

## PAIN

# The nicotinic $\alpha 6$ subunit gene determines variability in chronic pain sensitivity via cross-inhibition of P2X2/3 receptors

Jeffrey S. Wieskopf,<sup>1</sup> Jayanti Mathur,<sup>2</sup> Walrati Limapichat,<sup>3</sup> Michael R. Post,<sup>3</sup> Mona Al-Qazzaz,<sup>4</sup> Robert E. Sorge,<sup>1</sup> Loren J. Martin,<sup>1</sup> Dmitri V. Zaykin,<sup>5</sup> Shad B. Smith,<sup>6</sup> Kelen Freitas,<sup>7</sup> Jean-Sebastien Austin,<sup>1</sup> Feng Dai,<sup>8</sup> Jie Zhang,<sup>2</sup> Jaclyn Marcovitz,<sup>1</sup> Alexander H. Tuttle,<sup>1</sup> Peter M. Slepian,<sup>1</sup> Sarah Clarke,<sup>1</sup> Ryan M. Drenan,<sup>9</sup> Jeff Janes,<sup>2</sup> Shakir Al Sharari,<sup>10</sup> Samantha K. Segall,<sup>6</sup> Eske K. Aasvang,<sup>11</sup> Weike Lai,<sup>8</sup> Reinhard Bittner,<sup>12</sup> Christopher I. Richards,<sup>13</sup> Gary D. Slade,<sup>14</sup> Henrik Kehlet,<sup>11</sup> John Walker,<sup>2</sup> Uwe Maskos,<sup>15</sup> Jean-Pierre Changeux,<sup>15</sup> Marshall Devor,<sup>16</sup> William Maixner,<sup>6</sup> Luda Diatchenko,<sup>6,17</sup> Inna Belfer,<sup>8</sup> Dennis A. Dougherty,<sup>3</sup> Andrew I. Su,<sup>18</sup> Sarah C. R. Lummis,<sup>4</sup> M. Imad Damaj,<sup>7</sup> Henry A. Lester,<sup>19</sup> Ardem Patapoutian,<sup>20</sup> Jeffrey S. Mogil<sup>1\*</sup>

Chronic pain is a highly prevalent and poorly managed human health problem. We used microarray-based expression genomics in 25 inbred mouse strains to identify dorsal root ganglion (DRG)-expressed genetic contributors to mechanical allodynia, a prominent symptom of chronic pain. We identified expression levels of *Chrna6*, which encodes the  $\alpha 6$  subunit of the nicotinic acetylcholine receptor (nAChR), as highly associated with allodynia. We confirmed the importance of  $\alpha 6^*$  ( $\alpha 6$ -containing) nAChRs by analyzing both gain- and loss-of-function mutants. We find that mechanical allodynia associated with neuropathic and inflammatory injuries is significantly altered in  $\alpha 6^*$  mutants, and that  $\alpha 6^*$  but not  $\alpha 4^*$  nicotinic receptors are absolutely required for peripheral and/or spinal nicotine analgesia. Furthermore, we show that *Chrna6*'s role in analgesia is at least partially due to direct interaction and cross-inhibition of  $\alpha 6^*$  nAChRs with P2X2/3 receptors in DRG nociceptors. Finally, we establish the relevance of our results to humans by the observation of genetic association in patients suffering from chronic postsurgical and temporomandibular pain.

## INTRODUCTION

Chronic pain in the clinic manifests itself mainly in the form of spontaneous pain and mechanical allodynia, a sensitized response to an innocuous stimulus. Patients suffering from the latter symptom sometimes find that clothing touching their skin or a light breeze is very

painful. Current medications are largely inadequate to treat such symptoms, and an in-depth understanding of molecular mechanisms of mechanical allodynia is still lacking.

Here, we used an unbiased approach to identify genes involved in mechanical allodynia. Specifically, we correlated the mechanical allodynia phenotypes of 25 inbred mouse strains with genome-wide gene expression levels in dorsal root ganglia (DRGs) of these strains. This expression genomics strategy has been adopted previously in pain research (1–4), but using a much smaller number of strains. We provide evidence for the expression of the *Chrna6* gene encoding the nicotinic  $\alpha 6$  subunit as a major determinant of variable mechanical allodynia after nerve injury.

Neuronal nicotinic acetylcholine receptors (nAChRs) are hetero- or homopentameric ligand-gated ion channels composed of  $\alpha$  ( $\alpha 2$  to  $\alpha 7$ ,  $\alpha 9$ , and  $\alpha 10$ ) and  $\beta$  ( $\beta 2$  to  $\beta 4$ ) subunits. They have been the target of analgesic drug discovery for many years, with progress being hindered by a narrow therapeutic window and side effects. Attention has been focused largely on  $\alpha 4\beta 2^*$  ( $\alpha 4$ - and  $\beta 2$ -containing) nAChRs (5), the most highly expressed subtype in the central nervous system (CNS), but effects on pain of  $\alpha 3^*$  (6),  $\alpha 7^*$  (7), and  $\alpha 9^*$  (8, 9) nAChRs have also been demonstrated. The  $\alpha 6^*$  nAChRs have been mysterious until the recent elucidation of their involvement in the mesolimbic dopaminergic system [where they activate dopamine neurons to cause locomotor hyperactivity (10)] and visual system [where they modulate glutamate and  $\gamma$ -aminobutyric acid release in the superior colliculus (11)]. The  $\alpha 6$  subunit is localized in sensory ganglia (12–15). There are no reported agonists that discriminate well between  $\alpha 6^*$  and  $\alpha 4^*$  nAChRs, raising the possibility that the  $\alpha 6$  subunit plays an unappreciated role in nicotinic analgesia in the spinal cord or periphery.

<sup>1</sup>Department of Psychology and Alan Edwards Centre for Research on Pain, McGill University, Montreal, Quebec H3A 1B1, Canada. <sup>2</sup>Genomic Institute of the Novartis Research Foundation, San Diego, CA 92121, USA. <sup>3</sup>Division of Chemistry and Chemical Engineering, California Institute of Technology, Pasadena, CA 91125, USA. <sup>4</sup>Department of Biochemistry, University of Cambridge, Cambridge CB2 1QW, UK. <sup>5</sup>National Institute of Environmental Health Sciences, National Institutes of Health, Research Triangle Park, NC 27709, USA. <sup>6</sup>Center for Neurosensory Disorders, University of North Carolina at Chapel Hill, Chapel Hill, NC 27599, USA. <sup>7</sup>Department of Pharmacology and Toxicology, Virginia Commonwealth University, Richmond, VA 23219, USA. <sup>8</sup>Departments of Anesthesiology and Human Genetics, University of Pittsburgh, Pittsburgh, PA 15213, USA. <sup>9</sup>Department of Medicinal Chemistry and Molecular Pharmacology, Purdue University, West Lafayette, IN 47907, USA. <sup>10</sup>Department of Pharmacology, King Saud University, P. O. Box 2457, Riyadh 11451, Saudi Arabia. <sup>11</sup>Section for Surgical Pathophysiology, Rigshospitalet, Copenhagen University, 2100 Copenhagen, Denmark. <sup>12</sup>Department of Surgery, Marienhospital Stuttgart, 70199 Stuttgart, Germany. <sup>13</sup>Department of Chemistry, University of Kentucky, Lexington, KY 40506, USA. <sup>14</sup>Department of Dental Ecology, School of Dentistry, University of North Carolina at Chapel Hill, Chapel Hill, NC 27599, USA. <sup>15</sup>Neurobiologie Intégrative des Systèmes Cholinergiques, CNRS UMR 3571, Département de Neuroscience, Institut Pasteur, 75724 Paris, France. <sup>16</sup>Department of Cell and Developmental Biology, Institute of Life Sciences and Center for Research on Pain, Hebrew University of Jerusalem, Jerusalem 91904, Israel. <sup>17</sup>Faculty of Dentistry, Department of Anesthesia, and Alan Edwards Centre for Research on Pain, McGill University, Montreal, Quebec H3A 1G1, Canada. <sup>18</sup>Department of Molecular and Experimental Medicine, The Scripps Research Institute, La Jolla, CA 92037, USA. <sup>19</sup>Division of Biology and Biological Engineering, California Institute of Technology, Pasadena, CA 91125, USA. <sup>20</sup>Department of Molecular and Cellular Neuroscience, The Scripps Research Institute, and Howard Hughes Medical Institute, La Jolla, CA 92037, USA.

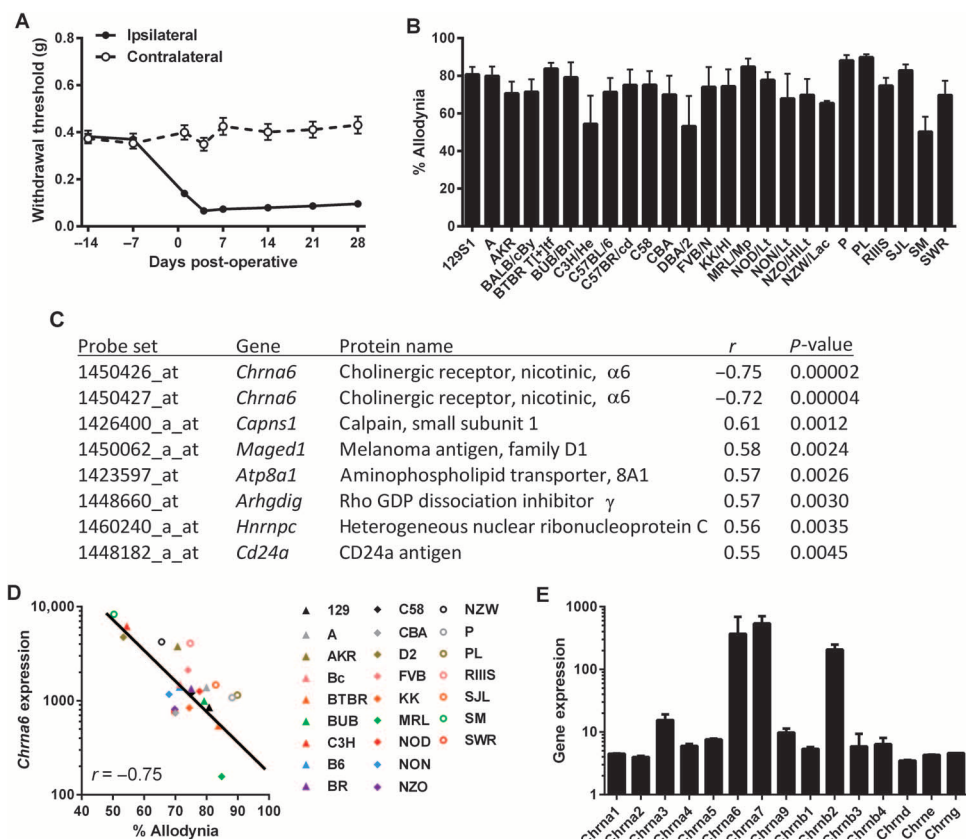
\*Corresponding author. E-mail: jeffrey.mogil@mcgill.ca

## RESULTS

**Association between dorsal root ganglion *Chrna6* expression and neuropathic mechanical allodynia in mice**

Mechanical allodynia induced by spared nerve injury (SNI) was quantified in 25 inbred mouse strains using von Frey filaments, and compared to basal DRG expression of 45,101 mRNA transcripts using microarray gene expression profiling (Affymetrix MOE430v2 chip) in these same 25 strains. All strains displayed ipsilateral allodynia (Fig. 1, A and B, and fig. S1), but highly significant effects of strain ( $F_{24,89} = 9.1$ ,  $P < 0.001$ ) and a strain  $\times$  repeated measures interaction ( $F_{144,534} = 2.2$ ,  $P < 0.001$ ) were observed. That is, strains displayed different extents and time courses of allodynia (fig. S1). A significant strain  $\times$  sex  $\times$  repeated measures interaction ( $F_{144,534} = 1.4$ ,  $P < 0.05$ , Greenhouse-Geisser corrected) was also evinced. This interaction appeared to be largely a result of robust sex differences in the SM/J (see below) and C3H/HeJ strains (16).

Correlating the overall allodynia data (Fig. 1B) with the Affymetrix chip data revealed eight correlations at  $P < 0.005$  (uncorrected) (Fig. 1C).

**Fig. 1. Correlation of DRG expression of *Chrna6* and mechanical allodynia after SNI in inbred mice.**

SNI surgery was performed on 25 inbred mouse strains, and withdrawal thresholds of the ipsilateral hind paw to von Frey fiber stimulation were measured. (A) Symbols ( $n = 139$ ) represent mean  $\pm$  SEM paw withdrawal threshold on each testing day. (B) Bars ( $n = 4$  to 6 mice per strain) represent mean  $\pm$  SEM percentage of maximum possible allodynia (see Supplementary Materials and Methods). (C) Top eight correlations (Pearson's  $r$ ; all  $P < 0.005$ ) between the strain means shown in (B) and basal DRG expression levels of  $\approx 45,000$  probe sets in these same strains. (D) Correlation between allodynia and basal DRG expression (in arbitrary units) of probe set 1450426\_at (*Chrna6*); symbols represent individual strain values. Strain abbreviations: Bc, BALB/cBy; B6, C57BL/6; BR, C57BR/cd; D2, DBA/2. (E) Average expression across all strains of all *Chrm\** genes. Bars represent basal DRG expression (in arbitrary units)  $\pm$  SD.

The top two highest correlations (in either direction), genome-wide, were with two different probes for *Chrna6* ( $r = -0.75$  and  $r = -0.72$ ) (Fig. 1, C and D) such that higher expression of *Chrna6* was associated with less development of allodynia. These associations were both significant at  $P < 0.05$  using false discovery rate correction for multiple comparisons; no other associations were significant, and thus no attempts were made to evaluate the candidacy of the other genes in Fig. 1C. The *Chrna6* correlations were sex-dependent, being considerably higher in males ( $r = -0.67$ ,  $-0.63$ ) than in females ( $r = -0.37$ ,  $-0.33$ ). Table S1 shows strain-dependent expression of all *Chrm\** genes coding for nicotinic subunits; none other than *Chrna6* featured suggestive correlations with allodynia or baseline nociception. As can be appreciated by the error bar, *Chrna6* displayed more genotypic variation in DRG expression than did any other subunit gene (Fig. 1E), with a coefficient of variation  $\approx 50\%$  higher than the next most variable subunit (*Chrb3*) and more than 10-fold higher than *Chrna4* (see table S1).

