**Behavioral/Systems/Cognitive**

**BAMBI (Bone Morphogenetic Protein and Activin Membrane-Bound Inhibitor) Reveals the Involvement of the Transforming Growth Factor-β Family in Pain Modulation**

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Transforming growth factors-β (TGF-βs) signal through type I and type II serine–threonine kinase receptor complexes. During ligand binding, type II receptors recruit and phosphorylate type I receptors, triggering downstream signaling. BAMBI [bone morphogenetic protein (BMP) and activin membrane-bound inhibitor] is a transmembrane pseudoreceptor structurally similar to type I receptors but lacks the intracellular kinase domain. BAMBI modulates negatively pan-TGF-β family signaling; therefore, it can be used as an instrument for unraveling the roles of these cytokines in the adult CNS. BAMBI is expressed in regions of the CNS involved in pain transmission and modulation. The lack of BAMBI in mutant mice resulted in increased levels of TGF-β signaling activity, which was associated with attenuation of acute pain behaviors, regardless of the modality of the stimuli (thermal, mechanical, chemical/inflammatory). The nociceptive hyposensitivity exhibited by *BAMBI−/−* mice was reversed by the opioid antagonist naloxone. Moreover, in a model of chronic neuropathic pain, the alldynic responses of *BAMBI−/−* mice also appeared attenuated through a mechanism involving δ-opioid receptor signaling. Basal mRNA and protein levels of precursor proteins of the endogenous opioid peptides *proopiomelanocortin* (POMC) and *proenkephalin* (PENK) appeared increased in the spinal cords of *BAMBI−/−*. Transcript levels of TGF-βs and their intracellular effectors correlated directly with genes encoding opioid peptides, whereas BAMBI correlated inversely. Furthermore, incubation of spinal cord explants with activin A or BMP-7 increased POMC and/or PENK mRNA levels. Our findings identify TGF-β family members as modulators of acute and chronic pain perception through the transcriptional regulation of genes encoding the endogenous opioids.

**Introduction**

The transforming growth factor-β (TGF-β) superfamily of cytokines includes activins, TGF-βs, and bone morphogenetic proteins (BMPs). TGF-βs signal through heteromeric combinations of type I receptors, also termed activin-like kinases (ALK), and type II receptors, both of which are serine/threonine kinases (Wrana et al., 1994). Ligand binding leads to downstream propagation of the signal through canonical phosphorylation of Smad proteins, which are translocated into the nucleus to regulate the transcription of TGF-β target genes (Feng and Derynck, 2005; Massagué et al., 2005), as well as through noncanonical signaling, notably via TGF-β-activated kinase 1 (TAK1) (Yamaguchi et al., 1995; Zhang, 2009). TGF-β signaling is modulated by BAMBI [BMP and activin membrane-bound inhibitor], a transmembrane protein that is structurally similar to type I receptors but lacks the intracellular kinase domain. BAMBI acts as a pseudoreceptor that inhibits pan-TGF-β family signaling by preventing the formation of active receptor complexes (Onichtchouk et al., 1999; Yan et al., 2009). BAMBI can also function as a positive regulator of the Wnt/β-catenin pathway (Lin et al., 2008).

Little is known about the physiological functions regulated by BAMBI or the pathological consequences of an imbalance between BAMBI and TGF-βs. During embryogenesis, BAMBI would act as a negative feedback regulator of BMP signaling (Onichtchouk et al., 1999; Tsang et al., 2000), in a context of cooperative function within the BMP-4 synexpression group (Grotewold et al., 2001; Karaunava et al., 2004). However, deletion of the BAMBI gene has no apparent phenotypic consequences affecting embryonic development (Chen et al., 2007). BAMBI deregulation has been involved in the pathogenesis of various human pathologies, such as tumor progression and metastatic capability (Togo et al., 2008; Fritzmann et al., 2009; Khin et al., 2009), pathological fibrosis (Seki et al., 2007; Harada et al., 2009), and metastatic bone formation (Kitazawa et al., 2005).
The biological processes regulated by the TGF-β family of cytokines in the rodent developing nervous system have been extensively studied (Böttner et al., 2000). In the adult CNS, more or less strong signals for several members and receptors of the TGF-β family are still detectable (Funaba et al., 1997; Meiller et al., 1997; Böttner et al., 2000; Unsicker and Strelau, 2000; Chen et al., 2003; Fan et al., 2003; Mikawa et al., 2006), supporting a role in the regulation of nociceptive processing under physiological and pathological conditions. Herein, we show that BAMBII is expressed in mice nociception-relevant areas and that targeted deletion of BAMBII results in attenuation of both acute pain behaviors and neuropathic pain development after nerve injury. The mechanism involves the transcriptional regulation by members of the TGF-β family of genes encoding endogenous opioid peptides.

