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Kv7.2 regulates the function of peripheral sensory neurons

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Abstract

The Kv7 (KCNQ) family of voltage-gated K⁺ channels regulates cellular excitability. The functional role of Kv7.2 has been hampered by the lack of a viable *Kcnq2*-null animal model. In this study, we generated homozygous *Kcnq2*-null sensory neurons using the Cre-Lox system; in these mice, Kv7.2 expression is absent in the peripheral sensory neurons, whereas the expression of other molecular components of nodes (including Kv7.3), paranodes, and juxtaparanodes is not altered. The conditional *Kcnq2*-null animals exhibit normal motor performance, but have increased thermal hyperalgesia and mechanical allodynia. Whole cell patch recording technique demonstrates that *Kcnq2*-null sensory neurons have increased excitability and reduced spike frequency adaptation. Taken together, our results suggest that the loss of Kv7.2 activity increases the excitability of primary sensory neurons.

Keywords

KCNQ; M-current; dorsal root ganglion; nociceptors; Kv7

INTRODUCTION

The Kv7 (KCNQ) family of K⁺ channels is made up of five members, Kv7.1 to Kv7.5 (Jentsch, 2000, Delmas and Brown, 2005, Brown and Passmore, 2009). Many peripheral nervous system (PNS) and central nervous system neurons express Kv7.2, Kv7.3, and Kv7.5, and Kv7.1 and Kv7.4 have recently been found in certain neuronal populations (Goldman et al., 2009, Heidenreich et al., 2011). In addition, Kv7.1 and Kv7.4 are prominently expressed in the cochlea, and Kv7.1 is expressed in the heart (Brown and Passmore, 2009). Kv7

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channels contribute to the normal resting membrane potential, and also form the non-inactivating M-current, which is modulated by muscarinic agonists (Brown and Adams, 1980, Wang et al., 1998). Their importance in maintaining normal cellular excitability is demonstrated by the effects of dominant mutations of *KCNQ1*, *KCNQ2*, *KCNQ3*, and *KCNQ4* – all decrease the Kv7 current and cause hereditary diseases in a cell autonomous manner (Singh et al., 1998) – in the heart (*KCNQ1*), brain (*KCNQ2* and *KCNQ3*), and inner ear (*KCNQ1* and *KCNQ4*) (Jentsch, 2000, Brown and Passmore, 2009). One *KCNQ2* mutation causes neuromyotonia (Dedek et al., 2001), a form of peripheral nerve hyperexcitability, which is likely the result of diminished Kv7.2 and Kv7.3 current at nodes of Ranvier, where Kv7.2 and Kv7.3 are localized (Devaux et al., 2004). A conserved ankyrin-G binding motif located at the C-termini mediates the localization of Kv7.2 and Kv7.3 at nodes and axon initial segments (AIS) (Pan et al., 2006).

Previous investigations of the role of Kv7 in regulating neuronal excitability and nociceptive behaviors utilized pharmacological M-channel blockers and/or enhancers (Passmore et al., 2003, Yue and Yaari, 2004, Rivera-Arconada and Lopez-Garcia, 2006, Lang et al., 2008, Roza and Lopez-Garcia, 2008). Because homozygous *Kcnq2*-null mice die at birth due to pulmonary atelectasis, investigators studied heterozygous *Kcnq2*-knockout mice (Watanabe et al., 2000, Yang et al., 2003, Otto et al., 2006, Tzingounis and Nicoll, 2008), *Kcnq3*-null mice (Tzingounis and Nicoll, 2008), or mice expressing a dominant-negative human *KCNQ2* mutation as a transgene (Peters et al., 2005). None of these studies, however, satisfy the need to develop an animal model with a complete absence of Kv7.2 expression. In addition, previous studies of heterozygous *Kcnq2*-knockout mice found that reduction of Kv7.2 expression have a decreased seizure threshold (Watanabe et al., 2000, Yang et al., 2003).

In the present study, we characterize mice in which *Kcnq2* has been deleted in all somatic sensory neurons, using the Cre-Lox system (Sauer and Henderson, 1988, Nagy, 2000). These mice are viable and their myelinated sensory axons have a normal ultrastructure and normal molecular composition of nodes (including Kv7.3), paranodes, and juxtaparanodes. *Kcnq2* mutant mice showed signs of thermal hyperalgesia and mechanical allodynia, and *Kcnq2*-null dorsal root ganglia (DRG) neurons showed increased excitability and reduced spike-frequency adaptation. Taken together, our results suggest that Kv7.2 regulates neuronal excitability, and a reduction of Kv7.2 expression could lead to altered nociception.

MATERIALS AND METHODS

All procedures involving rodents were approved by the Institutional Animal Care and Use Committee of the University of Pennsylvania. Except when specified, all chemicals were from Sigma.

Generation of conditional *Kcnq2*-null mice

A floxed allele of *Kcnq2* was designed (Figure 1A) to delete exons 3–5; these correspond to 463 bp (amino acid 130–285) of mouse *Kcnq2* cDNA (GeneBank AF490773; (Wen and Levitan, 2002)); this deletion has been shown to result in a functional null allele (Watanabe et al., 2000). Two DNA fragments (*EcoRV* and *XhoI*) were cloned from R1 ES-cells genomic DNA, and used to generate a targeting vector in which a FRT-neomycin- FRT-loxP

cassette was inserted into a unique *XhoI* site, with flanking 7kb and 3.4kb of homologous genomic DNA. An additional loxP site was inserted in an *EcoRI* site just upstream of the third exon. The targeting vector also includes a diphtheria toxin (DT) gene as a negative selection marker against ES clones which have randomly integrated the targeting vector. The linearized targeting vector was electroporated into R1 embryonic stem cells, selected with G418, and colonies were picked and analyzed by Southern blot. Positive ES cells were used to generate chimeras and then heterozygous mice (Gollan et al., 2003, Poliak et al., 2003). The presence of the targeted locus was confirmed by polymerase chain reaction (PCR) analysis of tail DNA. Heterozygous mice were first mated with mice carrying the FLP recombinase (129S4/SvJaeSor-Gt(Rosa)26Sortm1(FLP1)Dym/J, RRID:IMSR_JAX:003946; (Farley et al., 2000)), to remove the *Neo* cassette and to create a floxed *Kcnq2* allele.

We crossed mice expressing *Pax3-Cre* (RRID:IMSR_JAX:005549) with mice carrying the *Rosa26* reporter gene (RRID:IMSR_JAX:003504). We examined X-gal expression in the brains, spinal cords, and lumbar DRG of 3 *Pax3-Cre*-positive//*Rosa26* mice and 3 *Pax3-Cre*-negative//*Rosa26* littermates, all 1-month-old, as previously described (Feltri et al., 1992, Arroyo et al., 1998). The mice were perfused with 0.5% glutaraldehyde in 0.1M phosphate buffer (PB; pH 7.4), their cerebra, cerebelli, and spinal cords were dissected and cut into slabs with a razor blade, and the resulting sections along with L4 and L5 DRG were fixed for 3 hours at 4°C, stained in X-Gal (Roche Diagnostic) at 37°C for 24–48 hours, rinsed in 0.1M PB, then re-fixed in 3% glutaraldehyde (in 0.1M PB) at 4°C overnight. The samples were photographed using a Nikon Coolpix 5000 camera mounted on a Leica MZ16 FA stereomicroscope. The DRG were osmicated, dehydrated, and embedded with the Embed 812 kit (EMS). Semi-thin sections (1 µm thick) were photographed using a cooled Hamamatsu camera mounted on Leica DMR light microscope.