Conventional haplotype mapping was also performed, correlating mechanical allodynia strain means with about 156,000 genomic haplotypes (17). Of the top 10 correlated haplotypes genome-wide (see table S2), 2 of them were located just upstream

of the *Chrna6* gene on mouse chromosome 8. Other potentially associated genes include *Kcnv1*, *Ubc*, *Aldh7a1*, *Gfra2*, and *Chrna3* (located very near *Chrna6* on chromosome 14).

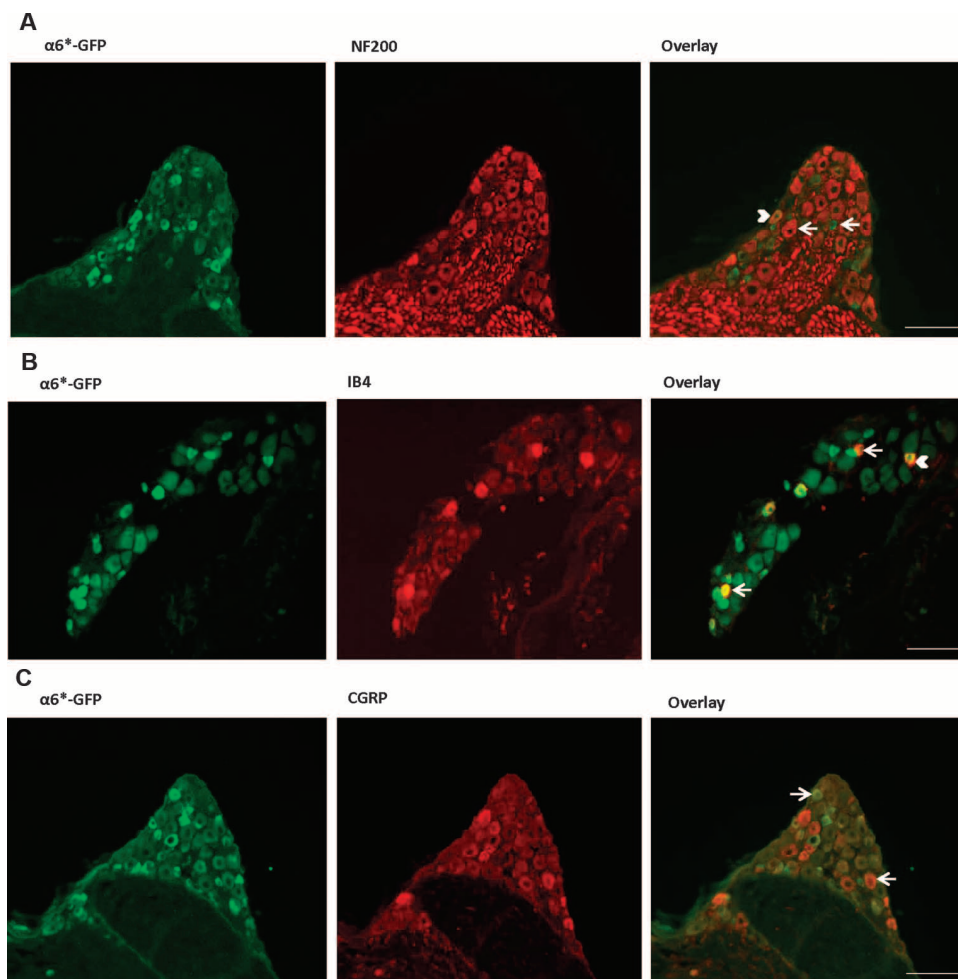
**Expression of *Chrna6* in a subset of DRG neurons**

Quantitative polymerase chain reaction (qPCR) experiments revealed detectable expression of *Chrna6* mRNA in whole brain, DRG, and eye, but not lung (Table 1). However, relative expression levels varied in these tissues, with DRG expression  $>10$ -fold higher than expression in whole brain and  $>2$ -fold higher than that in the eye (both  $P < 0.001$ ). In situ hybridizations performed at the Allen Institute for Brain Science showed the presence of *Chrna6* expression in small- to medium-diameter DRG neurons (www.brain-map.org), and this was confirmed in  $\alpha 6^*$ -GFP BAC transgenic mice (Fig. 2). DRGs were double-stained with green fluorescent protein (GFP) and neuronal sensory marker antibodies, including neurofilament 200 (NF200), which marks mostly myelinated A $\beta$  neurons, as well as isolectin B4 (IB4) and calcitonin gene-related peptide (CGRP), which mark distinct populations of nociceptive neurons. Sixty-six percent of the GFP-positive neurons also expressed NF200 (203 of 306 out of 840 total neurons counted), whereas 37% of NF200-positive neurons expressed GFP (123 of 336). Twenty-six percent of the GFP population also stained for IB4 (135 of 512 out of 1286), whereas 58% of IB4 population expressed GFP

(169 of 293). Finally, 8% of the GFP neurons also expressed CGRP (39 of 496 from a total of 1213), whereas 35% of CGRP-positive neurons expressed GFP (91 of 258). These results suggest that *Chrna6* is expressed in various functionally distinct DRG subtypes. A previous study observed a similar range of colabeling of *Chrna6* mRNA and peptidergic nociceptor-related (CGRP or transient receptor potential, V1) immunoreactivity in rat trigeminal ganglion (18).

**Table 1. Expression of *Chrna6* in multiple tissues.** Values represent mean  $\pm$  SEM expression normalized to *Actb* ( $\beta$ -actin). n.d., not detectable.

Tissue	Expression
DRG	0.42 $\pm$ 0.06
Eye	0.16 $\pm$ 0.01
Whole brain	0.04 $\pm$ 0.02
Lung	n.d.



**Fig. 2. *Chrna6* mRNA expression in a subset of DRG neurons.** (A to C) DRG neurons from adult  $\alpha 6^*$ -GFP mice were stained with antibodies against GFP (green) and sensory neuron markers NF200 (A), IB4 (B), and CGRP (C) (all red). Arrows indicate neurons expressing either GFP or the sensory marker. Filled arrowheads indicate neurons coexpressing both markers. Scale bars, 100  $\mu$ m.

We also confirmed genotype- and sex-dependent *Chrna6* expression in the DRG in three mouse strains using qPCR. As was observed in the gene chip experiment (fig. S2A), *Chrna6* expression appeared to be robustly strain-dependent and, in one strain, strongly sex-dependent, with male SM/J mice displaying >30-fold higher expression than female mice of the same strain (fig. S2B). We also observed a marked sex difference in SM/J mice in the development of mechanical allodynia after SNI (female > male; sex  $\times$  repeated measures:  $F_{6,78} = 3.7$ ,  $P < 0.005$ ) (fig. S2C).

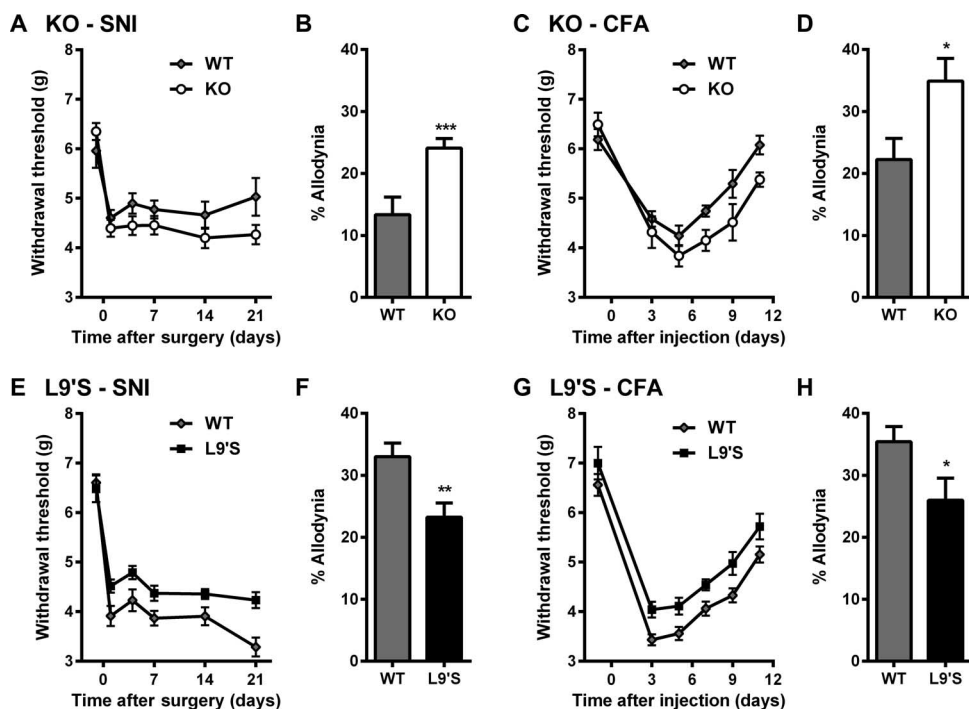
### Nerve injury-induced down-regulation of *Chrna6* and mechanical allodynia

In an independently performed experiment to identify genes associated with chronic pain (2), microarray gene expression profiling (also using the Affymetrix MOE430v2 chip) was performed in the DRGs of five inbred mouse strains after sham surgery or spinal nerve ligation (SNL), another common preclinical assay of neuropathic pain associated with mechanical allodynia. The two *Chrna6* probe sets appeared in the top 10 highest fold regulations by SNL compared to sham surgery (fig. S3A), exhibiting down-regulation of 67.3- and 41.6-fold, respectively. The down-regulation was highly correlated ( $r = 0.90$ ,  $P < 0.05$ ) with basal DRG expression (fig. S3B), but even after down-regulation by SNL, a strongly negative correlation ( $r = -0.93$ ) between *Chrna6* expression and allodynia was observed in the three strains tested behaviorally (fig. S3C), suggesting that *Chrna6* expression protects against allodynia after nerve injury as well.

### Confirmation of *Chrna6* involvement in mutant mice

To provide causal evidence of the involvement of the  $\alpha 6$  subunit in neuropathic pain, we tested transgenic *Chrna6* null mutant mice (19) and *Chrna6* gain-of-function L9'S mutant mice (10) for mechanical allodynia after SNI. To investigate whether  $\alpha 6^*$  nAChRs play a similar role in chronic inflammatory pain, we also tested these mutants for mechanical allodynia after intraplantar complete Freund's adjuvant (CFA) injection (Fig. 3). All genotypes displayed expected time courses of allodynia. For both SNI and CFA, *Chrna6* knockout (KO) mice showed higher overall levels of allodynia than did wild-type (WT) mice ( $t_{21} = 3.2$ ,  $P < 0.005$  and  $t_{10} = 2.5$ ,  $P < 0.05$ , respectively). For both SNI and CFA, *Chrna6* L9'S mutant mice displayed less allodynia overall than did their WT controls ( $t_9 = 3.0$ ,  $P = 0.01$  and  $t_{16} = 2.2$ ,  $P < 0.05$ , respectively). There were no significant genotype  $\times$  sex interactions observed in any data set. In an experiment performed independently, in a different





**Fig. 3. Differential mechanical allodynia after nerve injury and chronic inflammation in *Chrna6* mutant mice.** (A to H) Increased mechanical allodynia after SNI surgery (A and B) and CFA injection (C and D) in *Chrna6* KO mice, and decreased mechanical allodynia after SNI (E and F) and CFA (G and H) in *Chrna6* L9'S gain-of-function mutants. In all graphs, symbols ( $n = 5$  to 12 mice per genotype) represent mean  $\pm$  SEM paw withdrawal threshold (g) on each testing day; bars represent mean  $\pm$  SEM percentage of maximum possible allodynia (see Materials and Methods). \* $P < 0.05$ , \*\* $P < 0.01$ , \*\*\* $P < 0.001$  compared to other genotype. A replication of the KO data, using a different neuropathic assay, can be found in fig. S4.

laboratory, using *Chrna6* KO mice and another neuropathic assay [chronic constriction injury (CCI)], the increased mechanical allodynia of KO mice was confirmed (fig. S4A). A separate head-to-head experiment using CCI and CFA in *Chrna6* and *Chrna4* KO mice confirmed the significantly increased allodynia in *Chrna6* KOs, but revealed no differences between *Chrna4* KOs and their WT controls (fig. S4, B and C). The  $\alpha 6$  subunit appears to play a highly specific role in the modulation of mechanical allodynia because *Chrna6* KO mice showed statistically equivalent responses to WT mice on a battery of acute and tonic nociceptive assays (fig. S5).

### Relevance of $\alpha 6$ to anti-allodynic effects of nicotine

Nicotine itself exerts anti-allodynic effects after both inflammatory and neuropathic injuries (5). We tested the ability of systemic, intracerebroventricular, intrathecal, and peripheral (intraplantar) (–)–nicotine to reverse mechanical allodynia produced by either SNI or CFA in WT, KO, and L9'S mice. Although potency and efficacy varied by route of administration, nicotine was significantly and dose-dependently effective against both types of allodynia in WT mice by all injection routes (Fig. 4 and table S3). Gain-of-function L9'S mutants showed similar or significantly increased efficacy, but *Chrna6* KO mice displayed no significant nicotine-induced anti-allodynia in either assay by any route. We then performed a head-to-head comparison of supraspinal, spinal, and peripheral nicotine-induced anti-allodynia (25  $\mu$ g, intracerebroventricular; 17  $\mu$ g, intrathecal; 50  $\mu$ g, intraplantar) in *Chrna6* and *Chrna4* null mutants after neuropathic (CCI) or inflam-

matory (CFA) injury. All routes of administration produced robust reversal of both types of mechanical allodynia in both WT lines at these doses (Fig. 5). Supraspinal nicotine anti-allodynia was significantly reduced in *Chrna6* mutants (CCI:  $t_9 = 4.9$ ,  $P < 0.001$ ; CFA:  $t_{10} = 2.3$ ,  $P < 0.05$ ) and completely abolished in *Chrna4* mutants (CCI:  $t_8 = 4.1$ ,  $P < 0.01$ ; CFA:  $t_{10} = 3.5$ ,  $P < 0.01$ ) (Fig. 5A). By contrast, spinal nicotine anti-allodynia was abolished in *Chrna6* mutants (CCI:  $t_{20} = 5.7$ ,  $P < 0.001$ ; CFA:  $t_9 = 3.2$ ,  $P = 0.01$ ) and preserved in *Chrna4* mutants [CCI:  $t_8 = 1.9$ ,  $P =$  not significant (ns); CFA:  $t_{10} = 2.0$ ,  $P =$  ns] (Fig. 5B). Similarly, anti-allodynia resulting from injection of nicotine directly into the hind paw was abolished in *Chrna6* mutants (CCI:  $t_{10} = 6.4$ ,  $P < 0.001$ ; CFA:  $t_{10} = 3.7$ ,  $P < 0.01$ ) and preserved in *Chrna4* mutants (CCI:  $t_8 = 0.6$ ,  $P =$  ns; CFA:  $t_{10} = 1.7$ ,  $P =$  ns) (Fig. 5C). These data suggest that nicotine blocks mechanical allodynia in a wholly  $\alpha 6$ -specific manner, except supraspinally, where both  $\alpha 6^*$  and  $\alpha 4^*$  nicotinic receptors appear to contribute.

