonal hyperfasciculation and pathfinding errors (Buhusi et al., 2008). Additionally, observations of neonatal L1CAM-deficient mice suggest a decrease of ankyrinB expression in thalamocortical and corticothalamic axons, together with axonal hyperfasciculation and pathfinding errors (Wiencken-Barger et al., 2010), raising questions about the precise role of the intracellular L1CAM sequence.

In the present study, we show that L1CAM codistributes with ankyrinG and ankyrinB and that the deletion of L1CAM decreases ankyrins on the subcellular levels (ankyrinG at the AIS, ankyrinB on axons; Fig. 7), and on the global protein level (as revealed by quantitative immunoblot analyses; Fig. 4, A and B), suggesting that L1CAM may actually recruit and stabilize ankyrins to specific subcellular localizations. In addition to previous studies, our data show that on human axons L1CAM is not only important for the expression of ankyrinB and axonal arborizations but it is also crucial to the levels of ankyrinG at the AIS and its proper function in Na⁺-mediated AP generation. It is tempting to speculate that L1CAM and ankyrins are interdependent subunits of a complex stabilizing each other, and thereby regulating axonal morphological and functional development. Evidence for this notion comes from our rescue experiments, suggesting that the integrity of the ankyrin-binding site is important for the stabilization of ankyrins (Fig. 10).

AnkyrinG is a key component of the AIS and has been shown to be responsible for clustering Na⁺ and K⁺ channels at the AIS (Zhou et al., 1998; Garrido et al., 2003; Kole and Stuart, 2012). In the absence of ankyrinG in mouse neurons, Na⁺ currents are highly reduced and the shape of the AP is altered (Zhou et al., 1998; Hedstrom et al., 2008; Barry...
This phenotype is similar to the Na\(^+\) channel phenotype we observed in L1CAM-deficient human neurons. AnkyrinG mutations in patients have been associated with severe cognitive deficits, autism, sleeping disorder, severe attention deficit hyperactivity disorder, and bipolar disorder (Ferreira et al., 2008; Schulze et al., 2009; Iqbal et al., 2013).

To date, a change in neuronal excitability has not been observed in L1CAM mouse mutants and could represent a...
feature that is more easily distinguished in human neurons (Dahme et al., 1997; Cohen et al., 1998; Fransen et al., 1998; Demyanenko et al., 1999; Law et al., 2003; Saghatelyan et al., 2004; Nakamura et al., 2010). However, a disintegration of the AIS, APs with modified waveform upon stimulation, abolition of spontaneous firing in Purkinje cells, and deficits in motor learning have been reported for mice deficient for neurofascin in adult cerebellar Purkinje cells. But in developing neurons, neurofascin was shown to be dispensable for AIS assembly, while ankyrinG staining intensities were unchanged (Zonta et al., 2011). Neurofascin is thought to be essential for the clustering of Na⁺ channels at the nodes of Ranvier, another excitatory region of the axon, by its binding to ankyrinG (Sherman et al., 2005).
We observed an ∼20–50% decrease in ankyrinG levels, possibly leading to a decrease in the concentration of voltage-gated Na⁺ channels at the AIS. This result might explain the rather modest reduction of Na⁺ currents and the lack of significant changes in the shape of APs. However, this points to an important aspect of L1 syndrome because the missing L1CAM–ankyrin interactions likely contribute to the clinical presentations of L1CAM and ankyrin mutations. Future work could use this model system to test ways of restoring neuronal excitability, for example, by reducing the activity of resting K⁺ channels at the AIS. It has been shown that pharmacological inhibition of Cyclin-dependent kinase 2 or 5 (Cdk2 or Cdk5) by roscovitine, enriched K⁺ channels at the AIS (Vacher et al., 2011). One possible way to reduce K⁺ channel activity would be to pharmacologically activate Cdk2 or Cdk5 function.