Materials and Methods

Animals

The experiments were performed in male mice, housed in a room kept at 22°C with and exposed for their whole lifespan to an inverted 12 h light/dark cycle (dark from 8:00 A.M. to 8:00 P.M.). Food and water were provided ad libitum. This study was approved by the Cantabria University Institutional Laboratory Animal Care and Use Committee, and the experiments were performed in accordance with the Declaration of Helsinki and the European Communities Council Directive (86/609/EEC) and the guidelines of the Committee for Research and Ethical Issues of International Association for the Study of Pain (Zimmermann, 1983). Every effort was made to minimize the number of animals used and their suffering.

Generation of BAMBII−/− mice

Homologous recombination in embryonic stem cells was used to create a 129Sv × C57BL/6 mouse line, in which exons 2 and 3 of the BAMBII gene were deleted (Fig. 1A). The remaining exon 1 contains the 5′ untranslated region and the first 14 amino acids, which are part of the signaling sequence characteristic of membrane-targeted or -secreted proteins. Mutant mice were backcrossed with C57BL/6 up to the F6 generation. Interbreeding of BAMBII−/− mice resulted in homozygous mutant mice. The experiments were performed in adult (14–16 weeks) BAMBII+/−, BAMBII−/−, and BAMBII−−/− mice. Genotyping was performed on genomic DNA extracted from tissue punched from the ear by PCR, following standard protocols with TaqDNA Polymerase (Invitrogen) (Fig. 1B). The primers were designed to amplify a 398 bp fragment containing parts of the deleted region (forward, TGTGAT-AGCGGTTCCCATTGC; reverse, CCAGATA- AAAGTGTCGCTTGACG) for the wild-type allele and a 524 bp fragment containing part of the neomycin cassette (forward, 5′-TTTCGCAATGCAGACGGGTGG-3′; reverse, 5′-GGACAAAAAGACCCCTGGGAAG-3′) for the targeted allele. The existence of BAMBII transcripts in the CNS of wild-type mice as well as its absence in BAMBII−/− mice was confirmed by reverse transcription (RT)-PCR analyses (Fig. 1C). An increase in TGF-β1 family-mediated signaling in mutant mice, compared with their wild-type littermates, was reflected by the increased expression of phosphorylated Smad-1 and Smad-2 proteins observed in the brain cortex and spinal cord by Western blot analyses (Fig. 1D).

Behavioral studies

Behavioral studies were performed in 14-week-old mice (n = 8–12 per group) by observers blinded to the genotypes of the animals. Mice sensorimotor functions were evaluated in a battery of tests assessing motor activity, sensorimotor reflexes, equilibrium, and coordination as described previously (Martínez-Cué et al., 2005). Mice performed the behavioral tests during their activity (dark) phase, under dim red light.

Acute pain behavior studies

The warm-water tail-flick test was performed by immersing the tip of the tail (2–3 cm) into the water. The time it took to remove the tail from the water was measured (cutoff time, 60 s). In the hotplate test, mice were placed on a 50 or 52°C hotplate, and the delay to hindpaw licking and jumping were recorded (cutoff time, 120 s). In the Formalin test, mice received a 20 μl intraplantar injection of a 2% Formalin solution in the left hindpaw, and the time spent licking the paw was recorded within the first 5 min (first phase) and from 20 to 30 min after injection (second phase). The mechanical withdrawal threshold test was performed by applying an ascending series of calibrated von Frey hairs to the plantar surface of each hindpaw until a withdrawal response was observed. The results are expressed as percentage of paw withdrawals in response to stimuli of increasing strength (0.1–10 g) applied five times.
Chronic neuropathic pain behavior studies