### Electrophysiological measurement of $\alpha 6^*$ and P2X2/3 receptor interactions

An anti-allodynic effect of  $\alpha 6^*$  activation (after DRG gene expression) suggests a functional interaction between  $\alpha 6^*$  nAChRs

and another pain-relevant molecular target in the spinal cord or periphery. Several subtypes of nAChRs interact, both functionally and physically, with several subtypes of P2X receptors (20–22). We therefore considered the hypothesis that  $\alpha 6^*$  nAChRs interact with P2X2 and P2X3 receptors, known to be involved in pain (23, 24) and, like  $\alpha 6^*$  nAChRs (see Fig. 2), to be expressed in the IB4-positive subpopulation of nociceptors (24).

We tested three combinations of nAChR subunits ( $\alpha 6\beta 4$ ,  $\alpha 6\beta 4\beta 3$ , and  $\alpha 6\beta 2$ ), coexpressed with most of the possible combinations of P2X2 and P2X3 subunits (P2X2, P2X3, and P2X2/3 receptors). Most  $\alpha 6^*$  nAChRs yield very small agonist-induced current in heterologous expression experiments, vitiating accurate measurements; we overcame these problems by using gain-of-function  $\alpha 6$  subunits [ $\alpha 6$ (L9'S) for  $\alpha 6\beta 4$ ] (10), gain-of-function  $\beta 3$  subunits [ $\beta 3$ (V13'S) for  $\alpha 6\beta 4\beta 3$ ] (25), or the combination [ $\alpha 6$ (L9'S) $\beta 2$ (L9'S) $_{\text{LFM/AAQA}}$ ] (26). P2X2 receptors and P2X2/3 receptors express robustly in oocytes; the latter are activated selectively by  $\alpha, \beta$ me-ATP (adenosine triphosphate) (27).

With seven of the eight combinations of  $\alpha 6^*$  receptors and P2X receptors studied, we found functional interactions, in the form of cross-inhibition, between these two classes of ligand-gated receptors. In the first type of interaction, when ACh and ATP are co-applied, the agonist-induced currents are less than the sum of individual currents. This type of interaction was previously observed between P2X receptors and several other pentameric receptors. When  $\alpha 6^*$  nAChRs were expressed alone, they showed no direct responses to ATP; the addition of ATP (320  $\mu$ M) produced  $<15\%$  change in the ACh-evoked currents

**Fig. 4. Altered anti-allodynic potency and efficacy of nicotine in *Chrna6* mutant mice.** (A to H) Dose-response relationships for the ability of systemic [intraperitoneal (i.p.); A and B], intracerebroventricular (i.c.v.; C and D), intrathecal (i.t.; E and F), and peripheral [intraplantar (i.pl.); G and H] nicotine to reverse already developed (and maximal) mechanical allodynia produced by SNI (day 7 after surgery; A, C, E, and G) and CFA (day 3 after injection; B, D, F, and H). Symbols ( $n = 4$  to 8 mice per dose per genotype) represent mean  $\pm$  SEM percentage of maximum possible anti-allodynia, based on the pre-SNI/CFA and post-SNI/CFA withdrawal thresholds of each mouse (see Materials and Methods). Statistical analyses are shown in table S3.

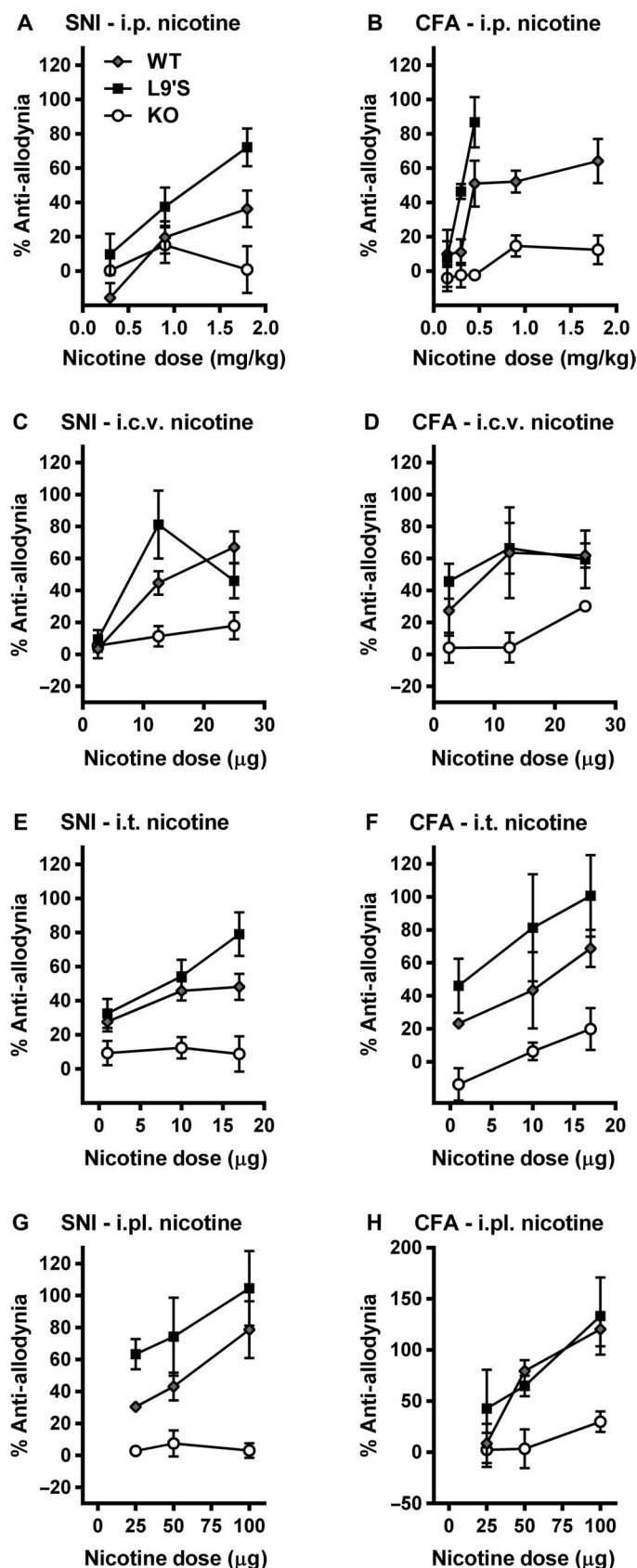
at any concentration. We found as well that P2X2, P2X3, or P2X2/3 currents were not affected by ACh (100  $\mu$ M). In four of the six cases where we could study dose-response relations, we found only minor (less than two-fold) changes in the  $EC_{50}$  (median effective concentration) values, and insignificant changes in the Hill coefficient, for each agonist when we coexpressed these receptors (table S4); an exception is described below. Despite lack of evidence of interactions from the shape of dose-response relations, when ACh and ATP are co-applied, the agonist-induced currents are less than the sum of individual currents (Fig. 6, A to C). This pattern was observed with all types of  $\alpha 6^*$  nAChR expressed with P2X2 (table S4), or with P2X2/3 receptors (Fig. 6, A to C). Cross-inhibition was also observed between  $\alpha 6(L9'S)\beta 4$  and P2X3 (K65A) receptors (28); the P2X3(K65A) mutation was used because it decreases the rate of desensitization (29).

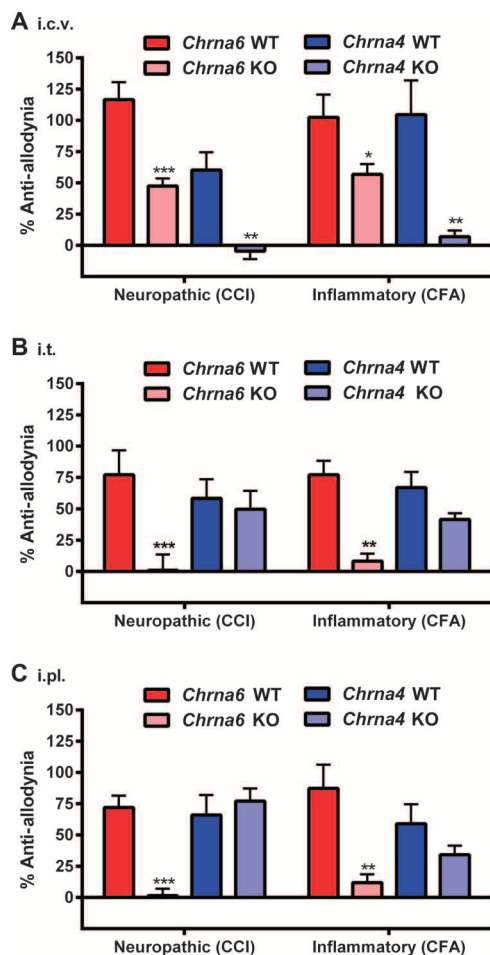
A second type of crosstalk occurs between  $\alpha 6\beta 2$  or  $\alpha 6\beta 4$  nAChRs and P2X receptors. The presence of coexpressed  $\alpha 6^*$  nAChRs changes the dose-response relation of the P2X3 receptor (28). This type of interaction has been previously reported only for the interaction between  $\alpha 3\beta 4$  nAChR and P2X2 (22). The  $EC_{50}$  of the P2X3 receptor is two to three times higher, and the response has decreased apparent cooperativity, revealed by a reduced Hill coefficient. As a result, responses to ATP in the concentration range 10 to 100  $\mu$ M are reduced by about half to two-thirds when normalized to maximal responses. These data are summarized in table S4 [see also (28)].

#### Fluorescence resonance energy transfer measurement of interaction between $\alpha 6^*$ and P2X3 receptors

We tested for physical interactions between  $\alpha 6\beta 4^*$  nAChRs and P2X receptors in cultured mouse cortical neurons, using fluorescence resonance energy transfer (FRET) as previously performed for P2X2 and  $\alpha 4\beta 2$  nAChRs (20). FRET typically reveals interactions between fluorophores that are less than 80 Å apart, implying a macromolecular complex. We tested for interactions between enhanced yellow fluorescent protein (eYFP)- and mCherry-labeled receptors, using fluorescence lifetime imaging microscopy (FLIM). Results show that P2X3 and  $\alpha 6\beta 4$  receptors physically interact, with a FRET efficiency of  $\approx 50\%$  (fig. S6) and a binding fraction of  $\approx 40\%$ .

We tested FRET with and without incorporation of a nonfluorescent  $\beta 3$  subunit into the  $\alpha 6\beta 4$  receptor (fig. S6, B and C). The incorporation of  $\beta 3$  did not alter the binding fraction or the FRET efficiency. To test whether the  $\beta 3$  simply did not become incorporated into the  $\alpha 6\beta 4$  receptor, we measured FRET in cells transfected with  $\alpha 6$ ,  $\beta 3$ -eYFP, and  $\beta 4$  subunits and with P2X3-mCherry receptors, where the fluorophores were located on the  $\beta 3$  and P2X3. This resulted in FRET efficiency of  $\approx 50\%$ , indicating that the incorporation of the  $\beta 3$  subunit does not change the FRET efficiency between P2X3 and  $\alpha 6$ -containing nAChRs. Because some of the electrophysiological data were obtained with the gain-of-function  $\alpha 6$  mutant, FRET imaging was also performed in the





**Fig. 5. Dependence of spinal and/or peripheral nicotine anti-allodynia on  $\alpha 6$ .** (A to C) Head-to-head comparison of supraspinal (25  $\mu$ g, intracerebroventricular) (A), spinal (17  $\mu$ g, intrathecal) (B), and peripheral (50  $\mu$ g, intraplantar) (C) nicotine anti-allodynia against neuropathic (CCI) and inflammatory (CFA) pain in *Chrna6* ( $\alpha 6^*$ ) and *Chrna4* ( $\alpha 4^*$ ) WT and KO mice tested using identical parameters at the peak of allodynia (14 days after CCI, 3 days after CFA). Bars ( $n = 5$  to 6 mice per genotype per injury) represent mean  $\pm$  SEM percentage of maximum possible anti-allodynia (see Materials and Methods). \* $P < 0.01$ , \*\* $P < 0.01$ , \*\*\* $P < 0.001$  compared to analogous WT.

mutant, expressed with  $\beta 4$ -eYFP subunit and P2X3-mCherry. The FRET obtained with  $\alpha 6$ (L9'S) $\beta 4$  receptor with P2X3 did not differ statistically from the data obtained with the WT  $\alpha 6$  subunit. A range of control experiments (including FRET determination of soluble eYFP and P2X3-mCherry, P2X3-eYFP and  $\alpha 6$ -mCherry, P2X3-eYFP and  $\beta 4$ -mCherry, and P2X3-eYFP and plasma membrane-anchored mCherry) were negative.