Overall, our study using cre/lox technology on human neurons presents a first step in understanding L1 syndrome at the neuronal level. Our results confirm a morphological phenotype previously observed in animal models (Dahme et al., 1997; Cohen et al., 1998; Demyanenko et al., 1999), but also present an additional component in the human mutant neuron’s reduced ability to generate APs.

MATERIALS AND METHODS

Viral constructs. The following lentiviral constructs were used: FUW-TetO-Ng2-T2A-puromycin expressing TetO-Ng2-T2A-puromycin cassette (TetO promoter drives expression of full-length mouse Ng2 and of puromycin via the cleavage-peptide sequence T2A; Fig. 3 B); F-Ubiquitin-W (FUW)-rTATA containing rTATA; FUW-TetO-EGFP expressing EGFP; FSW-NLS-mCherry expressing mCherry preceded by a nuclear localization sequence (under the control of neuron-specific human synapsin promoter) to monitor cell death; FUW-Flp to express Flp-recombinase; FUW-GFP::Cre to express Cre-recombinase to delete L1CAM exon 3 and create via frameshift a null-allele or GFP::ΔCre for the wild-type control; FUW-TetO-wild-type-L1CAM; FUW-TetO-R1166X-L1CAM; and FUW-TetO-S1224L-L1CAM. One adeno-associated virus (AAV) construct was used for gene targeting of X-chromosomal L1CAM allele of the male hESCs H1 (also shown in Fig. 3 B): the construct contains sequences from the region encoding exon 3 flanked by loxP sites and frt-sites flanked puromycin resistance cassette (Frt-PGK-Puro-SV40polyA-Frt) adjacent to the 5’ loxP site. For homologous recombination, the 5’ arm of the construct includes 1.6 kb of sequences located upstream of exon 3. The 3’ arm contains 1.4 kb of sequences located downstream of exon 3.

Virus generation. Lentiviruses were produced as previously described (Zhang et al., 2013). H1 ES cells (WiCell Research Resources) were maintained as feeder-free cells in mTeSR1 medium (Stem Cell Technologies). Mouse glial cells were cultured from the forebrain of newborn WT CD1 mice. In brief, newborn mouse forebrain homogenates were digested with papain and EDTA for 30 min, and cells were dissociated by harsh trituration to avoid growing of neurons, and then plated onto T75 flasks in DMEM supplemented with 10% FBS. Upon reaching confluence, glial cells were trypsinized and replated at lower density a total of three times to remove potential trace amounts of mouse neurons before the glia cell cultures were used for co-culture experiment with iN cells.

Gene targeting in ES cells. H1 ES cells were transduced by AAVs to generate conditional hemizygous mutant ES cells by flanking exon 3 (ENSE00003680685, coding the aa MEPPPV1TEQSPRRLVVFPTDDISLKEASGKPEV in the first Ig domain), which results in a frameshift and a premature stop codon upon cre recombination. Puromycin (1 µg/ml) was added and kept in mTeSR1 medium 2 d after transduction. The surviving ES cells were allowed to grow into colonies, and then individually picked. 7 correctly targeted colonies out of 76 colonies in total were confirmed by PCR screening using oligo sequences 5’-ACAGATGACATCGACCTC AAGTGTGAGG-3’ and 5’-GACTGGGAGATGGCGAGG ACTTG-3’, resulting in a 240-bp band for a correctly targeted clone or 185 bp for the untargeted WT. Correctly mutated alleles were confirmed after flp and cre recombination and iN differentiation.

