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. J. Comp. Neurol. 000:000–000, 2014.

INDEXING TERMS: KCNQ; M-current; dorsal root ganglion; nociceptors; Kv7

The Kv7 (KCNQ) family of K⁺ channels is made up of five members, Kv7.1–Kv7.5 (Jentsch, 2000; Delmas and Brown, 2005; Brown and Passmore, 2009). Many peripheral nervous system (PNS) and central nervous system (CNS) neurons express Kv7.2, Kv7.3, and K7.5; in addition, Kv7.1 and Kv7.4 have recently been found in certain neuronal populations (Goldman et al., 2009; Heidenreich et al., 2011). Kv7.1 and Kv7.4 are prominently expressed in the cochlea, and Kv7.1 is expressed in the heart (Brown and Passmore, 2009). Kv7 channels contribute to the normal resting membrane potential and also form the noninactivating 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 likely is 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 from 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 dominate-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 involves a decreased seizure threshold (Watanabe et al., 2000; Yang et al., 2003).

The present study characterizes 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 that 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 (St. Louis, MO).

Generation of conditional Kcnq2-null mice

A floxed allele of Kcnq2 was designed (see Fig. 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 Xhol) were cloned from R1 ES-cells genomic DNA and used to generate a targeting vector in which an FRT-neomycin-FRT-loxP cassette 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 homologous genomic DNA. An additional loxP site was inserted in a unique Xhol site, with flanking 7 kb and 3.4 kb of unique EcoRI sites. The targeting vector also includes a Cre recombinase expression 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 three Pax3-Cre-positive//Rosa26 mice and three Pax3-Cre-negative//Rosa26 littermates, all 1-month-old, as previously described (Feltir et al., 1992; Arroyo et al., 1998). The mice were perfused with 0.5% glutaraldehyde in 0.1 M phosphate buffer (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 Diagnostics, Indianapolis, IN) at 37°C for 24–48 hours, rinsed in 0.1 M PB, then refixed in 3% glutaraldehyde (in 0.1 M PB) at 4°C overnight. The samples were photographed with 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). Semithin sections (1 μm thick) were photographed with a cooled Hamamatsu camera mounted on a 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<sup>2<sup;++</sup>/pax3-Cre<sup>+</sup>}). 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 and heated to 85°C for 45 minutes to denature proteinase K, and then PCRs were then performed with REDTaq ReadyMix PCR mix, following the manufacturer’s protocol in a Bio-Rad DNA Engine Peltier thermal cycler. Three primers were used together: KCNQ2A (GGGGCAGTTGTCTAACCCTC), KCNQ2C (TAGTGTCAGTG; see Fig, 1B). After amplification, samples were separated in 1.5% agarose (GeneMate) gel in 1× TBS at 125 V for 1 hour. To detect the presence of Pax3-Cre in the control littermates (Kcnq2<sup>2<sup>++</sup>/pax3-Cre<sup>+/+</sup>}), three primers were used together: oIMR6977 (CTGCACTCAAGGGAGACG), and KCNQ2E (GGGGCAGTTGTCTAACCCTC), oIMR9074 (AGGGAAGCTTATGCCATG; see Fig, 1B). After amplification, samples were separated in 1.5% agarose (GeneMate) gel in 1× TBS at 125 V for 1 hour. To detect the presence of Pax3-Cre in the control littermates (Kcnq2<sup>2<sup>++</sup>/pax3-Cre<sup>+/+</sup>}), three primers were used together: oIMR6977 (CTGCACTCAAGGGAGACG), and KCNQ2E (GGGAGGCTTAGTGTCAGTG; see Fig, 1B). After amplification, samples were separated in 1.5% agarose (GeneMate) gel in 1× TBS at 125 V for 1 hour. To detect the presence of Pax3-Cre in the control littermates (Kcnq2<sup>2<sup>++</sup>/pax3-Cre<sup>+/+</sup>}), three primers were used together: oIMR6977 (CTGCACTCAAGGGAGACG), and KCNQ2E (GGGAGGCTTAGTGTCAGTG; see Fig, 1B). After amplification, samples were separated in 1.5% agarose (GeneMate) gel in 1× TBS at 125 V for 1 hour. To detect the presence of Pax3-Cre in the control littermates (Kcnq2<sup>2<sup>++</sup>/pax3-Cre<sup>+/+</sup>}), three primers were used together: oIMR6977 (CTGCACTCAAGGGAGACG), and KCNQ2E (GGGAGGCTTAGTGTCAGTG; see Fig, 1B). After amplification, samples were separated in 1.5% agarose (GeneMate) gel in 1× TBS at 125 V for 1 hour. To detect the presence of Pax3-Cre in the control littermates (Kcnq2<sup>2<sup>++</sup>/pax3-Cre<sup>+/+</sup>}), three primers were used together: oIMR6977 (CTGCACTCAAGGGAGACG), and KCNQ2E (GGGAGGCTTAGTGTCAGTG; see Fig, 1B). After amplification, samples were separated in 1.5% agarose (GeneMate) g...
Figure 1. Targeting strategy to generate an inducible deletion allele of *Kcnq2*. **A** shows the genomic arrangement of the first seven exons of wild-type *Kcnq2*. The external probes a and b are indicated as horizontal lines. The targeted allele contains two \( \text{loxP} \) sites (triangles facing downward) before the third and after the fifth exons. A cassette containing a neo gene under the herpes simplex \( \text{Tk} \) promoter (\( \text{PtkNeo} \); boxed), flanked by two FRT sites (triangles facing upward) 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 \( \text{PtkNeo} \) cassette by mating the targeted mice with a general FRT-deleter strain. The recombined allele lacks exons 3–5 and is generated after Cre-mediated deletion. **B** shows the expected sizes of the amplified DNA with the different primer pairs. **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 (\( \text{Kcnq2}^{fl/}\)). Samples were subjected to PCR with the indicated primers, and the reaction products were separated by gel electrophoresis. An \( \sim \)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. An \( \sim \)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. An \( \sim \)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.
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-micrometer-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 1 hour in TBS containing 5% fish skin gelatin and 0.5% Triton X-100, and incubated overnight at 4°C with various combinations 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, West Grove, PA; 1:200) at room temperature for 1 hour, washed with TBS, counterstained with 4,6-diamidino-2-phenylindole dihydrochloride (DAPI; Invitrogen, Carlsbad, CA), mounted with Vectashield (Vector Laboratories, Burlingame, CA), and 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.xhtml, RRID:rid_000096).