### Behavioral measurement of $\alpha 6^*$ and P2X2/3 receptor interactions

Because P2X3 receptors mediate both neuropathic and inflammatory pain (23, 24), and there is precedent for the ability of a protein (P2X7) to affect pain indirectly via down-regulation of P2X3 receptors (30), we assessed whether P2X3-dependent pain could be affected by activation of  $\alpha 6^*$  receptors, as suggested by the observed cross-inhibitions in a heterologous expression system. The P2X3 agonist,  $\alpha, \beta$ me-ATP, injected into the hind paw produced frank nocifensive (licking) behav-

ior of equivalent intensity in all three genotypes (Fig. 7A). The pain behavior was dose-dependently reversed by systemic nicotine in WT ( $F_{3,18} = 9.1$ ,  $P < 0.001$ ) and L9'S mice ( $F_{3,11} = 6.1$ ,  $P = 0.01$ ), as well as by the P2X3 receptor antagonist A-317491 ( $t_{10} = 4.6$ ,  $P < 0.001$ ). In *Chrna6* KO mice, statistically significant reversal ( $F_{3,28} = 3.1$ ,  $P = 0.04$ ) was only achieved at the highest dose and to a lesser degree than in the other two genotypes ( $P < 0.01$ ) (Fig. 7A).  $\alpha, \beta$ me-ATP also produced A-317491-reversible mechanical allodynia of equal magnitude in the three genotypes, which was completely reversed by nicotine (0.9 mg/kg) in L9'S mice, partially reversed in WT mice, and unaffected by nicotine in KO mice ( $P < 0.05$  compared to L9'S) (Fig. 7B).

### CHRNA6 and variable chronic pain in humans

A human cohort of 429 adults who underwent herniotomy (31) was genotyped at three *CHRNA6* polymorphisms that cover haplotypic diversity in the gene locus to test the association of the gene with clinical pain. One promoter region single-nucleotide polymorphism (SNP) (rs7828365) was found to be associated with changes in pain susceptibility under a recessive inheritance model, in which the minor allele homozygote (TT) showed an increased risk of persistent pain at 6 months after surgery (odds ratio, 1.20; SEM, 1.1;  $P = 0.03$ ; Fig. 8A). Only eight TT homozygotes were present in the cohort, and thus the association  $P$  value was computed by a permutation  $t$  test, which is robust in the presence of small expected counts. To replicate this finding, we genotyped rs7828365 in another cohort where the clinical pain phenotype was thoroughly characterized: 159 Caucasian females with temporomandibular disorder (TMD) (32). Although only two individuals were TT homozygotes, in agreement with findings from the postsurgical pain cohort, these TT-carrying TMD patients experienced substantially higher intensity and greater duration of clinical head and orofacial pain symptoms on a normalized composite score incorporating multiple domains of the CPSQ (33). The pain increase was significant as tested by a permutation  $t$  test ( $P = 0.03$ ; Fig. 8B). This cohort was also tested for association with SNPs in the *CHRNA4* (three SNPs), *CHRNA5* (eight SNPs), and *CHRNB2* (two SNPs) genes; no  $P$  value was lower than 0.40.

To estimate the overall effect of the genotype TT on chronic pain in human subjects, we combined the results of the two human studies, yielding  $P = 0.002$ , which remained significant ( $P = 0.02$ ) after adjusting for three SNPs and three inheritance models examined.

### DISCUSSION

The current studies reveal, via expression genomics performed in the mouse, an unexpected role of the *Chrna6* gene and  $\alpha 6^*$  receptors in mediating mechanical allodynia after nerve injury or inflammation and in the reversal of such mechanical allodynia by nicotine. We show that (i)  $\alpha 6^*$  receptors are expressed in a subset of nociceptors within the DRG; (ii) *Chrna6* expression in the DRG correlates with mechanical allodynia across mouse strains; (iii) mutant mice showing null expression or overexpression of *Chrna6* display more and less mechanical allodynia, respectively; and (iv) nicotine's spinal and peripheral anti-allodynic effects are mediated by  $\alpha 6^*$  rather than  $\alpha 4^*$  receptors, although both subunits participate in supraspinal effects of nicotine. Further, we have defined a plausible mechanism (although not necessarily the only one) whereby  $\alpha 6^*$  receptors can ameliorate chronic pain, via cross-inhibition with P2X2/3 receptors facilitated by direct contact between the proteins, demonstrated in vitro and behaviorally. Nicotine

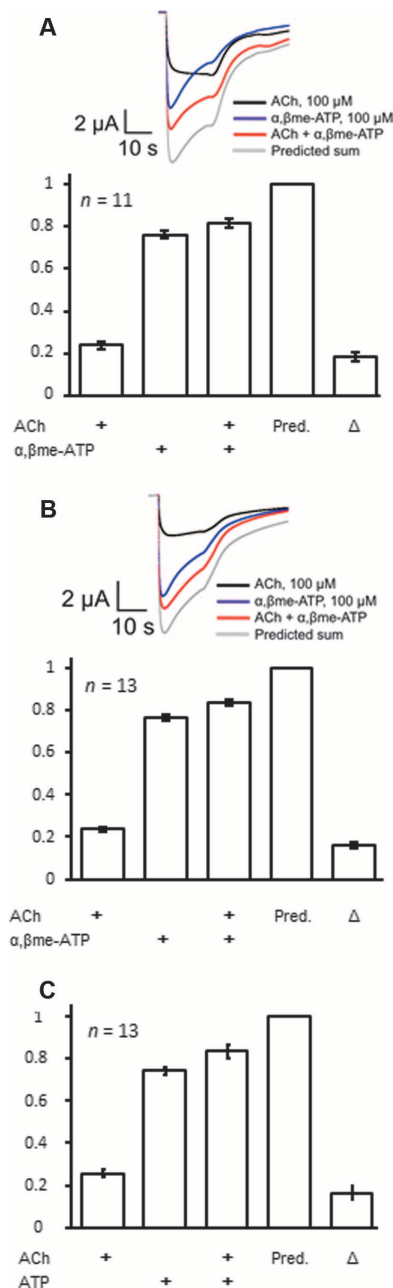


may be using this  $\alpha 6^*$ - and P2X2/3-dependent mechanism to produce pain relief, although the statistically significant (but incomplete) analgesia at a nicotine dose of 1.35 mg/kg in *Chrna6* null mutants suggests that alternative mechanisms may also be recruited.

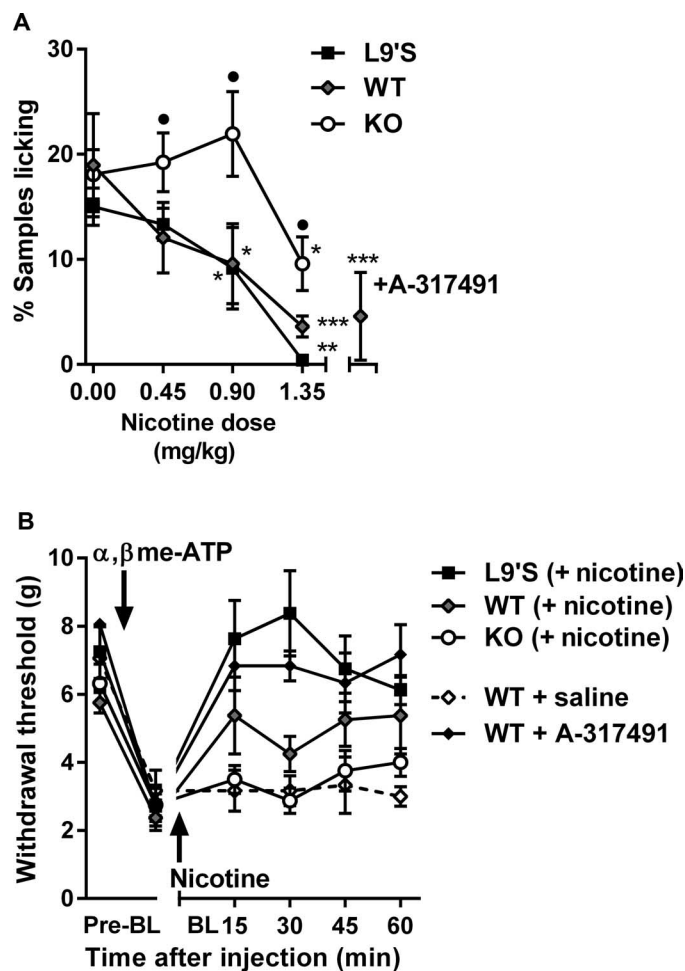
In addition, we have demonstrated the relevance of the *CHRNA6* gene in humans; the fact that TT homozygotes report substantially higher clinical pain in two very different chronic pain disorders reinforces the notion that  $\alpha 6^*$  receptors are playing a similar role, qualitatively, in pain biology in mice and humans. The rare frequency of the TT genotype suggests that *CHRNA6* is not a primary explanation of chronic pain variability in our species as it appears to be in the mouse. This is very likely simply due to species differences in frequencies of the trait-relevant alleles. The utility of rare variants in the validation of molecular targets for pain is well accepted (34, 35). Nonetheless, the highly limited sample size of TT homozygotes in the present study represents a limitation of the human genetic findings. The true role of *CHRNA6* in pain awaits investigation in more highly powered cohorts.

After the discovery of the remarkably high analgesic potency of the frog alkaloid epibatidine (36), pioneering investigations on epibatidine analogs were interpreted in terms of agonism of  $\alpha 4\beta 2^*$  nAChRs (37, 38). Preclinical and limited clinical evidence suggests that  $\alpha 4\beta 2^*$  subtypes play an important role in nociception.  $\alpha 4\beta 2^*$  nAChRs are expressed in many CNS regions that modulate pain transmission, and  $\alpha 4\beta 2^*$  agonists induce increases in inhibitory tone in the spinal cord. Several high-affinity  $\alpha 4\beta 2^*$  nAChR agonists were reported to have potent analgesic activity in rodent models of acute and chronic pain. Furthermore, evidence from studies using KO mice for the  $\alpha 4^*$  and  $\beta 2^*$  nicotinic subunits showed the dependency of nicotinic analgesia on this subtype (39), although this conclusion was largely based on nicotine inhibition of acute, thermal pain. Much less is known about the composition of other nAChRs mediating analgesia. Various other nAChR subunits—including  $\alpha 5$ ,  $\alpha 6$ ,  $\alpha 7$ ,  $\beta 3$ , and  $\beta 4$ —are incorporated into subtypes (for example,  $\alpha 4\alpha 6\beta 2^*$ ,  $\alpha 6\beta 2^*$ , and  $\alpha 6\beta 4^*$ ) that have been identified in the spinal cord and DRG tissues. Previous pharmacological and genetic evidence has challenged the assumption that the  $\alpha 4\beta 2$  subtype is the main analgesic target (40–43), as does the recently reported clinical trial failure of ABT-894 (44).

The present observations suggest that  $\alpha 6^*$  nicotinic receptors produce their analgesic effects by cross-inhibition of P2X2/3 receptors involving direct contact between the proteins. Other nAChR-P2X receptor interactions increase with the densities of the receptors (21). The details of the contacts, the receptor states involved, and the possible roles of ion flux are not fully known, but modifications to desensitization could play a role (28, 45). Purinergic receptors are important pain processing molecules known to be expressed on nociceptive small-diameter neurons in the DRG (46), with important roles having been demonstrated for P2X2/3 (23, 24, 47), P2X4 (48), and P2X7 (49). P2X2/3 receptors contribute to multiple pain modalities, including inflammatory pain (23, 24, 47, 50), neuropathic pain (51), visceral pain (52), musculoskeletal pain (53), cancer pain (54), and migraine (55). Presumably, the  $\alpha 6^*$  nicotinic receptors interacting in the periphery with P2X2/3 receptors are activated endogenously by acetylcholine, which exists abundantly in mammals both neuronally and nonneuronally, for example, in keratinocytes (56). Expression levels of choline acetyltransferase in mouse epidermis exceed that of any other tissue. Furthermore, experiments examining the pharmacokinetic profile of A-317491 in a rat skin-nerve preparation have suggested a peripheral site of action of P2X2/3 receptors on pain processing (57).

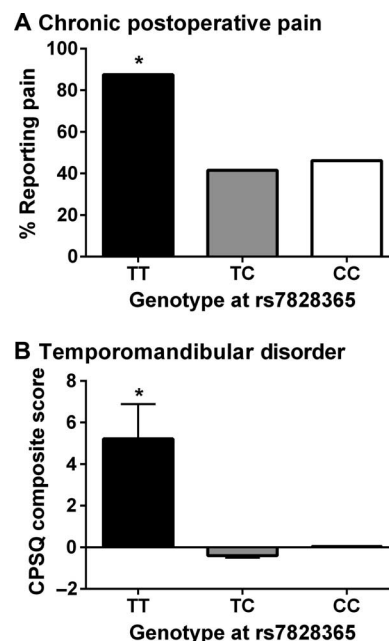


**Fig. 6. Electrophysiological detection of crosstalk between P2X and  $\alpha 6^*$  receptors in *Xenopus oocytes*.** (A) Coexpressed P2X2/3 and  $\alpha 6\beta 4$  receptors. (B) Coexpressed P2X2/3 and  $\alpha 6\beta 4\beta 3$  receptors. Exemplar inward currents are shown, evoked with 100  $\mu$ M  $\alpha, \beta$ me-ATP, 100  $\mu$ M ACh, or a mixture of 100  $\mu$ M  $\alpha, \beta$ me-ATP + ACh. (C) Coexpressed P2X2 and  $\alpha 6\beta 2$  receptors were tested with 100  $\mu$ M ATP, 10  $\mu$ M ACh, or a mixture of the two agonists. All graphs summarize experiments from  $n = 11$  to 13 cells. The “predicted” (Pred.) current for each cell is the arithmetic sum of the  $I_{\alpha, \beta$ me-ATP} and  $I_{ACh}$  currents. The “ $\Delta$ ” current for each cell is the predicted current minus the observed  $I_{\alpha, \beta$ me-ATP+ACh} current. Error bars represent SEM. To provide measurable responses, several subunits were mutated as described in Materials and Methods.



**Fig. 7. Modulation of P2X2/3 agonist-induced pain and hypersensitivity by nicotine in *Chrna6* mutant mice.** (A) Intrathecal administration of  $\alpha,\beta$ me-ATP produces nocifensive (licking) behavior inhibited by the P2X3 antagonist, A-317491 (300 nmol, intraplantar) and systemic nicotine in WT and L9'S but not KO mice (except at the highest dose). Symbols ( $n = 4$  to 11 mice per dose per genotype) represent mean  $\pm$  SEM samples featuring licking behavior (see Materials and Methods). \* $P < 0.05$ , \*\* $P < 0.01$ , \*\*\* $P < 0.001$  compared to zero dose within genotype.  $P < 0.05$  compared to other genotypes within dose. (B) Intrathecal administration of  $\alpha,\beta$ me-ATP produces mechanical allodynia reversed by A-317491 (300 nmol, intraplantar) and nicotine (0.9 mg/kg) in WT and L9'S but not KO mice. Symbols ( $n = 4$  mice per genotype or drug) represent mean  $\pm$  SEM paw withdrawal threshold at each time point.