Kcnq2 was deleted in sensory axons by crossing with mice that were heterozygous for both the floxed *Kcnq2* allele and *Pax3-Cre* (*Kcnq2*^{fl/+}//*Pax3-Cre*). We chose this approach because it generates relatively more mice of the desired genotype, and homozygous *Pax3-Cre* mice fail to develop past E18.5 according to Jackson Laboratory. All offspring were genotyped by PCR. Tail DNA was digested with DirectPCR (Viagen) and proteinase K overnight at 55°C, heated to 85°C for 45 minutes to denature proteinase K, then PCR were then performed with REDTaq ReadyMix PCR reaction mix following manufacturer's protocol in a Bio-Rad DNA Engine Peltier thermal cycler. Three primers were used together: KCNQ2A (GGGGCAGTTGTCTAACCCTC), KCNQ2C (TATGTGGTGTCTCCCCAGAAG), and KCNQ2E (GGGAGGCTCTAGTGTCTAGTG); see Figure 1B. After amplification, samples were separated in 1.5% agarose (GeneMate) gel in 1xTBS at 125V for 1 hour. To detect the presence of *Pax3-Cre* in the control littermates (*Kcnq2*^{+/+}), three primers were used together: oIMR6977 (CTGCACTCAAGGGACTCCTC), oIMR6978 (GTGAAGGCGAGACGAAAAAG), and oIMR9074 (AGGCAAATTTTGGTGACGG), following instruction provided by Jackson Labs.

Anatomical studies

One-month-old *Kcnq2*-mutant mice of either sex (*Kcnq2^{fl/fl}//Pax3-Cre*; n=3) and their control littermates (*Kcnq2^{+/+}//Pax3-Cre*; n=3) were anesthetized with ketamine/xylazine mix and killed by decapitation. Sciatic nerves, DRG (from L4–L6 spinal levels) with the ventral and dorsal roots attached, and femoral nerves (motor and sensory branches separately) were dissected, and quickly embedded in OCT mounting media cooled in an acetone/dry ice slurry. The sciatic and femoral nerve fibers were teased apart with fine needles, mounted on SuperFrost Plus glass slides (Fisher Scientific, Pittsburgh, PA), dried overnight, and stored at -20°C . Ten μm thick cryostat sections were thaw-mounted onto SuperFrost slides, and stored at -20°C . Teased fibers and OCT sections were immersed in -20°C acetone for 10 minutes, rinsed in Tris-buffered saline (TBS; pH 7.4), blocked at room temperature for one hour in TBS containing 5% fish skin gelatin and 0.5% Triton X-100, and incubated overnight at 4°C with various combination of primary antibodies diluted in blocking solution. The slides were washed with TBS, incubated with the appropriate FITC- and TRITC-conjugated donkey cross-affinity-purified secondary antibodies (Jackson ImmunoResearch, 1:200) at room temperature for one hour, washed with TBS, counterstained with 4',6-diamidino-2-phenylindole dihydrochloride (DAPI; Invitrogen), mounted with Vectashield (Vector Laboratories), then examined with a Leica DMR light microscope with a cooled Hamamatsu camera under the control of Openlab software (PerkinElmer, <http://www.perkinelmer.com/pages/020/cellularimaging/products/openlab.shtml>, RRID:rid_000096).

To analyze the structure of myelinated axons, one-month-old *Kcnq2* mutant mice of either sex (*Kcnq2^{fl/fl}//Pax3-Cre*; n=3) and littermate controls (*Kcnq2^{+/+}//Pax3-Cre*; n=3) were perfused with 0.9% NaCl, followed by fixative (2% glutaraldehyde/2% paraformaldehyde in 0.1M PB) for 10 minutes, then the femoral sensory and motor branches as well as the L4 and L5 DRG were dissected, fixed for 4 hours at 4°C , osmicated, dehydrated, and embedded with the Embed 812 kit. Semi-thin (1 μm) sections were stained with toluidine blue and examined as above; thin (80 nm thick) sections were stained with lead citrate and examined with a JEM-1010 transmission electron microscope (JEOL USA) outfitted with a digital camera.

Antibody Characterization

Please see Table 1 for a summary of all primary antibodies used. The KCNQ2N antiserum (Dr. Edward Cooper, Cat# KCNQ2N, RRID:AB_2312342) stained nodes and AISs of teased nerve fibers in an identical pattern as previously shown (Devaux et al., 2004; Pan et al., 2006), and is directed against residues 13–37 (GEKKLKVGFGVGLDPGAPDSTRDC) from the intracellular terminal region of human KCNQ2 (Cooper et al. 2001). The KCNQ3C antiserum (Dr. Edward Cooper, Cat# KCNQ3C, RRID:AB_2312343), which detects residues 578–604 (STPKHKKSQKGSFTFPSQQSPRNEPYc) of human KCNQ3 (Pan et al. 2006), also stained nodes in an identical pattern as previously described (Pan et al., 2006). To ascertain the specificity of the Kv7.2 antisera, HeLa cells were transiently transfected with the cDNA encoding human Kv7.2 or Kv7.3, (kindly provided by Dr. Edward Cooper) using the Lipofectamine 2000 (Invitrogen) method as previously described (Rasmussen et al., 2007), and immunostained 1 day post-transfection with the Kv7.2 and

Kv7.3 antisera. The KCNQ2N antiserum positively stained HeLa cells that were transiently transfected with the Kv7.2 cDNA but did not stain HeLa cells transfected with Kv7.3 cDNA; the KCNQ3C positively stained transfected HeLa cells expressing Kv7.3, but did not label transfected cells expressing Kv7.2 (data not shown).

The sequences of immunogen targeted by the KCNQ5 antiserum (Millipore (CHEMICON/Upstate/Linco), Cat# AB5599, RRID:AB_210806) is a stretch of 88 amino acids from the N-terminal sequence of KCNQ5 (M1–R88:GAAGLWVRSGAAAAAGAGGGRPGSGMKDVESGRGRVLLNSAAARGDGLLLLGTRAAALGGGGGLRESRRGKQGARMSLLGK), and were ascertained from both the manufacturer and from Caminos et al. (2007). The antiserum was able to positively label HeLa cells transfected by the Kv7.5 cDNA (kindly provided by Dr. Thomas Jentsch) but did not label HeLa cells transfected by the Kv7.2 cDNA (data not shown).

The mouse panNav monoclonal antiserum (Sigma-Aldrich, Cat# S8809, RRID:AB_477552) targets multiple voltage-gated Na⁺ channels, and detects immunogen (CTEEQKKYYNAMKKLGSKK) from the intracellular III–IV loop of Na⁺ channels (Rasband et al., 1999). Its specificity was confirmed by Western blotting of rat brain extracts recognizing a 260 kDa protein (manufacturer's technical information), and has previously been shown to specifically stain the AISs and nodes of a wide range of nervous tissues (Rasband et al., 1999; Devaux et al., 2004; Pan et al., 2006); in our stainings it recognized the nodes of both rat and mouse sciatic nerves, as expected.

The mouse monoclonal Kv1.1 antiserum (NeuroMab, Cat# K20/78, RRID:AB_2312366) was prepared against a synthetic peptide representing amino acids 458–476 (CEEDMNNSIAHYRQANIRTG) of rat Kv1.1 (Rasband et al., 1998). Our Western blot analysis showed a strong band at around 85 kDa, and a weaker band at around 65 kDa, identical with the information provided by the manufacturer (data not shown). In our staining it labeled the juxtaparanodal component of myelinated axons, as had been demonstrated previously (Rasband et al., 2001).