Standard protocol for generation of iN cells from conditional mutant human ES cells and for activating the conditional mutations. iN cell generation has been previously using calcium phosphate. Lentiviruses were harvested in the medium 48 h after transfection, pelleted by centrifugation (49,000 g for 90 min), resuspended in MEM, aliquoted, and frozen at −80°C. Only virus preparations with >90% infection efficiency as assessed by EGFP expression or puromycin resistance were used for experiments. AAV-DJ was used to deliver the targeting construct for generation of cKO cells. AAV-DJ was produced in HEK293T cells by co-transfection of pHelper, pDJ, and AAV vector (8.5 µg of DNA per 75 cm² culture area) using calcium phosphate. Cells were harvested 72 h after transfection in PBS/1 mM EDTA and after one freezing/thawing cycle. AAVs were collected from cytoplasm using Benzonase nuclease at a final concentration of 50 U/ml at 37°C for 30 min. After clearing the suspension from cell debris by slow centrifugation (3,000 g for 30 min) AAVs were isolated after fast centrifugation (400,000 g for 120 min) in iodixanol (gradient from 15–60%) from the 40% layer and further concentrated using centricon concentrating tube (100,000 MWCO; EMD Millipore) according to the manufacturer’s suggested protocol.

Cell culture. Experiments were performed as previously described (Zhang et al., 2013). H1 ES cells (WiCell Research Resources) were maintained as feeder-free cells in mTeSR1 medium (Stem Cell Technologies). Mouse glial cells were cultured from the forebrain of newborn WT CD1 mice. In brief, newborn mouse forebrain homogenates were digested with papain and EDTA for 30 min, and cells were dissociated by harsh trituration to avoid growing of neurons, and then plated onto T75 flasks in DMEM supplemented with 10% FBS. Upon reaching confluence, glial cells were trypsinized and replated at lower density a total of three times to remove potential trace amounts of mouse neurons before the glia cell cultures were used for co-culture experiment with iN cells.
described (Zhang et al., 2013). In brief, targeted and flp-recombined (after blasticidin selection) human ES cells were treated with Accutase (Innovative Cell Technologies) and plated as dissociated cells in 24-well plates (10⁴ cells/well) on day −1 (Fig. 3 C). Cells were plated on Matrigel-coated (BD) coverslips in mTeSR1 containing 2 mM thiazovivin (BioVision). At the same time point, lentiviruses prepared as described in the paragraph "Virus generation"(0.3 µl/well of 24-well plate) were added. Two different types of lentiviruses were co-infected: the lentiviruses used for iN cell induction as described, and lentiviruses expressing either Cre-recombinase (to create a null allele) under control of the ubiquitin promoter or an inactive mutated ΔCre-recombinase for the WT control (Fig. 3 C). On day 0, the culture medium was replaced with N2/DMEM/F12/NEAA (Invitrogen) containing human brain-derived neurotrophic factor (BDNF; 10 ng/ml; PeproTech), human NT-3 (10 ng/ml; PeproTech), and mouse Laminin-1 (0.2 µg/ml; Invitrogen). Doxycycline (2 µg/ml; Sigma-Aldrich) was added on day 0 to induce TetO gene expression and retained in the medium until the end of the experiment. On day 1, a 24-h puromycin selection (1 µg/ml) period was started. On day 2, mouse glia cells were added in neurobasal medium supplemented with B27/Glutamax (Invitrogen) containing BDNF, NT3, and Laminin-1; Ara-C (2 µM; Sigma-Aldrich) was added to the medium to inhibit astrocyte proliferation. After day 2, 50% of the medium in each well was exchanged every 2 d. FBS (2.5%) was added to astrocyte proliferation. After day 2, 50% of the medium in each well was exchanged every 2 d. FBS (2.5%) was added to the culture medium on day 10 to support astrocyte viability, and iN cells were assayed after at least 21 d or as indicated.

Quantification of neuronal morphology. iN cells were sparsely transfected by calcium phosphate on day 19 with a TetO-EGFP expression vector. On day 21, cells were fixed and stained for EGFP and MAP2a/b to discriminate between dendrites and axons. Fixation at later time points would result in too abundant axonal growth. All MAP2- and EGFP-positive neurites were considered to be dendrites and all MAP2-negative but EGFP-positive neurites to be axons. Images were acquired using an A1RSi confocal microscope system (Nikon), with a 10× or 20× objective at room temperature. Images of ∼10–30 neurons per condition (n = 1 for axons; n = ~100 in case dendrites) were reconstructed using the MetaMorph neurite application, scoring for total dendritic length, dendritic branch points, total axonal length, axonal branch points, and soma area. In the case of old human neurons, cells were fixed after DIV180 and immunolabeled.