Our current data demonstrate that, in both chronic inflammatory and neuropathic pain models, nicotine blocks mechanical allodynia—a more important clinical symptom (58) than the acute thermal pain that has been often studied in this area—in an  $\alpha6^*$ -specific manner, and that the role of  $\alpha4^*$  in this phenomenon is limited or absent, except in the supraspinal compartment. It is therefore possible that the modest efficacy of some  $\alpha4\beta2^*$  agonists reported in animal models of chronic inflammatory pain (43) and initial clinical studies (38) may be related to their insufficient binding and/or functional activity at  $\alpha6^*$  subtypes, including both  $\alpha6\beta2^*$  and  $\alpha6\beta4^*$  subtypes studied here; Hone *et al.* (12) found that the latter have larger responses to ACh.



**Fig. 8. Human clinical pain is affected by a promoter SNP (rs7828365) in *CHRNA6*.** (A) Percentage of herniotomy patients reporting persistent pain 6 months after surgery, stratified by rs7828365 genotype (TT:  $n = 8$ ; TC:  $n = 82$ ; CC:  $n = 325$ ). (B) Chronic pain symptom questionnaire (CPSQ) composite pain scores (see Materials and Methods) of TMD patients stratified by rs7828365 genotype (TT:  $n = 2$ ; TC:  $n = 38$ ; CC:  $n = 117$ ). \* $P < 0.05$  compared to other genotypes.

A recent paper shows equal binding affinity of ABT-894 to  $\alpha6\beta2^*$  and  $\alpha6\beta4^*$  nAChRs in monkey striatum (59), but functional selectivity and efficacy were not reported. We believe that the refocusing of nAChR analgesic development on  $\alpha6^*$ -containing receptors could lead to much more efficacious compounds, which, due to the comparatively limited distribution of *Chrna6* expression, should display a favorable side effect profile compared to current drugs. Peripheral targeting of such compounds would make them even more attractive because our findings suggest that efficacy would be preserved, whereas side effects—for example, related to enhanced dopamine release (60) and engagement of brain reward pathways (61)—would be reduced yet further. Development of  $\alpha6^*$ -acting drugs for the treatment of Parkinson's disease and nicotine addiction have been hampered by the inability to achieve heterologous expression of  $\alpha6^*$ -containing receptors and designing ligands that discriminate  $\alpha6^*$  from  $\alpha3^*$ , but mutation-based improvements have recently been reported (62, 63). Our findings also reveal a potential side effect of  $\alpha6^*$  antagonists, which may be a concern as they are being considered for the treatment of tobacco addiction (64).

## MATERIALS AND METHODS

### Study design

This study represents a series of experiments using multiple techniques—including genetics, pharmacology, and electrophysiology—in *Xenopus* oocytes, mice, and humans. All in vivo studies and tissue harvests were performed in accordance with national and institutional guidelines, and were approved by animal care and use committees at McGill



University, Virginia Commonwealth University, and the California Institute of Technology. In pharmacological studies, mice were assigned to experimental groups using within-cage randomization. Blinding to genotype was in general not possible because of coat color variation; experimenters were, however, blinded to drug and dose. Power analyses were in general not possible because the effect size of genotype and drug effects were not predictable a priori; sample sizes in this study are consistent with norms in the field (65). Data from three mice in the strain survey were omitted from further analysis because they were identified as statistical outliers (Studentized residual >3).

## Mice

Mice in the 25-strain survey were naïve, adult (6- to 14-week-old) mice of both sexes obtained from The Jackson Laboratory. Strains included 129S1, A, AKR, BALB/cBy, BTBR *T+ tf*, BUB/Bn, C3H/He, C57BL/6, C57BR/cd, C58, CBA, DBA/2, FVB/N, KK/Hl, MRL/Mp, NOD/Lt, NON/Lt, NZO/HlLt, NZW/Lac, P, PL, RIIS, SJL, SM, and SWR (all “J” substrains). Mouse strains used in the Persson *et al.* (2) experiment, some of whose results are reanalyzed here, included AKR/J, C3H/HeJ, C57BL/6J, C58/J, and CBA/J. Subjects of all subsequent experiments were naïve, adult (6- to 14-week-old) C57BL/6J mice bred in our vivarium from breeders obtained from The Jackson Laboratory, mice with L9'S gain of function of  $\alpha 6^*$  nAChRs and their WT controls (10), transgenic *Chrna6* null mutant (KO) mice and their WT controls (19),  $\alpha 6^*$ -GFP BAC transgenic mice (11), or *Chrna4* ( $\alpha 4^*$ ) KO mice and their WT controls (66). In all cases, equal numbers of male and female mice were used. All mutants have been bred fully congenic (>10 generations) to C57BL/6. All mice were housed in standard polycarbonate cages in groups of two to five same-sex littermates in a temperature-controlled ( $20^\circ \pm 1^\circ\text{C}$ ) environment (14:10-hour light/dark cycle; lights on at 0700); tap water and food (Harlan Teklad 8604) were available ad libitum. All behavioral experiments were performed by male personnel.

## Microarray gene expression profiling

DRGs were dissected from naïve 2-month-old mice of both sexes ( $n = 3$  mice per sex per strain) between 0900 and 1200. Total RNA was isolated from tissues using TRIzol (Invitrogen) followed by RNeasy (Qiagen). RNA quality was examined on an Experion (Bio-Rad) instrument. Complementary DNA (cDNA) and amplified antisense RNA were made from 1.4  $\mu\text{g}$  of pooled total RNA, using the Affymetrix single amplification protocol. Affymetrix MOE430v2 arrays were hybridized, washed, stained, and scanned using standard Affymetrix protocols.

Quality control was achieved as follows. Total RNA samples were only used for pools if the 28S ribosomal RNA (rRNA) bands were at least twice the intensity of 18S rRNA bands. Arrays were only included in the final data set if the following metrics from the Affymetrix MAS5 algorithm were met: (i) percent present calls  $\geq 40$  and (ii) scaling factors, percent present calls, and background were all within 2 SDs of the mean. Signal intensity histograms, displayed in the MAS5 software, contained no outliers when observed visually. Hierarchical clustering of samples was also examined for outliers that might indicate arrays and RNA with good quality but from poorly dissected tissue; there were no such outliers.

## Haplotype mapping

Haplotype association mapping was performed as described (17, 67). Briefly, local haplotypes were computed for all inbred mouse strains by

analyzing a sliding window of SNP genotypes. Strains were grouped on the basis of haplotype group assignment, and the *F* statistic was used to quantify the association between that local haplotype and the phenotype of interest. The significance of that *F* statistic was computed nonparametrically on the basis of a weighted bootstrap method that accounted for the inherent population structure in the panel of inbred mouse strains (17).

## Immunofluorescence

Adult male  $\alpha 6^*$ -GFP transgenic mice were perfused with 4% paraformaldehyde, and DRGs from all levels were quickly dissected. The DRGs were postfixed overnight in 30% sucrose and embedded in optimum cutting temperature compound. DRGs were sectioned with a cryostat at 10- $\mu\text{m}$  thickness and mounted on Superfrost Plus slides and stored at  $-80^\circ\text{C}$ .

Frozen slides were air-dried at room temperature for 1 hour. Slides were blocked for 1 hour at room temperature in phosphate-buffered saline (PBS) plus Triton X-100-containing 3% bovine serum albumin and 10% goat serum or donkey serum, and overnight at  $4^\circ\text{C}$  with primary antibodies diluted in the blocking solution. The primary antibodies used were as follows: rabbit anti-GFP (1:500, Life Technologies, catalog no. A11122), goat anti-GFP (1:150, LifeSpan Biosciences, catalog no. LS-C48996), rabbit anti-CGRP (1:1000, Abcam, catalog no. ab-47027), and rabbit anti-NF200 (1:500, Sigma, catalog no. N4142). The sections were then washed three times in PBS with Triton X-100 and incubated at room temperature for 1 hour with secondary antibodies conjugated to Alexa 488 or Alexa 568 fluorochromes (Life Technologies) diluted 1:200 in blocking solution. To detect IB4 staining, GS-IB4–Alexa 568 (Life Technologies, catalog no. I21412) was diluted 1:200 and incubated during secondary antibody incubations. Sections were then washed three times in PBS with Triton X-100 and mounted in SlowFade gold anti-fade medium with 4',6-diamidino-2-phenylindole (Life Technologies).

## Image acquisition and quantification

Fluorescence images were acquired using an AX70 microscope (Olympus). Images were taken using identical acquisition parameters, and raw images were analyzed with MetaMorph software. Neurons were considered GFP-positive if the mean fluorescence intensity, measured as arbitrary units, was higher than the mean background fluorescence. This was set as the threshold to include all the GFP-positive cells. Regions were drawn around the GFP-positive cells, and these regions were transferred over to the other sensory marker to image coexpressing neurons. Cells were considered positive for NF200, IB4, or CGRP if the mean fluorescence intensity was higher than the mean background fluorescence.

## Real-time qPCR

For tissue comparison experiments, DRGs were freshly isolated from adult male C57BL/6J mice and snap-frozen on dry ice, and total RNA was isolated using TRIzol treatment. Total RNA from all other tissues was purchased from Zyagen. For strain comparison experiments, DRGs from different inbred strains were isolated ( $n = 3$  mice per sex per strain) and treated similarly. Total RNA (200 ng) was used to generate the first-strand cDNA using the QuantiTect Reverse Transcription kit (Qiagen). A real-time TaqMan PCR assay for *Chrna6* (assay ID: Mm00517529\_m1) was purchased from Life Technologies, with a FAM reporter dye and a nonfluorescent quencher. FastStart Universal Probe Master Mix (Rox)

from Roche Diagnostics was used. The reaction was run, in triplicate, in the ABI 7900HT Fast Real-Time System using 0.5  $\mu$ l of the cDNA in a 10- $\mu$ l reaction as per the manufacturer's instructions.

Calibrations and normalizations were done using the  $2^{-\Delta\Delta C_T}$  method. The target gene was *Chrna6*, whereas the reference gene was *Actb* ( $\beta$ -actin). The calibrator for the tissue comparisons was the DRG; the calibrator for the strain comparisons was the DBA/2 strain.

### Oocyte expression and analysis

Rat  $\alpha 6$ , rat  $\beta 2$ , and mouse  $\beta 3$  nAChR subunits were in the pGEM vector, and rat  $\beta 4$  nAChR was in the pAMV vector. All P2X cDNAs were in the pcDNA3 vector. Site-directed mutagenesis was performed using the Stratagene QuikChange protocol and verified through sequencing. Circular cDNA was linearized and then used as a template for *in vitro* transcription. Stage V to VI *Xenopus laevis* oocytes were injected with 50 nl of mRNA solution. To express the  $\alpha 6\beta 4$  combination, we used a hypersensitive  $\alpha 6$  subunit containing a serine mutation at the leucine $9'$  in the M2 domain (residue 279). To express the  $\alpha 6\beta 4\beta 3$  combination, we used the WT  $\alpha 6$  and  $\beta 4$  in combination with the hypersensitive  $\beta 3$  containing a serine mutation at the valine $13'$  in M2 (residue 283). When  $\alpha 6\beta 4^*$  nAChR and P2X receptors were coexpressed, equal volumes of corresponding mRNA solutions were mixed before the oocyte injection. To express the  $\alpha 6\beta 2$  combination, we used the hypersensitive  $\alpha 6$  subunit, as well as a hypersensitive  $\beta 2$  subunit containing a serine mutation at the leucine $9'$  in M2 and two endoplasmic reticulum export-enhancing mutations (26). To study P2X $_3$ , we used the K65A mutation, which accelerated the rate of recovery from desensitization. The  $\alpha 6\beta 2$ P2X $_2/\alpha 6\beta 2$ P2X $_3$  mRNA injection ratios were 10:10:1 and 1:1:1, respectively, at 5 ng per oocyte total mRNA. P2X $_2/3$  was expressed by co-injection of 1:10 ratio of P2X $_2$ /P2X $_3$  mRNA. After mRNA injection, oocytes were incubated for 12 to 72 hours at 18°C in culture medium (ND96 $^+$  with 5% horse serum).

Two-electrode voltage-clamp recordings used the OpusXpress 6000A (Axon Instruments). For cross-inhibition experiments on P2X $_3$  (K65A), the concentration of ATP was 100  $\mu$ M for cells expressing P2X $_3$  (K65A) and  $\alpha 6\beta 4\beta 3$ (V13'S) or 320  $\mu$ M for P2X $_3$ (K65A) and  $\alpha 6$ (L9'S) $\beta 4$ . To investigate the cross-interaction between P2X $_2/3$  receptor and  $\alpha 6\beta 4^*$  nAChRs, the P2X $_2/3$  receptor was activated by 100  $\mu$ M  $\alpha, \beta$ me-ATP, and the  $\alpha 6^*$  nAChR by 100  $\mu$ M ACh. Peak currents from at least three traces were averaged from the same cell for data analysis.

All dose-response data were normalized to the maximal current ( $I_{\max} = 1$ ) of the same cell and then averaged.  $EC_{50}$  and Hill coefficient ( $n_H$ ) were determined by fitting averaged, normalized dose-response relations to the Hill equation. Dose-response relationships of individual oocytes were also examined and used to determine outliers.

For all cross-interaction data involving P2X $_2$  or P2X $_2/3$ , the predicted current from agonist co-application was calculated from the arithmetic sum of  $I_{\text{ACh}}$  and  $I_{\text{ATP}}$  (or  $I_{\alpha, \beta \text{me-ATP}}$ ) from the same cell. The actual, observed current upon co-application of the agonists was subtracted from the prediction value of the same cell, and this difference was designated as  $\Delta$ . All current data and  $\Delta$  were normalized to the prediction value of the same cell, and then the normalized data were averaged across seven or more cells from two or more batches of oocytes.