The mouse monoclonal contactin-associated protein (Caspr) antiserum (Peles et al., 1997, Cat# contactin-associated protein (Caspr), RRID:AB_2311776) was generated by Dr. Poliak by immunizing mice with a fusion protein composed of the extracellular domain of human Caspr (amino acids 1–1282; Peles et al., 1997) fused to the Fc region of human immunoglobulin G, and stained a single band of 180kD molecular weight on Western blot (Poliak et al., 1999); in our staining it recognized the paranodes of mouse sciatic nerve, identical to previous descriptions (Poliak et al., 1999, Poliak et al., 2003, Ogawa et al., 2010).

Behavior Testing

Three-month-old *Kcnq2*-mutant (*Kcnq2*^{fl/fl}/*Pax3-Cre*; 5 males and 4 females) and their littermates (*Kcnq2*^{+/+}/*Pax3-Cre*; 5 males and 4 females), derived from 3 litters were studied. Mechanical allodynia was measured using Chaplan's up-down threshold method (Chaplan et al., 1994, Hubbard and Winkelstein, 2005, Lee et al., 2008, Quinn et al., 2010). Mice were confined in plexiglass enclosure placed on a wire-mesh platform and allowed to

acclimate for at least 30 minutes before each test. Three rounds of testing were performed over three consecutive days, with each round comprised of five stimulations of either the right or the left mid-plantar hindpaw in a random order, with a series of ascending von Frey filament strength (0.4 g, 0.6 g, 1.0 g, 1.4 g, 2.0 g; Stoelting Co.) held perpendicular for 6–8 seconds against the skin with enough force to cause slightly buckling. A positive response is recorded for sharp paw withdrawal or if immediate flinching was observed upon removal of filament. Ambulation was considered an ambiguous response and in such case the stimulus was repeated. If two consecutive filament strengths elicited a withdrawal response, the lower of the two filament strengths was recorded as the threshold. Any mouse that failed to display a response with the highest filament strength was recorded as having a threshold of 2.0 g. Testing in the opposite direction (descending filament strength) was also performed during each round to confirm the withdrawal threshold. The average threshold of the three rounds was recorded for each mouse.

The thermal nociceptive response was assessed using a paw thermal stimulator system (UARDG, University of California San Diego; otherwise referred to as Hargreaves' apparatus) as previously described (Hargreaves et al., 1988, Dirig et al., 1997). Four rounds were performed over four consecutive days. Briefly, the mice were allowed to acclimate on the glass plate of the apparatus (maintained at 30°C) for at least 30 minutes, then either the left or the right mid-plantar hindpaw was randomly heated with a thermocouple set at 5 amperes. A timer is automatically started with the thermal source, and response latency is defined as the time required for the hindpaw to show an abrupt withdrawal (maximum 20 seconds). Paw withdrawal is automatically detected by an array of photodiode motion sensors mounted on the stimulus tower that stops the timer and terminates the stimulus. Stimulus current is monitored continuously. Six trials were performed during each round, with a minimum of 5 minutes between each trial to allow the hindpaws to return to normothermic baseline (Dirig et al., 1997). The average threshold of the four rounds was recorded for each mouse.

Motor function was measured with a rotarod apparatus (Ugo Basile, Stoelting Co.) as previously described (Wood et al., 2005, Oliveira et al., 2006). Briefly, the rotarod has a 3-cm diameter rotating rod raised 16 cm above a platform and divided into five sections for testing multiple mice simultaneously. Mice were acclimated on the first day by allowing them to run on the rotarod with the slowest rotation speed. Three rounds of testing were performed during the three subsequent days, with three trials during each round. For each trial, mice were placed on the rotarod, and the rotation speed was gradually increased from 4 to 40 rpm over the course of 5 minutes. Each trial ended when mice fell off (maximum 300 seconds), and the latency to fall was recorded for each trial. Test was considered valid if mice ran forward on the rotarod for at least 10 seconds. Mice were given 1-hour rest between each trial. The average time to fall of each mouse was used as the outcome.

Whole cell patch recording

DRG neurons were dissected and cultured from three-month-old *Kcnq2* mutant mice of either sex (*Kcnq2*^{fl/fl}//*Pax3-Cre*; n=5) and their littermate controls (*Kcnq2*^{+/+}//*Pax3-Cre*; n=5) from 2 litters, as previously described (Malin et al., 2007). Lumbar DRGs were rapidly

removed, transferred to ice-cold $\text{Ca}^{2+}/\text{Mg}^{2+}$ -free HBSS (GIBCO), then incubated first in papain solution (60U papain; Worthington), 3 μl saturated NaHCO_3 , 1 mg L-cysteine free base, 1.5 ml $\text{Ca}^{2+}/\text{Mg}^{2+}$ -free HBSS (GIBCO) for 10 minutes at 37°C, then in collagenase/dispase solution (0.1 U/ml collagenase 0.8 U/ml dispase (Roche), 3 ml $\text{Ca}^{2+}/\text{Mg}^{2+}$ -free HBSS) for 1 hour at 37°C. Neurons were dissociated by trituration using fire-polished glass Pasteur pipettes, suspended in F12 medium (GIBCO) containing 10% FCS (Invitrogen) and 1% penicillin/streptomycin (Invitrogen), then plated onto laminin/poly-D-lysine (Beckton Dickinson) coated coverslips (Fisher). DRG neurons adhered to coverslips and were maintained in culture for 12–48 hours after plating at 37°C prior to recording.

Whole cell patch-clamp techniques (Hamill et al., 1981, Lancaster et al., 2001) were employed with an Axopatch 200B amplifier and Axon Instruments PCLAMP 9 software (pClamp, <http://www.moleculardevices.com/products/software/pclamp.html>, RRID:rid_000085). Patch pipettes (1–4 M Ω) were fabricated from glass capillaries (MTW150F-4, World Precision Instruments). Pipettes were filled with a variant of a solution described previously (Ikeda et al., 1986), with a composition of (in mM) 140 KCl, 2 MgATP, 10 N-[2-hydroxyethyl] piperazine-N9-[2-ethanesulfonic acid] (HEPES), 11 ethylene glycolbis(b-aminoethyl ether)-N,N,N9,N9-tetraacetic acid (EGTA), and 2 CaCl_2 ; titrated to pH 7.3 with KOH, and to 314 mOsm with sucrose. Pipette voltage offset was neutralized prior to the formation of a giga-seal. Membrane input resistance (R_{in}), series resistance (R_{s}), and capacitance (C_{m}) were determined from current transients elicited by 5 mV depolarizing steps from a holding potential of –60 mV, delivered using the Membrane Test application of PCLAMP9. Criteria for cell inclusion in the study were as follows: R_{s} 10 M Ω , R_{in} 100 M Ω , and stable recording during the entire experiment. Cover slips were superfused (2–4 ml/min) continuously during recording at 34–36°C extracellular solution (composition in mM: 10 glucose, 140 NaCl, 3 KCl, 0.6 MgCl_2 , 2.5 CaCl_2 , 10 HEPES, titrated to pH 7.4 with TRIS base, to 325 mOsm with sucrose if needed). Tetraethylammonium (TEA) was fully dissolved in the extracellular solution prior to application; 3 mM TEA was applied for at least 2 minutes before determining its effects on membrane properties. To prevent bias in cell selection or analysis, the electrophysiologist was blinded to genotypes of the cells (control versus *Kcnq2*-null) during the experiments and measurements of cell properties. A total of 25 neurons from mutant mice, and 25 neurons from control littermates, were recorded and analyzed.