Antibody characterization

See Table 1 for a summary of all primary antibodies used. The KCNQ2N antiserum (Dr. Edward Cooper; catalog No. KCNQ2N, RRID:AB_2312342) stained nodes and AISs of teased nerve fibers in a pattern identical to that previously shown (Devaux et al., 2004; Pan et al., 2006) and is directed against residues 13–37 (GEKKLKVGFVGLDPGAPDSTRDC) from the intracellular terminal region of human KCNQ2 (Cooper et al. 2001). The KCNQ3C antiserum (Dr. Edward Cooper; catalog No. KCNQ3C, RRID:AB_2312343), which detects residues 578–604 (STPKHKKSQKGSAFTFPSQQSPRNEPYc) of human KCNQ3 (Pan et al. 2006), also stained nodes in a pattern identical to that 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 cDNA encoding human Kv7.2 or Kv7.3 (kindly provided by Dr. Edward Cooper) using the Lipofectamine 2000 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 cDNA encoding human Kv7.2 or Kv7.3 (kindly provided by Dr. Edward Cooper) using the Lipofectamine 2000 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 cDNA encoding human Kv7.2 or Kv7.3 (kindly provided by Dr. Edward Cooper) using the Lipofectamine 2000 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 cDNA encoding human Kv7.2 or Kv7.3 (kindly provided by Dr. Edward Cooper) using the Lipofectamine 2000 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 cDNA encoding human Kv7.2 or Kv7.3 (kindly provided by Dr. Edward Cooper) using the Lipofectamine 2000 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 cDNA encoding human Kv7.2 or Kv7.3 (kindly provided by Dr. Edward Cooper) using the Lipofectamine 2000 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 cDNA encoding human Kv7.2 or Kv7.3 (kindly provided by Dr. Edward Cooper) using the Lipofectamine 2000 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 cDNA encoding human Kv7.2 or Kv7.3 (kindly provided by Dr. Edward Cooper) using the Lipofectamine 2000 method as previously described (Rasmussen et al., 2007) and immunostained 1 day post-transfection with the Kv7.2 and Kv7.3 antisera.
The sequences of immunogen targeted by the KCNQ5 antiserum (Millipore Chemicon/Upstate/Linco; catalog No. AB5599, RRID:AB_210806) are a stretch of 88 amino acids from the N-terminal sequence of KCNQ5 (M1-R88:GAAGLWVRSGAAAAAGGGRPGSGMK DVESGRGVRVLNSAAARGDGLLLLCTRAAALGGGGGGLRES RRGKOGARMSLLGK) and were ascertained from both the manufacturer and from Caminos et al. (2007). The antiserum was able to label positively 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 monoclonal anti-serum (Sigma-Aldrich; catalog No. S8809, RRID:AB_477552) targets multiple voltage-gated Na\(^+\) channels and detects immunogen (CTEEQKYYNAMKKLGSKK) 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 stain specifically 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 sciatric nerves, as expected.