Sciatic nerve crush injury. To induce the development of neuropathic pain, mice were subjected to sciatic nerve crush injury (Bester et al., 2000) under inhalation anesthesia with isoflurane (induction, 3%; surgery, 1.5%). The left common sciatic nerve was exposed via blunt dissection through the biceps femoris muscle. The nerve was isolated from surrounding connective tissue at the midtibial level and crushed for 7 s using smooth forceps. The muscle and skin layers were closed under aseptic conditions. Control cohorts (sham-operated mice) underwent the same surgical procedure, but the nerve was exposed and left intact. The time course of neuropathic pain development, manifested as mechanical hyperalgesia and tactile allodynia (painful responses to normally innocuous stimuli), was assessed by the application of von Frey filaments to the plantar surface of the hindpaw before surgery and at days 3, 7, 10, 14, and 18 after crush injury of the sciatic nerve. A series of mice, subjected to sciatic injury for 14 d, were treated with either the nonselective opioid antagonist naloxone (NLX) (1 mg/kg; Sigma) or the selective antagonists of μ-opioid [β-funaltrexamine (β-FNA); 15 mg/kg; Tocris Bioscience], δ-opioid (naltrindole; 10 mg/kg; Tocris Bioscience), and κ-opioid [norbinaltorphimine (nor-BNI); 10 mg/kg; Tocris Bioscience] receptors before measuring mechanical responsiveness to von Frey filaments. 

Spared nerve injury model of neuropathic pain. An additional set of experiments was performed using the “spared nerve injury” (SNI) model (Decosterd and Woolf, 2000) of neuropathic pain. Mononeuropathy was induced according to the method originally described in the rat (Decosterd and Woolf, 2000) and reproduced in the mouse (Bourquin et al., 2006). Mice were anesthetized with isoflurane (induction, 3%; surgery, 1.5%). The left common sciatic nerve was blunt dissected through the biceps and femoris muscle, freed of connective tissue, and the three peripheral branches (sural, common peroneal, and tibial nerves) of the sciatic nerve were exposed. The tibial and common peroneal nerves were ligated and transacted together. The sural nerve was carefully preserved by avoiding any nerve stretch or nerve contact with surgical tools. The muscle and skin layers were closed under aseptic conditions. Control cohorts (sham-operated mice) underwent the same surgical procedure, but the sciatic nerve was exposed and left intact. The time course of mechanical allodynia-like behavior was measured in the non-injured sural nerve skin territory from days 1 to 15 after surgery by the application of von Frey filaments.

Results

RNA isolation, cDNA synthesis, and real-time PCR

Total RNA from tissue was obtained by TRizol extraction (Invitrogen). One microgram of the isolated RNA was reverse transcribed into cDNA with an RT-PCR kit (Fermentas), according to the instructions of the manufacturer. Quantitative, real-time PCR was conducted on an MX-3000P Stratagene thermocycler using specific TaqMan expression assays (Applied Biosystems) and Universal PCR Master Mix (Takara). Results were normalized to the housekeeping genes glyceraldehyde-3-phosphate dehydrogenase or ribosomal 18S subunit expression and measured in parallel in each sample. Results are expressed as ΔΔCt. Duplicate transcript levels were determined in three independent experiments.

Western blot

Whole-cell lysates were prepared from the cerebral cortices, hippocampus, and spinal cords. Equal amounts of whole-cell protein were resolved on 12.5% SDS-PAGE and transferred to a polyvinylidene difluoride membrane. The following primary antibodies were used: anti-phospho-Smad-2 (1:1000; catalog #3101; Cell Signaling Technology); anti-phospho-Smad-1 (1:1000; a gift from Dr. C. H. Heldin, Ludwig Institute for Cancer Research, Uppsala University, Uppsala, Sweden), and anti-proopiomelanocortin (POMC) (1:1000; ab32893, Abcam). Immunoreactivity was detected with GE Healthcare ECL Advanced detection reagents and visualized by GE Healthcare Hyperfilm-ECL. Films were scanned, and optical densities were determined by using the Scion Image software (Scion Corporation).

Radioimmunoassay

After decapsulation, the spinal cords of the mice were extracted by hydro-extrusion and immediately frozen in liquid nitrogen. Peptides were extracted as described. Radioimmunoassay (RIA) of Leu-enkephalin and Dynorphin A was performed using commercial kits (Phoenix Pharmaceuicals), following the recommendations of the manufacturer. Each sample was assayed in triplicate.

Spinal cord explants stimulation

Mice were anesthetized and decapitated. The spinal cord was extracted by hydro-extrusion (De Sousa and Horrocks, 1979) and dissected through the middle line to obtain two hemi-spinal cords. Each hemi-spinal cord was placed onto a Millicell-CM culture plate (Millipore Corporation) and inserted into a six-well plate. Spinal cords were incubated for 1 h in 1.5 ml of DMEM (Sigma) with or without recombinant BMP-7 (20 ng/ml; R & D Systems) or activin (20 ng/ml; R & D Systems), in a humidified atmosphere of 5% CO₂, at 37°C. Each hemi-spinal cord served as the control for its complementary half. RNA isolation, retrotranscription to cDNA, and real-time PCR were performed as described previously.