Statistical Tests

Other than the following exceptions, all comparisons between two samples were explored using unpaired two-samples Student's t-test, with equal variance first established with the equality of variance test. All statistical tests were performed with SAS 9.2 (Statistical Analysis System, http://www.sas.com/en_us/software/sas9.html, RRID:nif-0000-31484), with all data given as mean $\pm$ standard error of mean (SEM). Comparison of Kv7.2- and Kv7.3-positive nodes in different peripheral nerves was tested with two-way ANOVA; motor function results over the course of nine sessions were analyzed with repeated measures ANOVA; two-way repeated measures ANOVA was used to compare mutant and control neurons before and after 3 mM TEA across multiple rheobases, followed by the

Tukey test to identify specific significant differences. The minimum *a priori* criteria for statistical significance is $p < 0.05$.

RESULTS

Generation of conditional *Kcnq2*-null mice

We generated sensory neurons lacking *Kcnq2* expression using the Cre-Lox system (Sauer and Henderson, 1988, Nagy, 2000). Mice containing a floxed allele of *Kcnq2* (Fig. 1A) were crossed with mice expressing a *Pax3-Cre* transgene, which is expressed by cells derived from the neural crest (Li et al., 2000). To determine whether *Pax3-Cre* deleted exons 3–5 of *Kcnq2*, we examined three *Kcnq2^{fl/fl}//Pax3-Cre* mice by amplifying genomic DNA with PCR and appropriate primers (Fig. 1A–B). As shown in Figure 1C, while both the floxed and the recombined *Kcnq2* alleles were detected in the DRG and trigeminal ganglia, only the floxed allele was detected in the liver, indicating that the *Kcnq2* allele was being selectively deleted in neural-crest-derived tissues.

We examined the expression of the *Pax3-Cre* by crossing the mice with *Rosa26 lacZ* reporter mice, in which Cre removes a floxed stop cassette so that *lacZ* is expressed (Soriano, 1999). Using litters that contained mice of the informative genotypes, we stained the brain, spinal cord, and DRG for beta-galactoside activity with X-gal as the chromogen. None of the samples taken from the control animals showed X-gal staining, whereas in *Pax3-Cre//Rosa26* mice, there was robust staining in much of the cerebellum, regions of the cerebrum (including the cortex and the thalamus), the dorsal part of the spinal cord (including the dorsal roots), and the DRG (data not shown). The ventral part of the spinal cord and the ventral roots were unstained, suggesting that motor neurons do not express *Pax3-Cre*. Semi-thin sections demonstrated that virtually all DRG neurons from the *Pax3-Cre*-positive mice showed X-gal staining (Fig. 2), indicating that DRG neurons and/or their embryonic precursors express *Pax3-Cre*.

Primary sensory neurons in conditional *Kcnq2*-null mice lacked Kv7.2

To determine whether *Pax3-Cre* resulted in the loss of Kv7.2 expression in sensory neurons, we examined three mutants (*Kcnq2^{fl/fl}//Pax3-Cre*) and three control littermates (*Kcnq2^{+/+}//Pax3-Cre*). We immunostained unfixed teased nerve fibers from femoral sensory and motor branches, and from dorsal and ventral roots, as well as sections of the lumbar DRG, for Kv7.2 or Kv7.3, combined with a mouse monoclonal antibody against voltage-gated Na⁺ channels (panNav). As shown in Figure 3, in the control mice, all nodes of Ranvier of both the motor and sensory branches of the femoral nerve were Kv7.2- and panNav-positive, agreeing with our previously published results (Devaux et al., 2004, Pan et al., 2006). In mutant animals, all nodes were panNav-positive, but none of the nodes of the femoral sensory branch, and only some nodes of the femoral motor branch, were Kv7.2-positive. The lack of nodal staining of the *Kcnq2*-null sensory axons supports for the specificity of the Kv7.2 antiserum, which also selectively labeled HeLa cells transiently transfected to express human Kv7.2, but did not label transfected cells expressing Kv7.3 (data not shown). Similarly, none of the nodes in the dorsal roots (which are purely sensory), and all of the

nodes in the ventral roots (which are purely motor), were Kv7.2-positive (Fig 4). Another Kv7.2 antiserum (provided by Dr. Jérôme Devaux) gave similar results (data not shown).

We also immunostained sections of lumbar DRG and trigeminal ganglia with the Kv7.2 antiserum. As shown in Figure 5, in control animals, the Kv7.2 antiserum predominately labeled larger diameter DRG neurons (most of which were panNav-negative), while the panNav antibody primarily labeled smaller diameter neurons. In contrast, in the mutants, we did not detect robust Kv7.2 staining in any neurons, while the panNav antibody retained its staining pattern. Notably, similar to the teased fibers, nodes in the *Kcnq2*-null DRG were Kv7.2-negative. We also found similar results in the trigeminal ganglia (data not shown), with a loss of Kv7.2-immunostaining in the neurons and nodes of mutant animals. Thus, we conclude that *Pax3-Cre* deleted *Kcnq2* from virtually all primary sensory neurons.

In vitro studies have shown that Kv7.2 and Kv7.3 can form heteromeric channels, and that Kv7.2 increases cell membrane expression of Kv7.3 (Wang et al., 1998, Selyanko et al., 2001, Shah et al., 2002, Maljevic et al., 2003, Gomez-Posada et al., 2010). Because Kv7.2 and Kv7.3 are co-localized at nodes *in vivo* (Pan et al., 2006, Schwarz et al., 2006), we investigated whether nodal expression of Kv7.3 requires Kv7.2. As shown in Figures 4 and 6, all nodes of mutant mice showed robust Kv7.3 immunostaining – the femoral motor and sensory branches, as well as the dorsal and ventral roots. Thus, nodal expression of Kv7.3 does not require Kv7.2.

To quantify these results, we calculated the proportion of Kv7.2- or Kv7.3-positive nodes in each of the four nerve branches (femoral sensory, femoral motor, ventral root, and dorsal root) out of all panNav-positive nodes. As shown in Figure 7, the proportion of Kv7.2-positive nodes in the dorsal root ($1.4 \pm 0.06\%$), femoral sensory branch ($0.48 \pm 0.48\%$), and femoral motor branch ($47 \pm 1.6\%$) of the mutant animals were significantly less than in control littermates ($99 \pm 0.55\%$, $98 \pm 0.53\%$, $96 \pm 0.95\%$ respectively). As expected, the proportion of Kv7.2-positive nodes in the ventral root is not different between the mutants ($99 \pm 0.60\%$) and the controls ($97 \pm 0.41\%$), and the proportions of Kv7.3-positive nodes in all four nerve branches was found to be nearly 100% in both mutant and control mice.

We examined the structure of *Kcnq2*-null myelinated axons. Teased fibers from mutant sciatic nerves (in which approximately 20% of myelinated axons are motor and the remaining are sensory; (Schmalbruch, 1986), were immunostained for Kv7.2 and either panNav, Caspr, or Kv1.1. As shown in Figure 8, the localization of selected nodal (panNav), paranodal (Caspr) and juxtaparanodal (Kv1.1) components were the same for myelinated axons from either Kv7.2-positive (motor axons) or Kv7.2-negative (sensory axons) nodes. The motor and sensory axons of the femoral nerves and dorsal roots of the mutant animals appeared normal in semi-thin cross-sections (data not shown). Longitudinal thin sections of femoral sensory nerves were visualized by electron microscopy. The paranodal loops, nodal microvilli, and nodal axolemma all appeared normal (data not shown).