Immunofluorescence and immunoblotting experiments. Immunofluorescence experiments were performed essentially as previously described (Zhang et al., 2013). In brief, cultured iN cells were fixed in 4% paraformaldehyde and 4% sucrose in PBS for 20 min at room temperature, washed three times with PBS, and incubated in 0.2% Triton X-100 in PBS for 10 min at room temperature. Cells were blocked in PBS containing 5% goat serum for 1 h at room temperature. Primary antibodies were applied overnight at 4°C, cells were washed in PBS three times, and fluorescently labeled secondary antibodies (Alexa Fluor; 1:1,000) were applied for 1 h at room temperature. The following antibodies were used in immunocytochemistry experiments: MAP2 a/b (Sigma-Aldrich; 1:1,000), MAP2 a/b (AB5622; EMD Millipore; 1:500), MAP2 (CPA-680; EnCor; 1:2,000), Synapsin (E028; 1:1,000), ankyrinG (106/36; Neuromab; 1:500), ankyrinB (105/13; Neuromab; 1:500), Neurofilament-L (171002; Syntactic Systems; 1:500), and GFP (Invitrogen; A1122; 1:2,000). To label L1CAM on human neurons, in some cases (Fig. 7) a post-fix staining using polyclonal Rb antibody (10140-R004-50; Sinobiological) was performed. In all other cases, a live staining was performed. The human-specific clone UJ127.11 (Sigma-Aldrich) was given into fresh culturing media for 20 min at 37°C (10 µg/ml). Next, cells were fixed and washed as described in this paragraph. Images were taken using an A1RSi confocal microscope system (Nikon), with a 10×, 20×, or 60× objective at room temperature. All quantitative immunoblotting experiments were performed with fluorescently labeled secondary antibodies (LiCor; 1:5,000). Samples were separated by SDS-PAGE under reducing conditions and transferred onto nitrocellulose membranes. Blots were blocked in Tris-buffered saline containing 0.1% Tween 20 (Sigma-Aldrich) and 5% fat-free milk for 2 h at room temperature. The blocked membrane was incubated in blocking buffer containing the primary antibody overnight at 4°C, followed by three to five washes. The washed membrane was incubated in blocking buffer containing secondary antibody for 2 h at room temperature. Blots were scanned with the Odyssey-system (LiCor), followed by quantification with ImageStudio software (LiCor). For immunodetection, the following antibodies were used: NeuN (ABN78; EMD Millipore), TuJ1 (MMS-435P; Covance), Complexin 1/2 (L668), SNAP25 (P913), Synaptophysin-2 (P939), Synapsin (E028), Syntaxin-1 (438B), β-Actin (A1978; Sigma-Aldrich), Synaptophysin (Synaptic Systems, 7.2), Gβδ-dissociation inhibitor (Synaptic Systems; GDI; 81.2), calmodulin-associated serine/threonine kinase (BD; clone 7/CASK), L1CAM (UJ127.11; Sigma-Aldrich), L1CAM (ab123990; Abcam), SynCAM (T2412), MAP2 a/b (AB5622; EMD Millipore; 1:500), MAP2 a/b/c/d (Sigma-Aldrich; SABA300658; 1:500), Neurofilament-L (171002; SySy; 1:500), Neurofilament-H (Abcam; RT97; 1:500), ankyrinG (Neuromab; 106/20; 1:500), ankyrinB (105/13; Neuromab; 1:500), Tau-5 (EMD Millipore; MAB361; 1:1,000), PSD95 (Abcam; ab76115; 1:500), GluR1 (Abcam; ab1504; 1:500), CReB (Cell Signaling Technology; 86B10; 1:1,000), phospho-CReB (Cell Signaling Technology; 87G3; 1:1,000), Erk1/2 (Cell Signaling Technology; 137F5; 1:1,000), phospho-Erk1/2 (Cell Signaling Technology; E10; 1:1,000), JNK (Cell Signaling Technology; 9252; 1:1,000), phospho-JNK (Cell Signaling Technology; G9; 1:1000), Akt (Cell Signaling Technology; SG3; 1:1,000), phospho-Akt (Cell Signaling Technology; 9271; 1:1,000).
Quantification of synaptic density and survival. For synaptin puncta analyses, images were acquired using a A1R Si confocal microscope system (Nikon; 60× objective at room temperature), and the puncta density was determined using the software NIS-Elements (Nikon). Analyses of the survival of iN cells was directly monitored using images of iN cell nuclei expressing mCherry at the same position of the culture dish taken every other day, using a DFC400 digital camera (Leica) attached to a DMIIL LED inverted microscope with a 5× objective (Leica), driven by Application Suite image-acquisition software (Leica). Number of cells was determined using ImageJ software (National Institutes of Health). For each experimental condition, the mean of the pictures from 20–30 culture wells (of a 96-well culture plate) was considered as \( n = 1 \). In total, a mean of \( n = 3 \) was calculated.