For all cross-interaction data on the P2X $_3$ (K65A) receptor, co-application of the agonists used the "prolonged plus brief pulse" protocol (28). Averaged ATP-evoked peak current during ACh application ( $I_{\text{ATP}}^*$ ) was subtracted from averaged ATP-evoked current in

the absence of ACh ( $I_{\text{ATP}}$ ) from the same cell to obtain a  $\Delta^*$ . All current data and  $\Delta^*$  were normalized to ( $I_{\text{ATP}}$ ) and averaged across eight or more cells from two or more batches of oocytes.

### Neuronal cultures

Cortical neurons were extracted from day 17 mouse embryos and plated on 35-mm MatTek polylysine-coated glass bottom culture dishes in a neuronal medium containing Neurobasal, B27 (Invitrogen), and GlutaMAX supplemented with 3% equine serum. Neurons were plated at a density of 60,000 cells per dish. On day 4 of culture, neurons were treated with 1  $\mu$ M cytosine arabinoside. Neurons were maintained via 50% exchange with feeding medium (Neurobasal, B27, and GlutaMAX) twice per week. On day 7 in culture, plasmids were mixed in 100  $\mu$ l of Opti-MEM, although 4  $\mu$ l of Lipofectamine 2000 was mixed with a separate 100- $\mu$ l aliquot of Opti-MEM. After 5 min at 22°C, the separate solutions were mixed together and kept at room temperature for an additional 25 min. Neurons were transfected with 500 ng of each nAChR plasmids ( $\alpha 6$ ,  $\beta 3$ , and  $\beta 4$ ) and 1000 ng of P2X $_3$  plasmid WT or labeled with fluorescent protein. After 3 hours at 37°C, transfection medium was replaced with neuronal feeding medium.

### FRET analysis

Mouse E17 cortical neurons were transiently transfected with the indicated constructs on day 5 in culture, and measurements were made 1 to 3 days later. Before an imaging session, cell culture medium was replaced with phenol red to  $n_2$ -independent Leibovitz (L-15) medium (Invitrogen). FRET was analyzed by FLIM using a 60 $\times$  oil immersion objective on a C1si laser-scanning confocal microscope (Eclipse; Nikon) equipped with a 60r-scanning confocal microscope. Samples were scanned at a rate of 6  $\mu$ s per pixel for a 256  $\times$  256-pixel image. A 480-nm picosecond pulsed diode laser (PDL 800-D, PicoQuant GmbH) provided the excitation light (40 MHz), and emitted light was directed to a single-photon photomultiplier (SPCM-AQR SPAD, PerkinElmer). A time-correlated single-photon counting module and event timer (PicoHarp 300, PicoQuant GmbH) was used to record photon arrival times. Histograms of the time delay between the laser excitation pulse and photon arrival events were fit to exponential decays to extract fluorescence lifetimes for each pixel using PicoHarp 2.0, SymPhoTime software. The extracted lifetimes were used to determine the FRET efficiency ( $E$ ), where  $E = 1 - \tau_{\text{da}}/\tau_{\text{d}}$  ( $\tau_{\text{d}}$  is donor lifetime in the absence of the acceptor and  $\tau_{\text{da}}$  is donor lifetime in the presence of the acceptor). Binding fractions were determined from the coefficients of each exponential component in the fit.

### Nociceptive assays

**von Frey test.** In the strain survey, mice were tested on the von Frey test using the up-down staircase method of Dixon (68). Mice were placed on a metal mesh floor within small Plexiglas cubicles (9  $\times$  5  $\times$  5 cm high), and a set of eight calibrated von Frey fibers (Stoelting Touch Test Sensory Evaluator Kit #2 to #9; ranging from 0.007 to 1.40 g of force) were applied to the plantar surface of the hind paw until they bowed. The presence or absence of a withdrawal response in the next 3 s was scored, and the next fiber to be applied was determined. In all subsequent experiments, an automated von Frey test was used (Ugo Basile Dynamic Plantar Aesthesiometer). In this assay, pressure is gradually increased by the device until the mouse withdraws its hind paw; the maximal pressure at that point is displayed. We have found this method to feature less variability than

the up-down technique. Relative strain sensitivities are preserved using both methods (J. S. Mogil, unpublished data). In all experiments, measurements were taken in both ipsilateral and contralateral hind paws. Except in Fig. 1A, only ipsilateral hind paw responses are presented. There were no significant main effects of surgery, genotype, or drug on contralateral hind paw withdrawal thresholds in any experiment.

**Neuropathic surgeries.** After testing on two separate occasions (averaged) for baseline mechanical sensitivity as described above, some mice received experimental surgeries featuring damage to peripheral nerves serving the hind paw. In different studies, either the SNI (69, 70), SNL (71), or CCI (72) was used. In the SNI, we spared the sural nerve, and thus von Frey testing occurred on the lateral aspect of the hind paw. Mice were retested for mechanical sensitivity on post-operative days 1, 4, 7, 14, 21, and 28 in experiments evaluating allodynic severity, and on day 7 in experiments evaluating drug anti-allodynia. In the latter, (–)-nicotine (Sigma) was injected either systemically (0.15 to 1.8 mg/kg, intraperitoneally), intracerebroventricularly [2.5 to 25  $\mu$ g, (73)], intrathecally [1 to 17  $\mu$ g, (74)], or subcutaneously into the mid-plantar hind paw (25 to 100  $\mu$ g, intraplantar) immediately after “baseline” testing on day 7, and retested 15, 30, 45, and 60 min later. In the experiment shown in Fig. 5E, mice were retested 5, 15, and 30 min after intrathecal nicotine injection.

**Inflammatory assay.** After testing on two separate occasions (averaged) for baseline mechanical sensitivity as described above, mice were injected with CFA (50%; Sigma) into one hind paw. Mice were retested 3, 5, 7, 9, and 11 days after injection in experiments evaluating allodynic severity, and on day 3 in experiments evaluating drug anti-allodynia. In the latter, (–)-nicotine (Sigma) was injected either systemically (0.30 to 1.8 mg/kg, intraperitoneally), intracerebroventricularly [2.5 to 25  $\mu$ g, (73)], intrathecally [1 to 17  $\mu$ g, (74)], or subcutaneously into the mid-plantar hind paw (25 to 100  $\mu$ g, intraplantar) immediately after baseline testing on day 3, and retested 15, 30, 45, and 60 min later. In the experiment shown in Fig. 5E, mice were retested 5, 15, and 30 min after intrathecal nicotine injection.

**$\alpha,\beta$ me-ATP-induced pain behaviors.** In some experiments, mice pretreated 20 min earlier with nicotine (0 to 1.35 mg/kg, intraperitoneally) or A-317491 (300 nmol, intraplantar; Tocris Bioscience) were injected with 40 nmol of  $\alpha,\beta$ me-ATP (Tocris) into one hind paw, and nocifensive licking/biting behaviors were measured over the next 60 min by sampling the first 10 s of every 1-min time period. In other experiments, mice were tested for mechanical sensitivity as described above immediately before and 15 min after 40 nmol  $\alpha,\beta$ me-ATP (to confirm the presence of mechanical allodynia), followed immediately by systemic injection of nicotine (0.9 mg/kg) or A-317491 (300 nmol, intraplantar). Mechanical sensitivity was then measured at 15, 30, 45, 60, 90, and 120 min after drug.

**Pain test battery.** Details of the battery of acute and tonic assays are provided in Mogil *et al.* (65).

**Quantification of allodynia and anti-allodynia.** Allodynia over the multiple testing days was calculated as area over the withdrawal threshold  $\times$  time curve using the trapezoidal rule; percentage of maximum possible allodynia (% allodynia) was calculated for each mouse as compared to a hypothetical subject with the same baseline threshold and maximal allodynia (that is, a threshold of 0 g) at all post-surgery or post-CFA time points.

Drug anti-allodynia over 60 min was calculated as area under the curve using the trapezoidal rule, with respect to the pre-injury (pre-surgery or pre-CFA) baseline and the pre-drug (post-surgery or post-CFA)

baseline. Percentage of maximum possible anti-allodynia (% anti-allodynia) was calculated for each mouse as compared to a hypothetical subject with the same pre-injury and pre-drug baseline thresholds and complete resolution of allodynia at all post-drug time points.

## Human clinical cohorts

**Persistent post-herniotomy pain cohort.** This prospective cohort was composed of 429 Danish ( $n = 242$ ) and German ( $n = 187$ ) adult male patients of Caucasian origin (mean age, 55.1 years; SD, 13.3) who underwent open or laparoscopic transabdominal preperitoneal elective groin hernia repair (31). The main outcome for association analysis was the presence of moderate/severe postoperative 6-month pain (yes, 46.6%; no, 53.4%). There was no difference in pre-operative nociceptive function assessed by quantitative sensory testing between the Danish and the German cohort (31). Genotype-phenotype analysis was done using a prespecified regression equation, incorporating our assumption that one or two copies of the rare allele would affect the pain score in different genetic models, and adjusted by the following covariates: patients' age, surgery type, and activity assessment scale (AAS) score (0% if no pain-related activity impairment was reported, and 100% for maximum impairment) at baseline. All subjects donated a blood sample for DNA extraction; 14 samples could not be confidently assigned to a genotype. The study was approved by local ethics committees (Hørsholm Hospital, Denmark, and Centre for Minimal Invasive Surgery, Germany).

**TMD cohort.** Subjects were non-Hispanic white females ( $n = 159$ ), aged 18 to 60 years (mean, 36.8 years), recruited for a case-control study at the University of North Carolina (UNC) Orofacial Pain Clinic between 2005 and 2009. As described previously (32), TMD cases had to report facial pain for at least 5 days during the previous 2 weeks and be diagnosed with TMD arthralgia or myalgia during a standardized clinical examination that used the Research Diagnostic Criteria for TMD (75). Study participants who completed the CPSQ and provided blood for DNA extraction were included in this analysis. The CPSQ is a self-report questionnaire designed to ascertain the presence and characteristics of multiple pain symptoms and the lifetime presence of multiple pain conditions (33). To derive a single composite value representing pain of the head and neck, seven individual responses (duration of facial pain, intensity of current facial pain, intensity of greatest pain in the last 6 months, intensity of average pain over the last 6 months, primary headache characteristics, percentage of lifetime suffering from primary headache, and count of comorbid pain conditions) were normalized by conversion to  $z$  scores and then summed. All subjects provided signed informed consent for study procedures including blood draw and genotypic assessment, and this study was approved by the UNC Biomedical Institutional Review Board.

## Human genotyping

Genomic DNA was extracted from each blood sample using QIAamp DNA Blood Kit (Qiagen). Three tagging SNPs were identified within *CHRNA6* gene locus using the Haploview Tagger program: rs892413 [minor allele frequency (MAF) = 0.21], rs1072003 (MAF = 0.18), and rs7828365 (MAF = 0.12). Tagging SNPs were genotyped using the 5' nuclease method (76) and predesigned ABI SNP assays. Allele-specific signals were distinguished by measuring endpoint 6-FAM or VIC fluorescence intensities at 508 and 560 nm, respectively; genotypes were generated using StepOnePlus System Software (Applied Biosystems). The genotyping error rate was directly determined by re-genotyping



25% of the samples, randomly chosen, for each locus. Data cleaning and analysis were implemented using PLINK software v1.07 (77). Standard genotyping quality filters were imposed (call rate >95%, Hardy-Weinberg equilibrium  $P < 5 \times 10^{-5}$ ).

### Statistical analyses

Statistical analyses were conducted using an  $\alpha$  level of 0.05. Analysis of variance (ANOVA) or  $t$  tests were performed as appropriate after determining the normality of the experimental data (Shapiro-Wilk test), followed by Tukey's or Dunnett's post hoc tests, as appropriate. One-tailed testing was used where a priori expectations of direction of effect (for example, analgesia from a known analgesic compound like nicotine) existed. Analgesic ED<sub>50</sub>s and associated 95% confidence intervals were calculated using the method of Tallarida and Murray (78) as implemented by the FlashCalc 40.1 macro (M. H. Ossipov, University of Arizona). In expression and haplotype genomic mapping studies, multiple testing was controlled using false discovery rate.

Because of the small expected counts for TT homozygotes observed in the two human samples, permutation  $t$  tests were used to assess significance of genetic associations (79). In the case of the heritometry sample, where the response is binary, the usage of the  $t$  statistic in permutations is equivalent to the permutation test based on the  $\chi^2$  statistic (80). Association  $P$  values for two human studies were combined by a modification of the inverse normal method (81), where study-specific directional  $P$  values are combined and the result is converted to a two-sided  $P$  value.

### SUPPLEMENTARY MATERIALS

www.sciencetranslationalmedicine.org/cgi/content/full/7/287/287ra72/DC1

Fig. S1. Hind paw withdrawal thresholds to von Frey fiber stimulation measured before (–14, –7 days) and after (1, 4, 7, 14, 21, and 28 days) SNI surgery.

Fig. S2. Sex differences in *Chrna6* DRG mRNA expression in SM mice and their correlation with sex differences in mechanical allodynia.

Fig. S3. Down-regulation of *Chrna6* by nerve injury and correlation with mechanical allodynia.

Fig. S4. Increased ipsilateral mechanical allodynia in *Chrna6* ( $\alpha 6$  KO) but not *Chrna4* null mutants ( $\alpha 4$  KO).

Fig. S5. No altered sensitivity of *Chrna6* KO mice in a battery of acute and tonic nociceptive assays.

Fig. S6. Physical contacts between P2X3 and  $\alpha 6\beta 4^*$  nAChRs revealed by FLIM.

Table S1. Affymetrix gene expression data of all *Chrn*\* probes and correlation with baseline nociceptive sensitivity and mechanical allodynia.

Table S2. Top 10 correlated haplotypes (by  $P$  value), genome-wide, with SNI-induced mechanical allodynia in 25 mouse strains.

Table S3. Nicotine anti-allodynic ED<sub>50</sub>s in all genotypes.

Table S4. Dose-response characteristics (EC<sub>50</sub> and Hill coefficients; mean  $\pm$  SEM) for various combinations of  $\alpha 6$ -containing nicotinic receptors and P2X receptors expressed in oocytes.