Finally, because we had previously demonstrated that Remak fibers, but not nodes of Ranvier, express Kv7.5 (King and Scherer, 2012), we wished to determine whether this pattern of expression is retained in *Kcnq2*-mutant mice. As shown in figure 9, in both the

femoral sensory and femoral motor branches, Remak fibers of both control and mutant mice showed robust Kv7.5 immunostaining. Nodes were Kv7.5-negative, indicating that Kv7.3 forms homomeric channels in the *Kcnq2*-null axons.

Behavioral testing of mutant and control mice

We tested 9 mutants (*Kcnq2^{fl/fl}//Pax3-Cre*) and 9 control littermates (*Kcnq2^{+/+}//Pax3-Cre*) from three litters in a battery of tests. Mutant and control mice performed the rotarod test equally well (Fig. 10A), and showed similar performance improvement over the nine testing sessions across three consecutive days (data not shown). The average time to fall for all sessions was not statistically different between the mutants (163 ± 13 seconds) and controls ($170. \pm 6.1$ seconds) groups ($P = 0.62$). To determine whether mutant mice have altered thermal hyperalgesia, we measured the withdrawal latency from noxious thermal stimuli using the Hargreaves' test over four consecutive days. As shown in Figure 10B, the mutant mice on average had significantly shorter withdrawal latency (3.7 ± 0.16 seconds) than did controls (4.3 ± 0.16 seconds; $p < 0.05$). To determine whether mutant mice have altered mechanical allodynia, we measured pain withdrawal threshold using von Frey filaments over three consecutive days. As shown in Figure 10C, the mutants had a significantly lower average withdrawal threshold (0.65 ± 0.07 g) compared to the controls (0.93 ± 0.09 g; $p < 0.05$). Taken together, our behavior tests suggest that the loss of Kv7.2 expression leads to increased thermal hyperalgesia and mechanical allodynia.

Electrophysiological studies of DRG neurons

We examined the electrophysiological characteristics of acutely isolated DRG neurons from the lumbar DRG from 5 mutants (*Kcnq2^{fl/fl}//Pax3-Cre*) and 5 controls (*Kcnq2^{+/+}//Pax3-Cre*) from two litters, using whole cell-patch clamp recordings (Table 2). From each genotype, we recorded from 25 neurons of various sizes; the average cell size recorded from each genotype (as estimated from membrane capacitance) was not statistically different between the two groups (39 ± 3.3 pF versus 45 ± 4.1 pF, respectively). The initial resting membrane potential was also not statistically different between the two groups (-58 ± 2.5 mV versus -56 ± 1.6 mV, respectively).

To investigate the fast and slow afterhyperpolarization (AHP) properties, we elicited single action potential (AP) with brief (3 ms) depolarizing current steps, and measured the peak undershoot (most negative potential during the AHP), as shown schematically in Figure 11A. Prior to applying TEA, the average fast AHP peak undershoot of mutant DRG neurons was 3 mV more negative than control neurons (-86 ± 0.97 mV versus -83 ± 1.3 mV, respectively; $p < 0.05$). After applying 3 mM TEA, the fast AHP peak magnitudes were not significantly different between the mutant and control neurons (-84 ± 1.2 mV versus -82 ± 0.97 mV, respectively; $p = 0.24$) (Figure 11B). In contrast, prior to 3 mM TEA, the average slow AHP peak undershoot of mutant neurons was 6 mV more positive than that of control neurons (-68 ± 1.2 mV versus -74 ± 1.9 mV, respectively; $p < 0.01$). After applying 3 mM TEA, the slow AHP peak undershoot of control neurons became 8 mV more positive (-66 ± 1.2 mV; $p < 0.01$), while the slow AHP peak undershoot of *Kcnq2*-null neurons did not change significantly (-67 ± 1.6 mV; Fig. 11C). Because 3 mM TEA should selectively block primarily the Kv7.2 homomers and Kv7.2/Kv7.3 heteromers and not Kv7.3 homomers

(Wang et al., 1998, Hadley et al., 2000, Lerche et al., 2000, Hadley et al., 2003), these results indicate that inhibiting Kv7.2 contributes to the decreased slow AHP peak undershoot observed in the mutant DRG neurons.

We examined the AP firing patterns and spike-frequency adaptation by injecting 500 ms depolarizing current steps into the DRG neurons. First, the threshold current (rheobase) for each neuron was determined using incremental (100 pA) 500 ms current steps (starting from 0.1 nA). Then we recorded the number of APs evoked by 500 ms depolarizing current steps of 1x, 2x, and 3x rheobase. To determine the responsiveness of DRG neurons to absolute (as opposed to relative threshold as described above) depolarizing stimuli, we also injected a series of 500 ms current steps from 0.4 nA to 3.6 nA (in increments of 0.4 nA). As shown in Figure 12, the *Kcnq2*-null neurons produced significantly more APs and exhibited less spike-frequency adaptation than did control neurons, and 3 mM TEA significantly increased the number of APs and decreased spike-frequency adaptation in control neurons, but not in *Kcnq2*-null neurons (numbers of AP from each group - control neurons: 1x rheobase 2.1, 2x rheobase 5.8, 3x rheobase 8.04, absolute 15.6; *Kcnq2*-null neurons: 1x rheobase 2.9, 2x rheobase 11.2, 3x rheobase 17.44, absolute 24.4; two-way repeated ANOVA). In addition, 3 mM TEA decreased the rheobase of control neurons by 49% (Table 1; 0.39 ± 0.07 nA from 0.77 ± 0.23 nA; $p < 0.05$), which is also statistically different than mutant neurons after TEA (0.88 ± 0.19 nA; $p < 0.05$). In contrast, the rheobase of the mutant neurons was not affected by TEA (0.88 ± 0.19 nA from 0.81 ± 0.16 nA; $p = 0.78$). In summary, mutant DRG neurons displayed increased excitability and decreased spike-frequency adaptation, and this hyperexcitability could be replicated in control neurons by 3 mM TEA.

DISCUSSION

Deleting *Kcnq2* enabled us to investigate directly the role of Kv7.2 in sensory neurons. As expected, deleting Kv7.2 had no discernible effect on the structure of myelinated axons, or on the motor performance of the mutant animals. There were modest effects on both acute thermal and mechanical nociceptive behaviors, and on the electrophysiological properties of sensory neurons. Without a direct experimental test, we cannot exclude the possibility that the lack of KCNQ2 in developing neurons contributes to the observed phenotype or that compensation may have diminished the effects of deleting *Kcnq2*.

Kv7.3 nodal expression does not depend on Kv7.2

Our data confirm that Kv7.2 and Kv7.3 are found at nodes (Devaux et al., 2004, Pan et al., 2006, Schwarz et al., 2006), although we document an even greater extent of their colocalization. This finding suggests that Kv7.2/Kv7.3 heteromers are the main Kv7 channels at PNS nodes. The localization of Kv7.3 at *Kcnq2*-null nodes demonstrates that Kv7.3 surface expression does not require Kv7.2, in contrast to previous in vitro studies (Schwake et al., 2000, Gomez-Posada et al., 2010). Because both Kv7.2 and Kv7.3 contain an ankyrin-G binding domain (Pan et al., 2006), it is reasonable to expect that Kv7.3 could be selectively retained at nodes and AIS. Indeed, the deletion of the ankyrin-G binding motif in Kv7.2 alone does not alter the AIS localization of Kv7.2/Kv7.3 heteromers (Rasmussen et al., 2007). In addition, the normal molecular components of nodes (panNav), paranodes

(Caspr), and juxtaparanodes (Kv1.1), as well as ultrastructure of *Kcnq2*-null sensory myelinated axons, shows that these features do not depend on Kv7.2. Finally, we found that Kv7.5 expression in Remak bundles is maintained in *Kcnq2*-null nerves. While Kv7.5 and Kv7.3 can form heteromeric channels *in vitro* (Schroeder et al., 2000), we did not detect Kv7.5-immunoreactivity at *Kcnq2*-null nodes, suggesting that in our mutant animals, the nodal Kv7 channels are comprised of Kv7.3 homomers.