Electrophysiology. All electrophysiological recordings were performed using whole-cell patch clamp. In brief, 5 wk after iN induction, on the day of recording, a coverslip containing relatively low-density induced human neurons was placed in a recording chamber mounted onto an Axioskop 2F upright microscope (Zeiss) equipped with DIC and fluorescence capabilities. Cells were approached under DIC with ~2 MΩ pipettes pulled from borosilicate glass (Warner Instruments, Inc.) using a vertical PC-10 puller (Narishige), and impaled until a high-resistance seal (GΩ) was formed between the recording pipette and the cell membrane. Next, whole-cell configuration was established by gentle application of negative pressure trough the recording pipette. The recording pipette contained (in mM): 125 K-gluconate, 20 KCl, 10 Hepes, 0.5 EGTA, 4 ATP-Magnesium, 0.3 GTP-Sodium, and 10 Na-Phosphocreatine; osmolarity, 312 mOsm, pH 7.2, adjusted with KOH. Electrical signals were recorded at 25 kHz with a two-channel Axoclamp 700B amplifier (Axon Instruments) and digitalized with a Digidata 1440 digitizer (Molecular Devices) that was, in turn, controlled by Clampex 10.1 (Molecular Devices). All recordings were performed at ~24°C.

For whole-cell voltage-clamp recordings, neurons were maintained at ~80 mV holding potentials. Series resistance varied between 4–10 MΩ. All iNs in which series resistance was higher than that were not included in the analysis. In all current-clamp experiments, the membrane potential was maintained approximately ~80 mV by constant injection of negative current through the recording pipette. Samples in the recording chamber were continuously perfused with oxygenated (95% O\(_2\)/5%CO\(_2\)) bath solution containing (in mM): 125 NaCl, 2.5 KCl, 1 MgCl\(_2\), 2 CaCl\(_2\), 25 glucose, 1.25 NaH\(_2\)PO\(_4\), 0.4 ascorbic acid, 3 myo-inositol, 2 Na-pyruvate, and 25 NaHCO\(_3\), pH 7.4, and 315 mOsm.

Data presentation and statistics. All data shown are means ± SEMs. Number of measured cells and independent experiments is indicated inside each bar, or mentioned in the figure legend. All statistical analyses were performed using either two-tailed Student’s t test or two-way ANOVA, comparing the test sample to the control sample examined in the same experiments.

Study approval. The present study was approved by Stem Cell Research Oversight (SCRO) at the Stanford University Research Compliance Office (SCRO 518: Studying brain diseases affecting synaptic transmission by using human iNs). Experiments involving animals were approved by the Stanford University Institutional Animal Care and Use Committee, Administrative Panel on Laboratory Animal Care Research Compliance Office.

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