Data analysis and statistics

Behavioral experiments were conducted blindly as to the genotype of the mice. All results are presented as mean ± SEM. Statistical differences between genotypes were analyzed with one-way, two-way, or repeated-measures ANOVA, followed by Bonferroni’s test. Individual time points were analyzed with unpaired t test. For all analyses, p < 0.05 was considered significant.

Results

Regional distribution of BAMBI in the nervous system

Representative in situ hybridization images at the mesencephalic and spinal cord levels are presented in Figure 2, A and B. Strong
Lack of BAMBI has no effect on general behavior

Mutant mice were fertile, appeared healthy, showed no apparent phenotypic defects, and displayed longevity that was indistinguishable from wild-type littersmates. Their sensorimotor abilities, locomotor activity, and coordination, measured in the open field and the rotarod tests, were similar to those of wild-type (data not shown). Lack of BAMBI reduces basal nocifensive responses in thermal, mechanical, and chemical/inflammatory tests of acute pain

We examined the behavioral responses of BAMBI−/− mice to noxious thermal (n = 12 per group), chemical/inflammatory (n = 8 per group), and mechanical (n = 12 per group) stimuli tests (Vierck, 2006). Two models of thermal nociception were used: the warm-water tail-immersion test that examines spinal-mediated responses, and the hotplate test that examines both spinal and supraspinal-mediated responses. In the tail-immersion test, BAMBI−/− mice experienced a significantly greater delay in tail withdrawal than the wild-type mice at 45, 47, and 49°C (Fig. 3A). Furthermore, there was a gene-dosage dependence of the response, because the reaction time of the BAMBI−/− mice occurred between wild-type and knock-out mice (Fig. 3B). The delays in licking behavior and jumping from a hotplate set at 50°C (Fig. 3C) and 52°C were also significantly longer in BAMBI−/− mice than in their wild-type littersmates.

The effect of BAMBI deletion on chemical pain behavior was assessed in the Formalin test. Injection of Formalin into the hindpaw induces a biphasic pain response; the first phase is thought to result from direct activation of TRPA1 cation channel in primary afferent sensory neurons, whereas the second phase has been proposed to reflect the combined effects ofafferent input and activity-dependent sensitization of CNS neurons within the dorsal horn (Coderre et al., 1990; Tjølsen et al., 1992; McNamara et al., 2007). Intradermal injection of Formalin into the hindpaw produced the typical biphasic paw-licking behavior in wild-type and BAMBI−/− mice; however BAMBI−/− mice showed a signific
Lack of BAMBI attenuates the expression of neuropathic pain responses

We assessed the potential involvement of BAMBI in chronic pain behaviors in two models of neuropathic pain that mimics peripheral nerve injury in humans: the crush injury of sciatic nerve (Bester et al., 2000) and the spared nerve injury (Decosterd and Woolf, 2000; Bourquin et al., 2006).

In the crush injury of sciatic nerve model, we evaluated the nocifensive response to the hindpaw tactile mechanical stimulation at graded intensities of stimuli for 18 d after crush (n = 10 per group). Wild-type mice displayed a progressive and significant decrease in the paw-withdrawal threshold, starting at day 7 and reaching the maximal allodynic response 14 d after crush. ANOVA indicated that BAMBI<sup>−/−</sup> mice were significantly less sensitive to mechanical stimuli compared with the wild-type group throughout the follow-up period (ANOVA; genotype: F(1,90) = 39.9, p < 0.001; time: F(5,90) = 48.9, p < 0.001; genotype × time: F(5,90) = 6.8, p < 0.001). In Figure 4 (top), we show representative graphs showing the reduced nocifensive responses of BAMBI<sup>−/−</sup> mice to hindpaw mechanical stimulation, at graded intensities of force on day 14 after injury compared with their wild-type littermates (genotype: F(1,69) = 80.7, p < 0.001; nerve injury: F(1,69) = 97.1, p < 0.001; genotype × nerve injury: F(1,69) = 25.0, p < 0.001).

In the spared nerve injury model of neuropathic pain (n = 7–9 animals per group) BAMBI<sup>−/−</sup> mice also exhibited a lower degree of mechanical allodynia than their wild-type littermates as indicated two-way ANOVA (genotype: F(1,104) = 90.8, p < 0.001; nerve injury: F(1,104) = 199.3, p < 0.001; genotype × nerve injury: F(1,104) = 13.1, p < 0.001) (Fig. 5).