Kv7.2 contributes to the regulation of neuronal excitability

Kv7 channels activate at subthreshold potentials and do not become inactivated, thereby contributing to the regulation of neuronal excitability (Brown and Passmore, 2009). Kv7 blockers (linopridine or XE991), dominant-negative Kv7.2 mutants, or decreased Kv7.2 expression have all been shown to increase excitability (decreased spike-frequency adaptation and/or increased number of action potentials) of hippocampal neurons (Aiken et al., 1995, Yue and Yaari, 2004, Gu et al., 2005, Peters et al., 2005, Shah et al., 2008) and of somatic and visceral sensory neurons (Passmore et al., 2003, Rivera-Arconada and Lopez-Garcia, 2005, Wladyka and Kunze, 2006, Wladyka et al., 2008, Zheng, 2013). A Kv7 enhancer (retigabine) produces the opposite effects (Lerche et al., 2000, Brueggemann et al., 2007). In isolated rat peripheral nerves, retigabine slows axonal conduction, and these effects can be reversed by application of linopridine or TEA (Devaux et al., 2004). Schwarz et al. (2006) showed that XE991 both abolishes the slow accommodation to the depolarization and the post-depolarization undershoot of action potential at nodes, as well as increases repetitive firing and decreases spike-frequency adaptation in rat motor axons. Taken together, these results indicate that Kv7 channels regulate both neuronal and axonal activity.

Our analysis of *Kcnq2*-null DRG neurons confirmed and extended these prior works. By using TEA at a Kv7.2-specific concentration (Wang et al., 1998, Hadley et al., 2000, Lerche et al., 2000, Hadley et al., 2003), we showed that Kv7.2 contributes to the spike-frequency adaptation of sensory neurons. Because TEA did not further decrease spike-frequency adaptation in *Kcnq2*-null neurons, decreased Kv7.2 activity is the most parsimonious explanation for the increased excitability of the *Kcnq2*-null neurons.

Classically, AHP can be subdivided into three phases: fast (1–5 ms), medium (50–200 ms), and slow (500 ms to several seconds) AHP (Madison and Nicoll, 1984, Storm, 1990, Gu et al., 2005). Because the slow activation speed of the M channels (tens of milliseconds), they do not contribute materially to the fast AHP (Brown and Passmore, 2009), which is instead considered to be mediated by the Big Potassium (BK) family of potassium channels (Storm, 1990). Indeed, our *Kcnq2*-null neurons exhibited only slightly more negative fast AHP amplitude compared to control neurons, and the application of TEA did not statistically change fast AHP amplitudes in both mutant and control neurons. The negative TEA results indicate that Kv7.2 does not play an appreciable role in the fast AHP. While we cannot explain why the fast AHP amplitude of our *Kcnq2*-null neurons was slightly more negative compared to the control neurons, the fact that Kv7.2-specific concentration of TEA did not change the fast AHP amplitude of the control neurons suggest that this difference may not be due to the lack of Kv7.2 activity.

On the other hand, the identity of the channel(s) that mediates the slow AHP is still uncertain, but decreases in Kv7.2 or Kv7.5 activity has been shown to reduce slow AHP in mouse hippocampal neurons (Tzingounis and Nicoll, 2008, Tzingounis et al., 2010). Our *Kcnq2*-null DRG neurons also displayed a decrease in slow AHP amplitude compared to control neurons, and 3 mM TEA was able to reduce the slow AHP amplitude of control neurons, while having no effect on mutant neurons. While a previous study have shown that 10 μ M XE991 application in hippocampal had actually enhanced slow AHP, other experiments showed that muscarine, which suppresses Kv7 channels (Brown and Adams, 1980), was able to inhibit slow AHP in the supraoptic neurons (Ghamari-Langroudi and Bourque, 2004, Hu and Mooney, 2005), which has also been shown to express Kv7 (Zhang et al., 2009). Therefore, while the exact contribution of Kv7 channels to slow AHP remained uncertain, our results raise the possibility that Kv7.2 activity does contribute, at least in part, to slow AHP in DRG neurons.

Increased mechanical allodynia and thermal hyperalgesia in *Kcnq2*-null mice

The results regarding the role of Kv7 channels on mechanical allodynia and thermal hyperalgesia are inconsistent. In one study, retigabine increases the tail withdrawal threshold to noxious thermal stimuli at a dose-dependent manner (Dost et al., 2004), but because this effect was not reversed by the co-administration of linopridine, it may be due to non-Kv7-specific effect of retigabine. This finding also conflicts with another study using different methodology, in which retigabine did not affect the withdrawal response from noxious thermal stimuli (Blackburn-Munro and Jensen, 2003). Similarly, intraplantar injections of XE991 into the rat hindpaws did not induce thermal hyperalgesia or mechanical allodynia (Linley et al., 2008). The above studies shared the important limitation that these enhancers and blockers likely act on most or all Kv7 subunits, and potentially on other channels (for example GABA_A (Otto et al., 2002)). Unmyelinated axons express Kv7.5 (King and Scherer, 2012), which could also be the site of these pharmacological agents.

In our current study we sought to minimize these confounding factors by analyzing a type of myelinated sensory axons that lacks Kv7.2 - the A-delta fibers. While technical limitations do not allow us to completely differentiate A-delta fiber mediated nociception from C-fiber mediated nociception, previous studies suggest that withdrawal reflex behavior from both acute noxious thermal stimuli (Price and Dubner, 1977, Dubner and Bennett, 1983, Yeomans and Proudfit, 1996, Hargreaves et al., 1998, Cuellar et al., 2010) and punctate mechanical stimuli (Dubner and Bennett, 1983, Koltzenburg et al., 1993, Ziegler et al., 1999) are both primarily mediated by A-delta fibers. Specifically, Yeomans and Proudfit (1996) found that radiant heating of rat hindpaw at a high rate of 6.5°C/second (°C/s) for 6 seconds primarily evokes A-delta fiber response, while a low rate of 0.9°C/s for 20 seconds primarily activates C-fibers response; notably, the C-fiber response at either heating rate did not begin until 5–8 seconds after onset of heating. Because the thermocouple of our Hargreaves apparatus had been shown to raise rat hindpaw temperature from 30°C to 49°C in 5 seconds (3.8°C/s) (Dirig et al., 1997), and because the average withdrawal latency of both mutant and control animals in our Hargreaves' test were less than 5 seconds (3.7 s and 4.3 s, respectively), our results suggest that the increased thermal hyperalgesia behavior of our *Kcnq2*-null mutant animal was, at least in part, due to increased A-delta fiber activity.