The mouse monoclonal anti-serum (NeuroMab; catalog No. K20/78, RRID:AB_2312366) was prepared against a synthetic peptide representing amino acids 458–476 (CEEDMNNLSAHYQANIRGT) of rat Kv1.1 (Rasband et al., 1998). Our Western blot analysis showed a strong band at about 85 kDa and a weaker band at about 65 kDa, identical to 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; catalog No. 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 180 kD molecular weight on Western blot (Poliak et al., 1999). In our staining it recognized the paranodes of mouse sciatric nerve, identical to previous descriptions (Poliak et al., 1999, 2003; Ogawa et al., 2010).

**Behavioral testing**

Three-month-old Kcnq2 mutant mice (Kcnq2\(^{+/\times}\)/Pax3-Cre; five males and four females) and their littermates (Kcnq2\(^{+/+}\)/Pax3-Cre; five males and four females), derived from three litters, were studied. Mechanical allodynia was measured using Chaplan’s up-and-down threshold method (Chaplan et al., 1994; Hubbard and Winkelstein, 2005; Lee et al., 2008; Quinn et al., 2010). Mice were confined in a 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 3 consecutive days, with each round comprising five stimulations of either the right or the left midplantar hindpaw in a random order, with a series of ascending von Frey filament strengths (0.4, 0.6, 1.0, 1.4, and 2.0 g; Stoelting Co., Wood Dale, IL) 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 cases 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 4 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 midplantar 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 3 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. The test was considered valid if mice ran forward on the rotarod for at least 10 seconds. Mice were given 1 hour of rest between each trial. The average time to fall for each mouse was used as the outcome.

Whole-cell patch recording

DRG neurons were dissected and cultured from 3-month-old Kcnq2 mutant mice of either sex (Kcnq2^+/-//Pax3-Cre; n = 5) and their littermate controls (Kcnq2^+/+//Pax3-Cre; n = 5) from two litters, as previously described (Malin et al., 2007). Lumen DRGs were rapidly removed, transferred to ice-cold Ca^{2+}/Mg^{2+}-free HBSS (Gibco, Grand Island, NY), then incubated for 1 hour at 37°C and then in collagenase/dispase solution (0.1 U/ml collagenase 0.8 U/ml dispase [Roche], 3 ml Ca^{2+}/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), then plated 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 Dickson)-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 (Molecular Devices, http://www.moleculardevices.com/products/software/pclamp.html, RRID:ribo_00085). Patch pipettes (1–4 MΩ) were fabricated from glass capillaries (MTW150F-4; World Precision Instruments, Sarasota, FL). Pipettes were filled with a variant of a solution described previously (Ikeda et al., 1986), with a composition (in mM) of 140 KCl, 2 MgATP, 10 Na^{2+}-[2-hydroxyethyl] piperazine-N,N'-[2-ethanesulfonic acid] (HEPES), 11 ethylene glycolbis[b-aminoethyl ether]-N,N,N',N'-tetraacetic acid (EGTA), and 2 CaCl2, titrated to pH 7.3 with KOH and to 314 mOsm with sucrose. Pipette voltage offset was neutralized prior to the formation of a gigaseal. Membrane input resistance (R_m), 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 MΩ, R_m ≥ 100 MΩ, and stable recording during the entire experiment. Coverslips were superfused (2–4 ml/minute) continuously during recording at 34–36°C extracellular solution (composition in mM: 10 glucose, 140 NaCl, 3 KCl, 0.6 MgCl2, 2.5 CaCl2, 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 vs. Kcnq2-null) during the experiments and measurements of cell properties. In total 25 neurons from mutant mice and 25 neurons from control littermates were recorded and analyzed.