The opioid antagonist naloxone reverses thermal and mechanical hypersensitivity in BAMBI<sup>−/−</sup> mice

The presence of BAMBI transcripts in the dorsal horn of the spinal cord and the periaqueductal gray matter, paradigmatic areas in which the endogenous opioiodergic system plays a key role in processing nociceptive information (Fields, 2000), directed us to examine whether the reduced sensitivity to pain in the mutant mice was linked to an opioid-mediated mechanism. For this purpose, a series of mice (n = 10 per group) were pretreated with the nonselective opioid receptor antagonist naloxone (1 mg/kg, i.p.) 30 min before performing the warm-water tail-immersion and von Frey tests. After naloxone treatment, BAMBI<sup>−/−</sup> mice showed a normalization in the withdrawal response to both thermal and mechanical stimuli, whereas no changes were evident in wild-type mice. Data are means ± SEM. *p < 0.05, **p < 0.01, and ***p < 0.001 versus BAMBI<sup>+/+</sup> mice (two-tailed Student’s t test).

The anti-allodynic phenotype of BAMBI<sup>−/−</sup> mice involves δ-opioid receptor activity

Fourteen days after sciatric nerve injury (Fig. 4, bottom), when maximal differences in neuropathic pain-related behaviors between genotypes were evident, naloxone administration (n = 10 per group) triggered mechanical hypersensitivity and allodynia in BAMBI<sup>−/−</sup> mice but did not influence significantly the nociceptive response in the wild-type group (genotype: F(1,69) = 16.6, p < 0.001; NLX treatment: F(1,69) = 11.5, p < 0.01; genotype × treatment: F(1,69) = 4.5, p < 0.05). To further define the opioid receptor subtype(s) responsible for the anti-allodynic phenotype, a series of BAMBI<sup>−/−</sup> mice, subjected to sciatric nerve injury for 14 d, were...
pretreated with selective antagonists of the different opioid receptor types, 60 min before analyzing the development of mechanical hypersensitivity. Subsequently, mice received a dose of naloxone to evaluate the degree of antagonism reached with each drug. As shown in Figure 6, the δ-selective antagonist naltrindole fully reversed the anti-allodynic phenotype ($F_{1,67} = 490, p < 0.001$), whereas the μ-opioid (β-FNA; $F_{1,67} = 48.5, p < 0.001$) and κ-opioid (nor-BNI; $F_{1,67} = 4.7, p < 0.05$) selective antagonists were barely effective.

Figures 4 and 5. Development of neuropathic pain in response to crush injury of sciatic nerve. Representative graphs showing the behavioral manifestations of neuropathic pain (mechanical allodynia) evaluated with von Frey monofilaments, on day 14 after SNI surgery in BAMBI$^{-/-}$ (filled circles) and wild-type (filled squares) mice. Values are the mean ± SEM percentage of hindpaw withdrawals elicited by mechanical stimuli of increasing strength, in wild-type (top graph) and BAMBI$^{-/-}$ mice (bottom graph) mice subjected to sham operation (open symbols) or SNI surgery (filled symbols). Two-way ANOVA indicates that BAMBI$^{-/-}$ mice were significantly less sensitive to mechanical stimuli compared with their wild-type littermates. Note also the difference in mechanical sensitivity between sham-operated BAMBI$^{-/-}$ and BAMBI$^{-/-}$ mice.

Basal expression of POMC mRNA and protein, as well as proenkephalin mRNA and the derived opioid peptide Leu-enkephalin, is increased in BAMBI$^{-/-}$ spinal cord

Given the regulatory function exerted by TGF-βs on gene transcription, we analyzed whether the spinal cord expression levels of genes encoding endogenous opioid peptides was affected by BAMBI deletion. Our results indicated that the basal level of POMC mRNA, the gene codifying for β-endorphin, was $2.46 \pm 0.30$-fold (95% confidence level, 1.63–3.29; $p = 0.022$, two-tailed $t$ test) higher in the BAMBI$^{-/-}$ mice ($n = 5$) than in the wild type ($n = 5$). We also observed a $1.64 \pm 0.11$-fold increase (95% confidence level, 1.32–1.97; $p = 0.007$, two-tailed $t$ test) in the basal level of proenkephalin (PENK) mRNA, the gene codifying for enkephalins, in BAMBI$^{-/-}$ mice compared with wild-type mice. Conversely, the basal level of prodynorphin (PDYN) mRNA, the gene codifying for dynorphins, was similar in both genotypes. To assess that increased transcription is translated to the protein level, we determined by Western blot or RIA the expression levels of the different opioid peptides. Spinal expression levels of POMC protein, determined by Western blot, and Leu-enkephalin, measured by RIA, were higher in BAMBI$^{-/-}$ than in wild-type mice (POMC: $+43.8 \pm 11.9\%$, $p = 0.034$; PENK: $+44.2 \pm 9.0\%$, $p = 0.011$). No differences between genotypes were observed in the spinal cord concentration of dynorphin, measured by RIA (data not shown).