Furthermore, Ji et al. (2007) reported that rats treated with spinal nerve ligation displayed decreased mechanical threshold of A-delta fibers, but not of C fibers, and simultaneously exhibited increased mechanical allodynia (as tested with von Frey filaments), suggesting that the increased mechanical allodynia of our *Kcnq2*-null animals may also be due to increase A-delta fiber activity. Lastly, Passmore et al. (2012) recently showed that M-current inhibition by XE991 (at concentration that is relatively selective against Kv7.2) enhanced the response of A-delta fibers, but not C fibers, to noxious heat stimulation. Taken together, the increased thermal hyperalgesia and mechanical allodynia exhibited by our *Kcnq2*-null mice suggest that the lack of Kv7.2 expression in the A-delta fibers may produce increased acute thermal and mechanical nociception. Because our rotarod test provided no evidence of different motor performance between our mutant and control animals, the increased nociceptive responses observed in the *Kcnq2*-null mice is unlikely to be a consequence of altered motor behavior. However, because our X-gal stainings indicate that *Pax3*-Cre expression may also be present in regions of the brain, the possibility exists that the loss of Kv7.2 expression within the central nervous system may also have played a role in the increased thermal hyperalgesia and mechanical allodynia exhibited by our *Kcnq2* mutant mice.

Taken together, our work raises the possibility that decreasing Kv7.2 activity can increase sensory neuronal excitability, and lead to increased perception of mechanical and acute thermal pain.

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Figure 1. Targeting strategy to generate an inducible deletion allele of *Kcnq2*

Panel A shows the genomic arrangement of the first seven exons of wild-type *Kcnq2*. The external probes *a* and *b* are indicated as horizontal green lines. The targeted allele contains two *loxP* sites (orange triangles facing down) before the third, and after the fifth exons. A cassette containing a *neo* gene under the herpes simplex *Tk* promoter (*PtkNeo*; yellow box), flanked by two FRT sites (purple triangles facing up) is inserted in a unique *XhoI* site in the intron between the fifth and the sixth exons. The targeting vector (not shown) is the same as the pictured targeted allele except that it includes a diphtheria toxin (DT) gene as a negative selection marker after the last *EcoRV* site. Mice carrying the floxed allele were obtained after the removal of the *PtkNeo* cassette by mating the targeted mice with a general FRT-deleter strain. The recombined allele lacks exons 3–5, is generated after Cre-mediated deletion. **Panel B** shows the expected sizes of the amplified DNA with the different primer pairs. **Panel C** shows the PCR results of DNA isolated from DRG, trigeminal ganglia, and

liver of *Kcnq2*-null mice and their control littermates, along with tail DNA from control and heterozygous floxed *Kcnq2* mice (*Kcnq2* fl/+). Samples were subjected to PCR with the indicated primers, and the reaction products were separated by gel electrophoresis. A ~270 bp band, corresponding to the predicted size of wild-type *Kcnq2* allele, was detected in both tail samples, as well as in all of the samples from the control littermates. A ~300 bp band, corresponding to the predicted size of the floxed *Kcnq2* allele, was detected in the heterozygous tail, as well as in all of the *Kcnq2*-null samples. A ~380 bp band, corresponding to the predicted size of the recombined *Kcnq2* allele, was detected in both the DRG and the trigeminal ganglia of the *Kcnq2*-null mice, but not in their liver.

Figure 2. Sensory neurons expressed *Pax3-Cre*

These are digital images of 1 μ m thick epoxy sections of DRG neurons from *Pax3-Cre* positive (n=3) and -negative (n=3) mice that also expressed the *Rosa26* reporter gene. The DRGs had been labeled with X-gal. Almost all of the *Pax3-Cre* positive neurons showed X-gal staining (two are indicated with arrows), whereas none of the neurons from control mice showed X-gal staining. Scale bar: 20 μ m.

Figure 3. Selective loss of *Kcnq2* from the nodes of sensory axons

These are digital images of unfixed teased fibers, double-labeled for Kv7.2 (red) and voltage-gated sodium channels (panNav; green). In the femoral sensory branch (columns A and B), Kv7.2 is found at all nodes of Ranvier (arrows) in control animals, but is not detected at any nodes (arrowheads) in *Kcnq2*-null animals. In the femoral motor branch (columns C and D), Kv7.2 is found at all nodes of Ranvier (arrows) in control animals, but is not detected at some nodes (arrowheads) in *Kcnq2*-null animals. Scale bar: 10 μ m.

Figure 4. Selective loss of *Kcnq2* from the nodes of sensory axons

These are digital images of unfixed mouse teased nerves, double-labeled for either Kv7.2 or Kv7.3 (red) and panNav (green). In the dorsal roots of mutant animals (columns A and B), Kv7.2 is not detected at any node (arrowheads), whereas Kv7.3 is found at every node (arrows). In the ventral roots of mutant mice (columns C and D), note that all nodes are Kv7.2- and Kv7.3-positive (arrows). Scale bar: 10 μ m.

Figure 5. Primary sensory neurons of *Kcnq2*-null mice do not express Kv7.2

These are images of unfixed sections of lumbar DRG, double-labeled with a rabbit antiserum against Kv7.2 (red) and a panNav monoclonal antibody (green). In control DRG (column A), the Kv7.2 antiserum labels large diameter neurons, one of which is indicated (arrow); these are relatively unlabeled by the panNav antibody. In contrast, in *Kcnq2*-null DRG (column B), the Kv7.2 antiserum does not label large neurons (arrow). In both control and *Kcnq2*-null DRG, the panNav monoclonal labels small neurons, two of which are indicated (arrowhead). In addition, all nodes of control neurons are doublelabeled by both Kv7.2 and panNav (some enclosed within a circle), while none of the nodes of mutant neurons are Kv7.2-positive. Scale bar: 20 μ m.

Figure 6. Kv7.3 is found at all nodes in *Kcnq2*-null mice

These are digital images of unfixed mouse teased nerves, double-labeled for Kv7.3 (red) and panNav (green). In the femoral sensory (columns A and B) and motor branches (columns C and D) of both mutant and control mice, all panNav positive nodes are also Kv7.3-positive (arrows). Scale bar: 10 μ m.

Figure 7. Selective loss of *Kcnq2*, but not *Kcnq3*, from the nodes of sensory axons

The bars represent average proportion of panNav-positive nodes that are Kv7.2- (panel A) or Kv7.3- (panel B) positive from the indicated source ($n=3$ for all samples). Note that ~0% of nodes in the dorsal root and femoral sensory branch, and 50% in the femoral motor branch, were Kv7.2-positive. Error bars represent SEM; * $p<0.05$; ** $p<0.01$ (two-way ANOVA).

Figure 8. The molecular components of nodes, paranodes and juxtaparanodes are maintained in *Kcnq2*-null sensory axons

These are digital images of teased fibers from unfixed mutant sciatic nerves, doublelabeled for Kv7.2 (red) and either panNav (column A), Caspr (column B), or Kv1.1 (column C); all green. The selected nodal (panNav), paranodal (Caspr) and juxtaparanodal (Kv1.1) components are the same for myelinated axons with Kv7.2- positive (motor axons) or Kv7.2-negative (sensory axons) nodes. Scale bar: 10 μ m.

Figure 9. Remak fiber expression of Kv7.5 is maintained in *Kcnq2* mutant mice

These are digital images of unfixed mouse teased nerves, double-labeled for Kv7.5 (red) and panNav (green). In the femoral sensory (columns A and B) and motor (columns C and D) branches of both mutant and wild-type mice, all panNav-positive Remak fibers are Kv7.5-positive (chevrons), while all panNav-positive nodes are Kv7.5-negative (arrowheads). Scale bar: 10 μ m.

Figure 10. Behavior testing of *Kcnq2*-null mice and control mice

Panel A: Mutant mice (n=9) and their control littermates (n=9) were subjected to three consecutive days of testing on a rotarod (three tests per day), with motor performance measured by the time it took the mice to fall off the rotarod. There was no statistical difference in the overall average time to fall between the mutant and control animals.