### REFERENCES AND NOTES

- X. Li, P. Sahbaie, M. Zheng, J. Ritchie, G. Peltz, J. S. Mogil, J. D. Clark, Expression genetics identifies spinal mechanisms supporting formalin late phase behaviors. *Mol. Pain* **6**, 11 (2010).
- A.-K. Persson, M. Gebauer, S. Jordan, C. Metz-Weidmann, A. M. Schulte, H.-C. Schneider, D. Ding-Pfennigdorff, J. Thun, X.-J. Xu, Z. Wiesenfeld-Hallin, A. Darvasi, K. Fried, M. Devor, Correlational analysis for identifying genes whose regulation contributes to chronic neuropathic pain. *Mol. Pain* **5**, 7 (2009).
- A.-K. Persson, X.-J. Xu, Z. Wiesenfeld-Hallin, M. Devor, K. Fried, Expression of DRG candidate pain molecules after nerve injury—A comparative study among five inbred mouse strains with contrasting pain phenotypes. *J. Peripher. Nerv. Syst.* **15**, 26–39 (2010).

- J. Nissenbaum, M. Devor, Z. Seltzer, M. Gebauer, M. Michaelis, M. Tal, R. Dorfman, M. Abitbul-Yarkon, Y. Lu, T. Elahipanah, S. delCanho, A. Minert, K. Fried, A.-K. Persson, H. Shpigler, E. Shabo, B. Yakir, A. Pisante, A. Darvasi, Susceptibility to chronic pain following nerve injury is genetically affected by *CACNG2*. *Genome Res.* **20**, 1180–1190 (2010).
- M. Vincler, Neuronal nicotinic receptors as targets for novel analgesics. *Expert Opin. Investig. Drugs* **14**, 1191–1198 (2005).
- T. Young, S. Wittenauer, J. M. McIntosh, M. Vincler, Spinal  $\alpha 3\beta 2^*$  nicotinic acetylcholine receptors tonically inhibit the transmission of nociceptive mechanical stimuli. *Brain Res.* **1229**, 118–124 (2008).
- S. D. AlSharari, K. Freitas, M. I. Damaj, Functional role of  $\alpha 7$  nicotinic receptor in chronic neuropathic and inflammatory pain: Studies in transgenic mice. *Biochem. Pharmacol.* **86**, 1201–1207 (2013).
- J. M. McIntosh, N. Absalom, M. Chebib, A. B. Elgoyhen, M. Vincler,  $\alpha 9$  nicotinic acetylcholine receptors and the treatment of pain. *Biochem. Pharmacol.* **78**, 693–702 (2009).
- S. Mohammadi, M. J. Christie,  $\alpha 9$ -Nicotinic acetylcholine receptors contribute to the maintenance of chronic mechanical hyperalgesia, but not thermal or mechanical allodynia. *Mol. Pain* **10**, 64 (2014).
- R. M. Drenan, S. R. Grady, P. Whiteaker, T. McClure-Begley, S. McKinney, J. M. Miwa, S. Bupp, N. Heintz, J. M. McIntosh, M. Bencherif, M. J. Marks, H. A. Lester, In vivo activation of midbrain dopamine neurons via sensitized, high-affinity  $\alpha 6^*$  nicotinic acetylcholine receptors. *Neuron* **60**, 123–136 (2008).
- E. D. W. Mackey, S. E. Engle, M. R. Kim, H. C. O'Neill, C. R. Wageman, N. E. Patzlaff, Y. Wang, S. R. Grady, J. M. McIntosh, M. J. Marks, H. A. Lester, R. M. Drenan,  $\alpha 6^*$  Nicotinic acetylcholine receptor expression and function in a visual salience circuit. *J. Neurosci.* **32**, 10226–10237 (2012).
- A. J. Hone, E. L. Meyer, M. McIntyre, J. M. McIntosh, Nicotinic acetylcholine receptors in dorsal root ganglion neurons include the  $\alpha 6\beta 4^*$  subtype. *FASEB J.* **26**, 917–926 (2011).
- L. Liu, G. Q. Chang, Y. Q. Jiao, S. A. Simon, Neuronal nicotinic acetylcholine receptors in rat trigeminal ganglia. *Brain Res.* **809**, 238–245 (1998).
- C. J. Keiger, J. C. Walker, Individual variation in the expression profiles of nicotinic receptors in the olfactory bulb and trigeminal ganglion and identification of  $\alpha 2$ ,  $\alpha 6$ ,  $\alpha 9$ , and  $\beta 3$  transcripts. *Biochem. Pharmacol.* **59**, 233–240 (2000).
- J. R. Genzen, W. Van Cleve, D. S. McGehee, Dorsal root ganglion neurons express multiple nicotinic acetylcholine receptor subtypes. *J. Neurophysiol.* **86**, 1773–1782 (2001).
- R. E. Sorge, M. L. LaCroix-Fralish, A. H. Tuttle, S. G. Sotocinal, J.-S. Austin, J. Ritchie, M. L. Chanda, A. C. Graham, L. Topham, S. Beggs, M. W. Salter, J. S. Mogil, Spinal cord Toll-like receptor 4 mediates inflammatory and neuropathic hypersensitivity in male but not female mice. *J. Neurosci.* **31**, 15450–15454 (2011).
- P. McClurg, J. Janes, C. Wu, D. L. Delano, J. R. Walker, S. Batalov, J. S. Takahashi, K. Shimomura, A. Kohsaka, J. Bass, T. Wiltshire, A. I. Su, Genomewide association analysis in diverse inbred mice: Power and population structure. *Genetics* **176**, 675–683 (2007).
- G. O. Dussor, A. S. Leong, N. B. Gracia, S. Kilo, T. J. Price, K. M. Hargreaves, C. M. Flores, Potentiation of evoked calcitonin gene-related peptide release from oral mucosa: A potential basis for the pro-inflammatory effects of nicotine. *Eur. J. Neurosci.* **18**, 2515–2526 (2003).
- N. Champiaux, Z.-Y. Han, A. Bessis, F. M. Rossi, M. Zoli, L. Marubio, J. M. McIntosh, J.-P. Changeux, Distribution and pharmacology of  $\alpha 6$ -containing nicotinic acetylcholine receptors analyzed with mutant mice. *J. Neurosci.* **22**, 1208–1217 (2002).
- B. S. Khakh, J. A. Fisher, R. Nashmi, D. N. Bowser, H. A. Lester, An angstorn scale interaction between plasma membrane ATP-gated P2X<sub>2</sub> and  $\alpha 6\beta 2$  nicotinic channels measured with fluorescence resonance energy transfer and total internal reflection fluorescence microscopy. *J. Neurosci.* **25**, 6911–6920 (2005).
- B. S. Khakh, X. Zhou, J. Sydes, J. J. Galligan, H. A. Lester, State-dependent cross-inhibition between transmitter-gated cation channels. *Nature* **406**, 405–410 (2000).
- D. A. Decker, J. J. Galligan, Molecular mechanisms of cross-inhibition between nicotinic acetylcholine receptors and P2X receptors in myenteric neurons and HEK-293 cells. *Neurogastroenterol. Motil.* **22**, 901–908 (2010).
- V. Souslova, P. Cesare, Y. Ding, A. N. Akopian, L. C. Stanfa, R. Suzuki, K. Carpenter, A. H. Dickenson, S. Boyce, R. Hill, D. Nebenius-Oosthuizen, A. J. H. Smith, E. J. Kidd, J. N. Wood, Warm-coding deficits and aberrant inflammatory pain in mice lacking P2X<sub>2</sub> receptors. *Nature* **407**, 1015–1017 (2000).
- D. A. Cockayne, S. G. Hamilton, Q.-M. Zhu, P. M. Dunn, Y. Zhong, S. Novakovic, A. B. Malmberg, G. Cain, A. Berson, L. Kassotakis, L. Hedley, W. G. Lachnit, G. Burnstock, S. B. McMahon, A. P. D. W. Ford, Urinary bladder hyporeflexia and reduced pain-related behaviour in P2X<sub>2</sub>-deficient mice. *Nature* **407**, 1011–1015 (2000).
- S. Dash, M. Bhakta, Y. Chang, R. J. Lukas, Identification of N-terminal extracellular domain determinants in nicotinic acetylcholine receptor (nAChR)  $\alpha 6$  subunits that influence effects of wild-type or mutant  $\beta 3$  subunits on function of  $\alpha 6\beta 2^*$ - or  $\alpha 6\beta 4^*$ -nAChR. *J. Biol. Chem.* **286**, 37976–37989 (2011).
- C. Xiao, R. Srinivasan, R. M. Drenan, E. D. Mackey, J. M. McIntosh, H. A. Lester, Characterizing functional  $\alpha 6\beta 2$  nicotinic acetylcholine receptors in vitro: Mutant  $\beta 2$  subunits improve membrane expression, and fluorescent proteins reveal responsive cells. *Biochem. Pharmacol.* **82**, 852–861 (2011).