As shown in Table 1, the spinal cord expression levels of genes encoding POMC, PENK, and PDYN were significantly and inversely related to BAMBI expression, whereas directly correlated with the mRNA levels of various TGF-β family members (TGF-β1, activin, and BMP-7) and with representative intracellular...
effectors from the TGF-β canonical (common SMAD4) and alternative (TAK1) signaling pathways (Wrana et al., 1994). These results led us to test whether exogenously administered recombinant TGF-βs influence transcriptional regulation of POMC and PENK under basal conditions. For this purpose, explanted spinal cords from wild-type mice were cultured in the presence of recombinant activin A (20 ng/ml) or BMP-7 (20 ng/ml) for 1 h, and POMC and PENK expression levels were assessed by quantitative RT-PCR. Because the TGF-β members acting physiologically on the spinal cord remain to be identified, we selected activin A and BMP-7 because they signal through the two different major Smad pathways in TGF-β superfamily signaling: activin/TGF-β-specific R-Smads (Smads 2 and 3) or BMP-specific R-Smads (Smads 1, 5, and 8). One hour incubation of spinal cord explants with activin A or BMP-7 significantly increased POMC mRNA levels [activin A: 2.7-fold vs untreated (95% confidence level, 1.57–3.38; two-tailed t test, p = 0.039); BMP-7: 1.74-fold vs untreated (95% confidence level, 1.63–1.84; two-tailed t test, p = 0.0001)]. Also, PENK mRNA levels increased significantly in explants incubated with BMP-7 versus untreated (1.64-fold; 95% confidence level: 1.32–1.97; two-tailed t test; p = 0.0079).

Discussion

To unravel novel roles for TGF-βs in the adult CNS, we have examined the expression pattern and the function of BAMBI, a pseudoreceptor that modulates negatively pan-TGF-β family signaling (Oichtchouk et al., 1999). We show that BAMBI transcripts are abundant in several CNS areas critically involved in pain processing (Fields, 2000; Todd and Koerber, 2006), such as the cingulate cortex, the mesencephalic periaqueductal gray, and the superficial laminae of the spinal cord dorsal horn. The presence of BAMBI protein and its preferential location in neurons was confirmed on the former structure by immunohistochemistry. At the peripheral level, in the dorsal root ganglion, BAMBI was also detected in virtually all sensory neurons but was absent in their satellite glia. The location of BAMBI expression is indicative of a role for TGF-β family members in nociceptive regulation, both central and peripherally. This suggestion is further supported by a recent report showing the antiallodynic effect exerted by recombinant TGF-β1 in a rat model of neuropathic pain (Echeverry et al., 2009). Thus, we aimed to assess possible changes in the responsiveness to painful stimuli, under conditions of enhanced TGF-β signaling. This objective directed us to develop a mutant mouse lacking BAMBI. As we expected, the absence of the inhibitory influence of BAMBI in knock-out mice resulted in increased levels of representative downstream effectors of the TGF-β signaling pathway, such as the phosphorylated forms of Smad1 and Smad2 transcription factors, in some regions of the CNS. Our first behavioral analyses was the absence in BAMBI−/− mice of phenotypic defects, affecting general motor and sensory functions, which could constitute confounding factors to interpret pain behaviors. Nevertheless, it should be noted that backcrossing to the sixth generation could be insufficient for complete abolition of potential “hitchhiking donor gene” confounds (Gerlai, 1996).