Statistical analysis

Other than the following exceptions, all comparisons between two samples were explored by unpaired two-samples Student’s t-test, with equal variance first established with the equality of variance test. All statistical tests were performed in 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 ± standard error of the 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 by 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 Kcnq2fl/fl/Pax3-Cre mice by amplifying genomic DNA with PCR and appropriate primers (Fig. 1A,B). As shown in Figure 1C, although 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 β-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. Semithin 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 (Kcnq2fl/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 the 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 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 labeled predominantly larger diameter DRG neurons (most of which were panNav negative), whereas the panNav antibody primarily labeled smaller diameter neurons. In contrast, for the mutants, we did not detect robust Kv7.2 staining in any neurons, and the panNav antibody retained its staining pattern. Notably, similar to the teased fibers, nodes in the Kcnq2-null DRG were

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**Figure 2.** Sensory neurons expressed Pax3-Cre. Digital images of 1-μm-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 μm.
Figure 3. Selective loss of Kcnq2 from the nodes of sensory axons. Digital images of unfixed teased fibers, double labeled for Kv7.2 (magenta) 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 μm.

Figure 4. Selective loss of Kcnq2 from the nodes of sensory axons. Digital images of unfixed mouse teased nerves, double labeled for either Kv7.2 or Kv7.3 (magenta) 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 μm.
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 colocalized 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) among all panNav-positive nodes. As shown in Figure 7, the proportions of Kv7.2-positive nodes in the dorsal root (1.4% ± 0.06%), femoral sensory branch (0.48% ± 0.48%), and femoral motor branch (47% ± 1.6%) of the mutant animals were significantly less than in control littermates (99% ± 0.55%, 98% ± 0.53%, 96% ± 0.95%, respectively). As expected, the proportion of Kv7.2-positive nodes in the ventral root is not different between the mutants (99% ± 0.60%) and the controls (97% ± 0.41%), and the proportions of Kv7.3-positive nodes in all four nerve branches were 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 panNav, Caspr, or Kv1.1. As shown in Figure 8, the localizations 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 semithin 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 nine mutants (Kcnq2^fl/fl///Pax3-Cre) and nine 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 3 consecutive days (data not shown). The average time to fall for all sessions was not statistically different between mutant (163 ± 13 seconds) and controls (170.6 ± 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 4 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 3 consecutive days. As shown in Figure 10C, the mutants had a significantly lower average withdrawal threshold (0.65 ± 0.07 g) compared with 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 five mutants (Kcnq2<sup>fl/fl</sup> /Pax3-Cre) and five controls (Kcnq2<sup>+/+</sup> /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 vs. 45 ± 4.1 pF, respectively). The initial resting membrane potential was also not statistically different between the two groups (258 ± 2.5 mV vs. 256 ± 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 vs. −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 the control neurons (−84 ± 1.2 mV vs. −82 ± 0.97 mV, respectively; P = 0.24; Fig. 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 vs. −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 ($-66 \pm 1.2 \text{ mV}$; $P < 0.01$), and the slow AHP peak undershoot of Kcnq2-null neurons did not change significantly ($-67 \pm 1.6 \text{ mV}$; Fig. 11C). Because 3 mM TEA should selectively block primarily the Kv7.2 homomers and Kv7.2/Kv7.3 heteromers but not Kv7.3 homomers (Wang et al., 1998; Hadley et al., 2000, 2003; Lerche et al., 2000), 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 $1 \times$, $2 \times$, and $3 \times$ 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: $1 \times$ rheobase 2.1, $2 \times$ rheobase 5.8, $3 \times$ rheobase 8.04, absolute 15.6; Kcnq2-null neurons: $1 \times$ rheobase 2.9, $2 \times$ rheobase 11.2, $3 \times$ rheobase 17.44, absolute 24.4; two-way repeated-measures ANOVA). In addition, 3 mM TEA decreased the rheobase of control neurons by 49% (Table 2; 0.39 ± 0.07 nA from 0.77 ± 0.23 nA; $P < 0.05$), which is also statistically different from the value for 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
Figure 8. The molecular components of nodes, paranodes and juxtaparanodes are maintained in Kcnq2-null sensory axons. Digital images of teased fibers from unfixed mutant sciatic nerves, double labeled for Kv7.2 (magenta) and 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 μm.