27. C. Virginio, R. A. North, A. Suprenant, Calcium permeability and block at homomeric and heteromeric P2X<sub>2</sub> and P2X<sub>3</sub> receptors, and P2X receptors in rat nodose neurones. *J. Physiol.* **510** (Pt. 1), 27–35 (1998).
28. W. Limapichat, D. A. Dougherty, H. A. Lester, Subtype-specific mechanisms for functional interaction between  $\alpha 6\beta 4^*$  nicotinic acetylcholine receptors and P2X receptors. *Mol. Pharmacol.* **86**, 263–274 (2014).
29. E. B. Pratt, T. S. Brink, P. Bergson, M. M. Voigt, S. P. Cook, Use-dependent inhibition of P2X<sub>3</sub> receptors by nanomolar agonist. *J. Neurosci.* **25**, 7359–7365 (2005).
30. Y. Chen, X. Zhang, C. Wang, G. Li, Y. Gu, L.-Y. M. Huang, Activation of P2X7 receptors in glial satellite cells reduces pain through downregulation of P2X3 receptors in nociceptive neurons. *Proc. Natl. Acad. Sci. U.S.A.* **105**, 16773–16778 (2008).
31. E. K. Aasvang, E. Gmaehle, J. B. Hansen, B. Gmaehle, J. L. Forman, J. Schwarz, R. Bittner, H. Kehlet, Predictive risk factors for persistent postherniotomy pain. *Anesthesiology* **112**, 957–969 (2010).
32. G. D. Slade, M. S. Conrad, L. Diatchenko, N. U. Rashid, S. Zhong, S. Smith, J. Rhodes, A. Medvedev, S. Makarov, W. Maixner, A. G. Nackley, Cytokine biomarkers and chronic pain: Association of genes, transcription, and circulating proteins with temporomandibular disorders and widespread palpation tenderness. *Pain* **152**, 2802–2812 (2011).
33. R. Ohrbach, R. B. Fillingim, F. Mulkey, Y. Gonzalez, S. Gordon, H. Gremillion, P. F. Lim, M. Riberio-Dasilva, J. D. Greenspan, C. Knott, W. Maixner, G. Slade, Clinical findings and pain symptoms as potential risk factors for chronic TMD: Descriptive data and empirically identified domains from the OPPERA case-control study. *J. Pain* **12** (Suppl. 11), T27–T45 (2011).
34. J. J. Cox, F. Reimann, A. K. Nicholas, G. Thornton, E. Roberts, K. Springell, G. Karbani, H. Jafri, J. Mannan, Y. Raashid, L. Al-Gazali, H. Hamamy, E. M. Valente, S. Gorman, R. Williams, D. P. McHale, J. N. Wood, F. M. Gribble, C. G. Woods, An *SCN9A* channelopathy causes congenital inability to experience pain. *Nature* **444**, 894–898 (2006).
35. E. Leipold, L. Liebmann, G. C. Korenke, T. Heinrich, S. Giesselmann, J. Baets, M. Ebbinghaus, R. O. Goral, T. Stodberg, J. C. Hennings, M. Bergmann, J. Altmüller, H. Thiele, A. Wetzel, P. Numborg, V. Timmerman, P. de Jonghe, R. Blum, H.-G. Schaible, J. Weis, S. H. Heinemann, C. A. Hubner, I. Kurth, A de novo gain-of-function mutation in *SCN11A* causes loss of pain perception. *Nat. Genet.* **45**, 1399–1404 (2013).
36. J. W. Daly, Nicotinic agonists, antagonists, and modulators from natural sources. *Cell. Mol. Neurobiol.* **25**, 513–552 (2005).
37. A. W. Bannon, M. W. Decker, M. W. Holladay, P. Curzon, D. Donnelly-Roberts, P. S. Puttfarcken, R. S. Bitner, A. Diaz, A. H. Dickenson, R. D. Porsolt, M. Williams, S. P. Americ, Broad-spectrum, non-opioid analgesic activity by selective modulation of neuronal nicotinic acetylcholine receptors. *Science* **279**, 77–81 (1998).
38. S. P. Americ, M. Holladay, M. Williams, Neuronal nicotinic receptors: A perspective on two decades of drug discovery research. *Biochem. Pharmacol.* **74**, 1092–1101 (2007).
39. L. M. Marubio, M. del Mar Arroyo-Jimenez, M. Cordero-Erausquin, C. Lena, N. Le Novere, A. de Kerchove d'Exaerde, M. Huchet, M. I. Damaj, J.-P. Changeux, Reduced antinociception in mice lacking neuronal nicotinic receptor subunits. *Nature* **398**, 805–810 (1999).
40. M. I. Damaj, M. Fei-Yin, M. Dukat, W. Glassco, R. A. Glennon, B. R. Martin, Antinociceptive responses to nicotinic acetylcholine receptor ligands after systemic and intrathecal administration in mice. *J. Pharmacol. Exp. Ther.* **283**, 1058–1065 (1998).
41. K. J. Jackson, M. J. Marks, R. E. Vann, X. Chen, T. F. Gamage, J. A. Warner, M. I. Damaj, Role of  $\alpha 5$  nicotinic acetylcholine receptors in pharmacological and behavioral effects of nicotine in mice. *J. Pharmacol. Exp. Ther.* **334**, 134–146 (2010).
42. M. I. Damaj, C. Fonck, M. J. Marks, P. Deshpande, C. Labarca, H. A. Lester, A. C. Collins, B. R. Martin, Genetic approaches identify differential roles for  $\alpha 6\beta 2^*$  nicotinic receptors in acute models of antinociception in mice. *J. Pharmacol. Exp. Ther.* **321**, 1161–1169 (2007).
43. B. Gao, M. Hierl, K. Clarkin, T. Juan, H. Nguyen, M. Valk, H. Deng, W. Guo, S. G. Lehto, D. Matson, J. S. McDermott, J. Knop, K. Gaida, L. Cao, D. Waldon, B. K. Albrecht, A. A. Boezio, K. W. Copeland, J. C. Harmange, S. K. Springer, A. B. Malmbert, S. I. McDonough, Pharmacological effects of non-selective and subtype-selective nicotinic acetylcholine receptor agonists in animal models of persistent pain. *Pain* **149**, 33–49 (2010).
44. D. J. Rowbotham, A. Arslanian, W. Nothaft, W. R. Duan, A. E. Best, Y. Pritchett, Q. Zhou, B. R. Stacey, Efficacy and safety of the  $\alpha 4\beta 2$  neuronal nicotinic receptor agonist ABT-894 in patients with diabetic peripheral neuropathic pain. *Pain* **153**, 862–868 (2012).
45. R. Hausmann, G. Bahrenberg, D. Kuhlmann, M. Schumacher, U. Braam, D. Bieler, I. Schlusche, G. Schmalzing, A hydrophobic residue in position 15 of the rP2X<sub>3</sub> receptor slows desensitization and reveals properties beneficial for pharmacological analysis and high-throughput screening. *Neuropharmacology* **79**, 603–615 (2014).
46. L. Vulchanova, M. S. Riedl, S. J. Shuster, G. Buell, A. Suprenant, R. A. North, R. Elde, Immunohistochemical study of the P2X<sub>2</sub> and P2X<sub>3</sub> receptor subunits in rat and monkey sensory neurons and their central terminals. *Neuropharmacology* **36**, 1229–1242 (1997).
47. D. A. Cockayne, P. M. Dunn, Y. Zhong, W. Rong, S. G. Hamilton, G. E. Knight, H.-Z. Ruan, B. Ma, P. Yip, P. Nunn, S. B. McMahon, G. Burnstock, A. P. D. W. Ford, P2X<sub>2</sub> knockout mice and P2X<sub>2</sub>/P2X<sub>3</sub> double knockout mice reveal a role for the P2X<sub>2</sub> receptor subunit in mediating multiple sensory effects of ATP. *J. Physiol.* **567**, 621–639 (2005).
48. S. Beggs, T. Trang, M. W. Salter, P2X<sub>4R+</sub> microglia drive neuropathic pain. *Nat. Neurosci.* **15**, 1068–1073 (2012).
49. R. E. Sorge, T. Trang, R. Dorfman, S. B. Smith, S. Beggs, J. Ritchie, J.-S. Austin, D. V. Zaykin, H. Vander Meulen, M. Costigan, T. A. Herbert, M. Yarkoni-Abitbul, D. Tichauer, J. Livneh, E. Gershon, M. Zheng, K. Tan, S. L. John, G. D. Slade, J. Jordan, C. J. Woolf, G. Peltz, W. Maixner, L. Diatchenko, Z. Seltzer, M. W. Salter, J. S. Mogil, Genetically determined P2X7 receptor pore formation regulates variability in chronic pain sensitivity. *Nat. Med.* **18**, 595–599 (2012).
50. P. Honore, J. Mikusa, B. Bianchi, H. McDonald, J. Cartmell, C. Faltynek, M. F. Jarvis, TNP-ATP, a potent P2X<sub>3</sub> receptor antagonist, blocks acetic acid-induced abdominal constriction in mice: Comparison with reference analgesics. *Pain* **96**, 99–105 (2002).
51. Y. Wang, X. Zhang, Q. L. Guo, W. Y. Zou, C. S. Huang, J. Q. Yan, Cyclooxygenase inhibitors suppress the expression of P2X<sub>3</sub> receptors in the DRG and attenuate hyperalgesia following chronic constriction injury in rats. *Neurosci. Lett.* **478**, 77–81 (2010).
52. M. E. Kiyatkin, B. Feng, E. S. Schwartz, G. F. Gebhart, Combined genetic and pharmacological inhibition of TRPV1 and P2X3 attenuates colorectal hypersensitivity and afferent sensitization. *Am. J. Physiol. Gastrointest. Liver Physiol.* **305**, G638–G648 (2013).
53. N. Noma, M. Shinoda, K. Honda, M. Kiyomoto, K. Dezawa, Y. Nakaya, O. Komiyama, Y. Imamura, K. Iwata, Interaction of IL-1 $\beta$  and P2X<sub>3</sub> receptor in pathologic masseter muscle pain. *J. Dent. Res.* **92**, 456–460 (2013).
54. M. Liu, H. Yang, D. Fang, J. J. Yang, J. Cai, Y. Wan, D. H. Chui, J. S. Han, G. G. Xing, Upregulation of P2X3 receptors by neuronal calcium sensor protein VILIP-1 in dorsal root ganglions contributes to the bone cancer pain in rats. *Pain* **154**, 1551–1568 (2013).
55. S. K. Hullugundi, M. D. Ferrari, A. M. Van den Maagdenberg, A. Nistri, The mechanism of functional up-regulation of P2X3 receptors of trigeminal sensory neurons in a genetic mouse model of familial hemiplegic migraine type 1 (FHM-1). *PLOS One* **8**, e60677 (2013).
56. I. Wessler, H. Kilbinger, F. Bittinger, R. Unger, C. J. Kirkpatrick, The non-neuronal cholinergic system in humans: Expression, function and pathophysiology. *Life Sci.* **72**, 2055–2061 (2003).
57. G. Wu, G. T. Whiteside, G. Lee, S. Nolan, M. Niosi, M. S. Pearson, V. I. Ilyin, A-317491, a selective P2X<sub>2</sub>/P2X<sub>3</sub> receptor antagonist, reverses inflammatory mechanical hyperalgesia through action at peripheral receptors in rats. *Eur. J. Pharmacol.* **504**, 45–53 (2004).
58. J. S. Mogil, Animal models of pain: Progress and challenges. *Nat. Rev. Neurosci.* **10**, 283–294 (2009).
59. D. Zhang, T. Bordia, M. McGregor, J. M. McIntosh, M. W. Decker, M. Quik, ABT-089 and ABT-894 reduce levodopa-induced dyskinesias in a monkey model of Parkinson's disease. *Mov. Disord.* **29**, 508–517 (2014).
60. Y. Wang, J. W. Lee, G. Oh, S. R. Grady, J. M. McIntosh, D. H. Brunzell, J. R. Cannon, R. M. Drenan, Enhanced synthesis and release of dopamine in transgenic mice with gain-of-function  $\alpha 6^*$  nAChRs. *J. Neurochem.* **129**, 315–327 (2014).
61. M. De Biasi, J. A. Dani, Reward, addiction, withdrawal to nicotine. *Annu. Rev. Neurosci.* **34**, 105–130 (2011).
62. A. H. Rasmussen, D. Strobaek, T. Dyhring, M. L. Jensen, D. Peters, M. Grunnet, D. B. Timmermann, P. K. Ahning, Biophysical and pharmacological characterization of  $\alpha 6$ -containing nicotinic acetylcholine receptors expressed in HEK293 cells. *Brain Res.* **1542**, 1–11 (2014).
63. A. J. Hone, M. Ruiz, M. Scadden, S. Christensen, J. Gajewiak, L. Azam, J. M. McIntosh, Positional scanning mutagenesis of  $\alpha$ -conotoxin PeIA identifies critical residues that confer potency and selectivity for  $\alpha 6\alpha 3\beta 2\beta 3$  and  $\alpha 3\beta 2$  nicotinic acetylcholine receptors. *J. Biol. Chem.* **288**, 25428–25439 (2013).
64. D. H. Brunzell, Preclinical evidence that activation of mesolimbic alpha 6 subunit containing nicotinic acetylcholine receptors supports nicotine addiction phenotype. *Nicotine Tob. Res.* **14**, 1258–1269 (2012).
65. J. S. Mogil, J. Ritchie, S. G. Sotocinal, S. B. Smith, S. Croteau, D. J. Levitin, A. K. Naumova, Screening for pain phenotypes: Analysis of three congenic mouse strains on a battery of nine nociceptive assays. *Pain* **126**, 24–34 (2006).
66. S. A. Ross, J. Y. Wong, J. J. Clifford, A. Kinsella, J. S. Massalas, M. K. Horne, I. E. Scheffer, I. Kola, J. L. Waddington, S. F. Berkovic, J. Drago, Phenotypic characterization of an  $\alpha 4$  neuronal nicotinic acetylcholine receptor subunit knock-out mouse. *J. Neurosci.* **20**, 6431–6441 (2000).
67. C. Wu, D. L. Delano, N. Mitro, S. V. Su, J. Janes, P. McClurg, S. Batalov, G. L. Welch, J. Zhang, A. P. Orth, J. R. Walker, R. J. Glynn, M. P. Cooke, J. S. Takahashi, K. Shimomura, A. Kohsaka, J. Bass, E. Saez, T. Wiltshire, A. I. Su, Gene set enrichment in eQTL data identifies novel annotations and pathway regulators. *PLOS Genet.* **4**, e1000070 (2008).
68. S. R. Chaplan, F. W. Bach, J. W. Pogrel, J. M. Chung, T. L. Yaksh, Quantitative assessment of tactile allodynia evoked by unilateral ligation of the fifth and sixth lumbar nerves in the rat. *J. Neurosci. Methods* **53**, 55–63 (1994).
69. I. Decosterd, C. J. Woolf, Spared nerve injury: An animal model of persistent peripheral neuropathic pain. *Pain* **87**, 149–158 (2000).
70. S. D. Shields, W. A. Eckert III, A. I. Basbaum, Spared nerve injury model of neuropathic pain in the mouse: A behavioral and anatomic analysis. *J. Pain* **4**, 465–470 (2003).
71. S. H. Kim, J. M. Chung, An experimental model for peripheral neuropathic produced by segmental spinal nerve ligation in the rat. *Pain* **50**, 355–363 (1992).
72. G. J. Bennett, Y.-K. Xie, A peripheral mononeuropathy in rat that produces disorders of pain sensation like those seen in man. *Pain* **33**, 87–107 (1988).

73. S. E. Laursen, J. K. Belknap, Intracerebroventricular injections in mice. Some methodological refinements. *J. Pharmacol. Methods* **16**, 355–357 (1986).
74. J. L. K. Hyliden, G. L. Wilcox, Intrathecal morphine in mice: A new technique. *Eur. J. Pharmacol.* **67**, 313–316 (1980).
75. S. Dworkin, L. LeResche, Research diagnostic criteria for temporomandibular disorders: Review, criteria, examinations and specifications, critique. *J. Craniomandib. Disord.* **6**, 301–355 (1992).
76. M. M. Shi, S. P. Myrand, M. R. Bleavins, F. A. de la Iglesia, High throughput genotyping for the detection of a single nucleotide polymorphism in NAD(P)H quinine oxidoreductase (DT diaphoresis) using TaqMan probes. *Mol. Pathol.* **52**, 295–299 (1999).
77. S. Purcell, B. Neale, K. Todd-Brown, L. Thomas, M. A. R. Ferreira, D. Bender, J. Maller, P. Sklar, P. I. W. de Bakker, M. J. Daly, P. C. Sham, PLINK: A toolset for whole-genome association and population-based linkage analysis. *Am. J. Hum. Genet.* **81**, 559–575 (2007).
78. R. J. Tallarida, R. B. Murray, *Manual of Pharmacologic Calculation* (Springer-Verlag, New York, 1981).
79. P. I. Good, *Permutation Tests: A Practical Guide to Resampling Methods for Testing Hypotheses* (Springer, New York, 2000).
80. R. B. D'Agostino, Relation between chi-squared and ANOVA tests for testing equality of  $k$  independent dichotomous populations. *Am. Statist.* **26**, 30–32 (1972).
81. D. V. Zaykin, Optimally weighted Z-test is a powerful method for combining probabilities in meta-analysis. *J. Evol. Biol.* **24**, 1836–1841 (2011).

**Acknowledgments:** We thank T. Earley, T. Miyamoto, and M. Petrus for assistance with DRG dissection. We thank S. Ranade and V. Uzzell for data analysis. **Funding:** Supported by the

Canadian Institutes for Health Research and the Louise and Alan Edwards Foundation (J.S.M.) and the NIH (D.A.D., H.A.L., and A.P.). **Author contributions:** The study was conceived by A.P. and J.S.M. and designed by J.S.W., L.D., I.B., M.I.D., H.A.L., A.P., and J.S.M. J.S.W., J. Mathur, W.L., M.R.P., M.A.-Q., R.E.S., L.J.M., K.F., J.-S.A., J.Z., J. Marcovitz, A.H.T., P.M.S., S.C., J.J., S.A.S., E.K.A., R.B., C.I.R., H.K., and J.W. collected data. D.V.Z., S.B.S., F.D., R.M.D., S.K.S., W.L., G.D.S., and A.I.S. analyzed data. U.M., J.-P.C., and M.D. provided reagents or data. W.M., L.D., I.B., D.A.D., S.C.R.L., M.I.D., H.A.L., A.P., and J.S.M. supervised the collection of data, contributed to its interpretation, and edited the manuscript. J.S.W. and J.S.M. wrote the manuscript. **Competing interests:** The authors declare that they have no competing interests. **Data and materials availability:** The expression genomics data for this study have been deposited into the GeneNetwork Database ([www.genenetwork.org](http://www.genenetwork.org)).

Submitted 7 July 2014

Accepted 11 March 2015

Published 13 May 2015

10.1126/scitranslmed.3009986

**Citation:** J. S. Wieskopf, J. Mathur, W. Limapichat, M. R. Post, M. Al-Qazzaz, R. E. Sorge, L. J. Martin, D. V. Zaykin, S. B. Smith, K. Freitas, J.-S. Austin, F. Dai, J. Zhang, J. Marcovitz, A. H. Tuttle, P. M. Slepian, S. Clarke, R. M. Drenan, J. Janes, S. Al Sharari, S. K. Segall, E. K. Aasvang, W. Lai, R. Bittner, C. I. Richards, G. D. Slade, H. Kehlet, J. Walker, U. Maskos, J.-P. Changeux, M. Devor, W. Maixner, L. Diatchenko, I. Belfer, D. A. Dougherty, A. I. Su, S. C. R. Lummis, M. Imad Damaj, H. A. Lester, A. Patapoutian, J. S. Mogil, The nicotinic  $\alpha 6$  subunit gene determines variability in chronic pain sensitivity via cross-inhibition of P2X2/3 receptors. *Sci. Transl. Med.* **7**, 287ra72 (2015).