Figure 6. Effects of specific opioid antagonists on neuropathic pain behavior in BAMBI−/− mice. Representative graphs showing the behavioral manifestations of neuropathic pain (mechanical allodynia) evaluated with von Frey monofilaments, on day 14 after sciatic nerve crush injury (triangles), and the effect of pretreatment with selective opioid antagonists before performing the von Frey test (open circles). The antagonistic effect achieved by subsequent naloxone administration is also shown (filled circles). Values are the mean ± SEM percentage of hindpaw withdrawals elicited by mechanical stimuli of increasing strength. The selective antagonist of δ-opioid receptors, naltrindole (top graph), fully reversed the anti-allodynic phenotype of BAMBI−/− mice, whereas the μ-opioid (β-FNA) and κ-opioid (nor-BNI) selective antagonists were much less effective.
When acute pain paradigms were tested, we observed that the absence of BAMBI conditioned lower nociception-related responses, regardless of the modality of the noxious stimuli used (thermal, mechanical, and chemical/inflammatory). There was gene-dosage dependence in the nocifensive responses to thermal and mechanical stimuli, as indicated by the behavior of the heterozygotes. Thus, the nocifensive responses of BAMBI<sup>−/−</sup> mice to noxious stimuli are consistent with the hypothesized role for TGF-βs in controlling physiological pain perception under basal conditions. Interestingly, the opioid antagonist naloxone was able to reverse the pain-related behaviors of BAMBI<sup>−/−</sup> mice to that of wild-type controls, indicating that the hypalgesia exhibited by BAMBI<sup>−/−</sup> mice under basal conditions can be attributed to an enhanced endogenous opioid tone.

In the chronic and pathological neuropathic pain model, our results indicate that, in BAMBI<sup>−/−</sup> mice, the release of TGF-β signaling from the inhibitory influence of BAMBI also attenuated pain hypersensitivity and allodynia induced by sciatic nerve injury over a follow-up period of 18 d. This anti-allodynic phenotype of BAMBI<sup>−/−</sup> mice was also evidenced in another model of peripheral mononeuropathy, such as the spared nerve injury. These results are in agreement with the aforementioned data of Echeverry et al. (2009) showing that chronic (14 d) intrathecal infusion of recombinant TGF-β1 prevented the development of allodynia and reversed previously established neuropathic pain in rats. The mechanism proposed by the authors involves an inhibitory effect on the spinal cord inflammatory response to nerve injury exerted by TGF-β1. In our study, naloxone completely restored the attenuated allodynic response of BAMBI<sup>−/−</sup> mice to that of wild type, suggesting that TGF-β signaling is critical in controlling the expression of abnormal nociceptive responses subsequent to nerve injury, by a mechanism fully dependent on ongoing opioid receptor activation. The results obtained with selective opioid antagonists indicate that the anti-allodynic phenotype is mainly dependent on δ-receptor activity.

The expression of BAMBI in dorsal horn overlaps with several type 1 receptors for TGF-β family members, such as ALK-3, ALK-4, and ALK-6, suggesting a functional implication of TGF-β signaling in this region. It is well known that the dorsal horn of the spinal cord is the zone in which nociceptive primary afferent axons terminate, establishing the first synapse in the ascending pathways that convey the sensory information from peripheral receptors to the brain. It contains local neuronal circuits that are involved in modulating incoming nociceptive information before its transmission up to the brain (Todd and Koerber, 2006). The superficial laminae of the dorsal horn contains intrinsic opioid inhibitory interneurons that express enkephalin, dynorphin, and β-endorphin and also receives afferent β-endorphinergic input from the arcuate and tractus solitarius nuclei (Tsou et al., 1986; Fields, 2000). Under physiological conditions, these opioid peptides exert an inhibitory control on the incoming nociceptive through receptors located both presynaptically on nociceptor terminals and postsynaptically on intrinsic dorsal horn neurons (Yaksh et al., 1982; Kondo et al., 2005; Pasternak, 2005). Under pathological conditions, the endogenous opioidergic system has been reported to counteract the development of neuronal hyper-excitability after intense noxious drive (Guan et al., 2006) and neuropathic pain-related states after nerve injury (Hao et al., 1998; Mansikka et al., 2004).

The precursors of endogenous opioid peptides are encoded by three different genes: the PENK gene that is translated into the protein precursor of enkephalins, the PDYN gene that encodes dynorphins, and the POMC gene that gives rise to β-endorphin. Our results indicate that basal levels of POMC mRNA and POMC protein, as well as PENK gene and the opioid peptide enkephalin, are higher in the spinal cords of BAMBI<sup>−/−</sup> mice than in those of wild type. Furthermore, the spinal cord expression levels of genes encoding precursor proteins of the opioid peptides directly correlated with members of the TGF-β family and with their intra-cellular effectors Smad4 and TAK1, whereas the relationship with BAMBI was inversely proportional. These results suggest that an increased transcriptional activity of endogenous opioids is likely mediating the reduced nociceptive responses in the absence of BAMBI. This interpretation is also supported by the transcriptional regulatory effect exerted by recombinant BMP-7 and activin A on POMC and PENK genes, in spinal cord explants. These results are consistent with previous reports showing in vitro that members of the TGF-β family regulate the transcription of genes coding for the endogenous opioid peptides in a variety of human and rodent cell preparations (Kamphuis et al., 1997; Kavelaars and Heijnen, 2000; Nudi et al., 2005), and in vivo (Bouret et al., 2001).