Panel B: Both mutant mice (n=9) and their control littermates (n=9) were subjected to four consecutive days (six trials per day) of thermal hyperalgesia measured by the withdrawal latency after thermal stimulation (on a Hargreaves' chamber). The overall average of all tests showed mutant animals have a statistically significant increase in thermal hyperalgesia.

Panel C: Both mutant mice (n=9) and control littermates (n=9) were subjected to three consecutive days of testing with a series of von Frey hair filaments (0.4 g, 0.6 g, 1.0 g, 1.4 g, 2.0 g; one up-and-down session per day). Mechanical allodynia determined by the lowest strength of hair filament capable of inducing a positive withdrawal response. The overall

average of all tests showed mutant animals have a statistically significant increase in mechanical allodynia.

For all three behavior tests, error bars represent SEM; n.s. = no significant difference; * $p < 0.05$ (unpaired two-sample Student's t-test with equal variance).

Figure 11. *Kcnq2*-null DRG neurons have a diminished slow afterhyperpolarization (AHP) that is not affected by 3 mM TEA

Panel A shows representative responses from a control DRG neuron given a brief (3 ms) depolarizing current step that generated a single action potential (AP), and a prolonged (500 ms) depolarizing current step that generated multiple APs, from which the magnitude of the fast and slow AHP peak undershoots, respectively, were measured.

Panel B shows that before 3 mM TEA, the average fast AHP peak undershoot of mutant neurons (n=25) was slightly more negative than that of control neurons (n=25). After applying 3 mM TEA, the fast AHP peak undershoot of both control and *Kcnq2*-null neurons did not change by a statistically significant amount.

Panel C shows that before 3 mM TEA, the average slow AHP peak undershoot of mutant neurons (n=25) was 6 mV more positive than that of control neurons (n=25). After applying 3 mM TEA for 2 minutes, the slow AHP peak undershoot of control neurons became 8 mV

more positive, while the slow AHP peak undershoot of *Kcnq2*-null neurons did not change by a statistically significant amount. * $p < 0.05$; ** $p < 0.01$ (2-way ANOVA).

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Figure 12. *Kcnq2*-null DRG neurons have decreased spike-frequency adaption that is not affected by 3 mM TEA

Single DRG neurons were injected with 500 ms depolarizing current steps of 1x, 2x, or 3x rheobase (A–F) from a holding level of between –50 to –60 mV before and after application of 3 mM TEA. Panels A–D show representative recordings of control (A&D) and mutant (B&E) neurons, before and after application of 3 mM TEA for 2 minutes. Panels C and F are summary diagrams showing the number of APs in mutant and control neurons, before and after applying TEA, respectively (* $p < 0.05$; two-ways repeated measures ANOVA). Notably, after TEA application, control DRG neurons also showed a decrease in spike frequency adaption, and generated statistically similar numbers of APs at each rheobase as compared to mutant (n.s. = no significant difference; two-ways repeated measures ANOVA). In contrast, TEA application did not reduce spike-frequency adaptation of the mutant DRG neurons. In addition, we also injected 500 ms incremental current at set current step from 0.4nA to 3.6nA (in increments of 0.4nA) in order to determine the responsiveness of DRG neurons to absolute (as opposed to relative threshold as described above) depolarizing stimuli, and the spike frequency adaption was also significantly reduced in mutant DRG neurons in comparison to control neurons before TEA application; after TEA application the control neurons generated statistically similar number of APs in response to absolute stimuli as compared to mutant neurons.

Table 1

List of Primary Antibodies Used

Name	Manufacturer	Dilution	Species	Type	Immunogen
KCNQ2N	A gift from Dr. Edward Cooper	1:200	Rabbit	Polyclonal	Human KCNQ2 N-terminus residues 13–37 (GEKKLVGFVGLDPGAPDSTRDC)
KCNQ3C	A gift from Dr. Edward Cooper	1:200	Rabbit	Polyclonal	Human KCNQ3 C-terminus residues 578–604 (STPKHKKSQKGSFTTPSQSPRNPY _C)
KCNQ5	Chemicon; AB5599	1:1000	Rabbit	Polyclonal	KCNQ5 N-terminus (M1–R88:GAAGLWVRSGAAAAAGAGGGRPGSGMKDVESGRGRVLLNSAAARGDGLLLLLCTRAAALGGGGGGLRESRRRKQKGARMSLLGK)
panNav	Sigma; clone K58/35 (S8809)	1:250	Mouse	Monoclonal	C ^T EEQKKYYNAMIKKLGSKK from the intracellular IIIIV loop of Na ⁺ channels
Kv1.1	NeuroMab; clone K20/78	1:250	Mouse	Monoclonal	Rat Kv1.1 amino acids 458–476 (CEEDMNNNSIAHYRQANIRTG)
Caspr	A gift from Dr. Eltor Peles; Clone 275	1:100	Mouse	Monoclonal	Contained within extracellular domain of human Caspr (amino acids 1–1282)

Table 2

Passive and active membrane properties of control and *Kcnq2*-null DRG neurons.

	control	<i>Kcnq2</i> -null
Passive properties		
C_m (pF)	39 ± 3.3	45 ± 4.1
Initial RMP (mV)	-58 ± 2.5	-56 ± 1.6
Pre-drug firing properties		
Fast AHP peak (mV)	-83 ± 1.3	$-86 \pm 0.97^*$
AP rheobase (nA)	0.77 ± 0.23	0.81 ± 0.16
1 times rheobase (#APs)	2.1 ± 0.38	2.9 ± 0.75
2 times rheobase (#APs)	5.8 ± 1.3	11 ± 2.8
3 times rheobase (#APs)	8.0 ± 1.3	$17. \pm 4.4^*$
Absolute stimulus (#APs)	16 ± 1.6	$25 \pm 5.2^*$
Slow AHP peak (mV)	-74 ± 1.9	$-68 \pm 1.2^{**}$
Post-drug firing properties		
Fast AHP peak (mV)	-82 ± 0.97	-84 ± 1.21
AP rheobase (nA)	0.39 ± 0.07	$0.88 \pm 0.19^*$
1 times rheobase (#APs)	4.0 ± 1.1	2.5 ± 0.66
2 times rheobase (#APs)	11 ± 2.0	9.2 ± 2.2
3 times rheobase (#APs)	13 ± 2.2	16 ± 2.9
Absolute stimulus (#APs)	21 ± 2.2	23 ± 3.9
Slow AHP peak (mV)	-66 ± 1.2	-67 ± 1.6

Twenty-five DRG neurons from five animals were recorded in the control and *Kcnq2*-null groups. Values are means $\pm$ SEM. C_m = membrane capacitance; RMP = resting membrane potential; AHP = afterhyperpolarization potential; fast AHP peak = AHP undershoot peak magnitude after single 3-ms stimulus; AP rheobase = either the minimum amount of current required to evoke a single AP, or 0.1 nA (whichever one was smallest); 1, 2, and 3 times rheobase = the number of APs fired by a DRG neuron during a 500-ms depolarizing current step of a magnitude 1, 2, or 3 times its rheobase, respectively; absolute = the maximum number of APs fired in response to a single 500-ms depolarizing current step of 0.1–0.9 nA magnitude (in 0.1-nA increments); slow AHP peak = AHP undershoot magnitude after 500-ms stimulus at 0.4nA. Values of C_m of the control DRG neurons were compared to mutant neurons using unpaired 2-sample Student's t-test with equal variance; AHP and AP rheobase were compared with 2-way ANOVA with Tukey test; numbers of APs were compared with two-way repeated measures ANOVA.

*
p<0.05;

**
p<0.01.