Figure 9. Remak fiber expression of Kv7.5 is maintained in Kcnq2 mutant mice. Digital images of unfixed mouse teased nerves, double labeled for Kv7.5 (magenta) 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), whereas all panNav-positive nodes are Kv7.5 negative (arrowheads). Scale bar = 10 μm.
Behavioral testing of Kcnq2-null mice and control mice. A: Mutant mice (n = 9) and their control littermates (n = 9) were subjected to 3 consecutive days of testing on a rotarod (three tests per day), with motor performance measured by the time it took for the mice to fall off the rotarod. There was no statistical difference in the overall average time to fall between mutant and control animals. B: Both mutant mice (n = 9) and their control littermates (n = 9) were subjected to 4 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 that mutant animals have a statistically significant increase in thermal hyperalgesia. C: Both mutant mice (n = 9) and control littermates (n = 9) were subjected to 3 consecutive days of testing with a series of von Frey hair filaments (0.4, 0.6, 1.0, 1.4, and 2.0 g; one up-and-down session per day). Mechanical allodynia was determined by the lowest strength of hair filament capable of inducing a positive withdrawal response. The overall average of all tests showed that 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).

TABLE 2.

Passive and Active Membrane Properties of Control and Kcnq2-Null DRG Neurons

<table>
<thead>
<tr>
<th></th>
<th>Control</th>
<th>Kcnq2-null</th>
</tr>
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<tbody>
<tr>
<td>Passive properties</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cm (pF)</td>
<td>39 ± 3.3</td>
<td>45 ± 4.1</td>
</tr>
<tr>
<td>Initial RMP (mV)</td>
<td>−58 ± 2.5</td>
<td>−56 ± 1.6</td>
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<tr>
<td>Predrug firing properties</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Fast AHP peak (mV)</td>
<td>−83 ± 1.3</td>
<td>−86 ± 0.97</td>
</tr>
<tr>
<td>AP rheobase (nA)</td>
<td>0.77 ± 0.23</td>
<td>0.81 ± 0.16</td>
</tr>
<tr>
<td>1× Rheobase (No. APs)</td>
<td>2.1 ± 0.38</td>
<td>2.9 ± 0.75</td>
</tr>
<tr>
<td>2× Rheobase (No. APs)</td>
<td>5.8 ± 1.3</td>
<td>11 ± 2.8</td>
</tr>
<tr>
<td>3× Rheobase (No. APs)</td>
<td>8.0 ± 1.3</td>
<td>17 ± 4.42</td>
</tr>
<tr>
<td>Absolute stimulus (No. APs)</td>
<td>16 ± 1.6</td>
<td>25 ± 5.2</td>
</tr>
<tr>
<td>Slow AHP peak (mV)</td>
<td>−74 ± 1.2</td>
<td>−68 ± 1.2</td>
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<tr>
<td>Postdrug firing properties</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Fast AHP peak (mV)</td>
<td>−82 ± 0.97</td>
<td>−84 ± 1.21</td>
</tr>
<tr>
<td>AP rheobase (nA)</td>
<td>0.39 ± 0.07</td>
<td>0.88 ± 0.19</td>
</tr>
<tr>
<td>1× Rheobase (No. APs)</td>
<td>4.0 ± 1.1</td>
<td>2.5 ± 0.66</td>
</tr>
<tr>
<td>2× Rheobase (No. APs)</td>
<td>11 ± 2.0</td>
<td>9.2 ± 2.2</td>
</tr>
<tr>
<td>3× Rheobase (No. APs)</td>
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<td>Absolute stimulus (No. APs)</td>
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<td>23 ± 3.9</td>
</tr>
<tr>
<td>Slow AHP peak (mV)</td>
<td>−66 ± 1.2</td>
<td>−67 ± 1.6</td>
</tr>
</tbody>
</table>

1Twenty-five DRG neurons from five animals were recorded in the control and Kcnq2-null groups. Values are means ± SEM. Cm, membrane capacitance; RMP, resting membrane potential; AHP, afterhyperpolarization potential; fast AHP peak, AHP undershoot peak magnitude after a single 3-ms stimulus; AP rheobase, either the minimum amount of current required to evoke a single AP or 0.1 nA (whichever was smallest); 1×, 2×, and 3× rheobase, the number of APs fired by a DRG neuron during a 500-ms depolarizing current step of a magnitude 1×, 2×, or 3× 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 0.4 nA. Values of Cm of the control DRG neurons was compared with that mutant neurons using unpaired two-sample Student’s t-test with equal variance; AHP and AP rheobase were compared with two-way ANOVA with Tukey test; numbers of APs were compared with two-way repeated-measures ANOVA.