Many studies support an antinociceptive action of enkephalins and β-endorphin by negatively modulating transmission of nociceptive information at the spinal level (Yaksh et al., 1982). Furthermore, POMC and PENK gene delivery to spinal cord dorsal horn induces analgesia, which is associated with increased production of β-endorphin or enkephalin by specific subsets of spinal neurons (Hao et al., 2003; Wu et al., 2004). In addition, it has been demonstrated that mice deficient in PENK showed exaggerated behavioral responses to painful stimuli (König et al., 1996). Also, transgenic mice, with a selective deficiency of β-endorphin, lack the opioid-mediated (naloxone reversible) analgesia induced by stress (Rubinstein et al., 1996). Therefore, our data suggest that hypalgesic-related responses in mice lacking BAMBI is attributable to an increased tonic activation of opioid receptors by their endogenous ligands. Particularly, the anti-allodynic phenotype is mainly dependent on δ-receptor signaling, which would be consistent with the relatively high affinity of enkephalins for this receptor (Dhawan et al., 1996).

Although our study was focused on spinal mechanisms, we cannot exclude the contribution of supraspinal and peripheral mechanisms in the altered pain processing displayed by BAMBI<sup>−/−</sup> mice. In fact, BAMBI localization in dorsal root ganglion neurons and the reduced pain behavior of BAMBI<sup>−/−</sup> mice in the Formalin test could point to a reduction in the excitability and postinjury

### Table 1. Pearson’s correlation coefficient values (r) obtained from the linear regression analyses correlating spinal cord mRNA expression levels of TGF-β1, BMP-2, BMP-7, activin, SMAD4, TAK1, and BAMBI with the expression of genes encoding POMC, PENK, and PDYN

<table>
<thead>
<tr>
<th>Gene</th>
<th>TGF-β1</th>
<th>BMP-2</th>
<th>BMP-7</th>
<th>Activin</th>
<th>SMAD4</th>
<th>TAK1</th>
<th>BAMBI</th>
</tr>
</thead>
<tbody>
<tr>
<td>POMC</td>
<td>0.65***</td>
<td>0.54*</td>
<td>0.47*</td>
<td>0.48*</td>
<td>0.57*</td>
<td>0.75***</td>
<td>0.54*</td>
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<tr>
<td>PENK</td>
<td>0.76***</td>
<td>0.65**</td>
<td>0.69**</td>
<td>0.65**</td>
<td>0.80***</td>
<td>0.71***</td>
<td>0.58*</td>
</tr>
<tr>
<td>PDYN</td>
<td>0.71***</td>
<td>0.72***</td>
<td>0.75**</td>
<td>0.80***</td>
<td>0.68**</td>
<td>0.82***</td>
<td>0.61**</td>
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</table>

WT, Wild type; KO, knock-out. *p < 0.05, **p < 0.01, and ***p < 0.001.
sensitization of primary sensory neurons (Tjølsen et al., 1992; McNamara et al., 2007) that play important roles in the initiation and maintenance of persistent pain.

Our present results and those of Echeverry et al. (2009) suggest that members of the TGF-β family modulate negatively pain perception under physiological and pathological conditions. However, the opposite response has been described for activin A that is released during peripheral inflammation, acting on sensory nerve terminals to increase the expression of calcitonin-gene related peptide in dorsal root ganglion neurons and leading to enhanced pain responses (Xu et al., 2005). In addition, subcutaneous injection of activin A induced immediate thermal hyperalgesia in rats, through a mechanism involving acute sensitization of TRPV1, which suggests a role for activin A in injury-induced hyperalgesia (Zhu et al., 2007). The extremely variable cellular responses elicited by many TGF-β superfamily members, depending on cell type and stimulation context, could account for these antagonistic effects, and the contribution of each of these mechanisms to nociception deserves further investigation.

In conclusion, the absence of BAMBI unravels a critical role for TGF-β signaling in nociceptive perception under physiological and neuropathic pathological conditions, through a mechanism involving transcriptional regulation of the opioid peptides. Moreover, our results suggest that specific members of the TGF-β family, their signaling effectors, and modulator molecules could be used as therapeutic targets for the design of novel analgesic drugs that may act by increasing the activity of the endogenous opioid system.

References