2P < 0.05.

3P > 0.01.
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, and 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 et al., 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 postdepolarization undershoot of action potential at nodes and 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 confirms and extends these prior works. By using TEA at a Kv7.2-specific concentration (Wang et al., 1998; Hadley et al., 2000, 2003; Lerche et al., 2000), we show 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). Indeed, our Kcnq2-null neurons exhibited only slightly more negative fast AHP amplitude compared with control neurons, and slow AHP amplitudes in either mutant or control neurons. The negative TEA results indicate that Kv7.2 does not play an appreciable role in the fast AHP. We cannot explain why the fast AHP amplitude of our Kcnq2-null neurons was slightly

Figure 11. Kcnq2-null DRG neurons have a diminished slow after-hyperpolarization (AHP) that is not affected by 3 mM TEA. 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. 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. 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, whereas the slow AHP peak undershoot of Kcnq2-null neurons did not change by a statistically significant amount. *P < 0.05, **P < 0.01 (two-way ANOVA).
more negative compared with the control neurons, but
the fact that Kv7.2-specific concentration of TEA did not
change the fast AHP amplitude of the control neurons
suggests that this difference might 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 unclear, but decreases in
Kv7.2 or Kv7.5 activity have 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 with mutant (n.s., no significant
difference; two-way 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 steps
from 0.4 to 3.6 nA (in increments of 0.4 nA) to determine the responsiveness of DRG neurons to absolute (as opposed to relative thresh-
hold, as described above) depolarizing stimuli, and the spike frequency adaptation was also significantly reduced in mutant DRG neurons in
comparison with control neurons before TEA application; after TEA application, the control neurons generated statistically similar numbers
of APs in response to absolute stimuli compared with mutant neurons.

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

The results regarding the role of Kv7 channels in
mechanical allodynia and thermal hyperalgesia are
inconsistent. In one study, retigabine increases the tail
withdrawal threshold to noxious thermal stimuli in a
dose-dependent manner (Dost et al., 2004), but
because this effect was not reversed by the coadminis-
tration of linopridine, it may be due to a 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 Jen-
sen, 2003). Similarly, intraplantar injections of XE991
into the rat hindpaws did not induce thermal hyperalge-
sia or mechanical alldynia (Linley et al., 2008). These
studies shared the important limitation that the enhancers and blockers likely act on most or all Kv7 subunits and potentially on other channels (for example, GABA<sub>A</sub>; Otto et al., 2002). Unmyelinated axons express Kv7.5 (King and Scherer, 2012), which could also be the site of these pharmacological agents.

Our current study was designed to minimize these confounding factors by analyzing a type of myelinated sensory axons that lacks Kv7.2, the A-delta fibers. Although 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 Proudfoot, 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 mediated primarily by A-delta fibers. Specifically, Yeomans and Proudfoot (1996) found that radiant heating of rat hindpaw at a high rate of 6.5°C/second for 6 seconds evokes primarily an A-delta fiber response, whereas a low rate of 0.9°C/second for 20 seconds activates primarily a C fiber 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/second; Dirig et al., 1997), and because the average withdrawal latencies of both mutant and control animals in our Hargreaves test were less than 5 seconds (3.7 and 4.3 seconds, 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 might also be due to increase A-delta fiber activity. Finally, Passmore et al. (2012) showed that M-current inhibition by XE991 (at a 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 the mutant and the 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 also 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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**CONFLICT OF INTEREST STATEMENT**

The authors declare that they have no conflicts of interest.

**ROLE OF AUTHORS**

All authors had full access to all the data in the study and take responsibility for the integrity of the data and the accuracy of the data analysis. Study concept and design: CHK, EP, SSS. Acquisition of data: CHK, EL, DS. Analysis and interpretation of data: CHK, EL, DS, EP, SSS. Drafting of the manuscript: CHK, SSS. Critical revision of the manuscript for important intellectual content: CHK, EL, DS, EP, SSS. Statistical analysis: CHK. Obtained funding: EP, SSS. Administrative, technical, and material support: CHK, EL, DS, EP, SSS. Study supervision: EP, SSS.

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