Tissue-nonspecific alkaline phosphatase promotes axonal growth of hippocampal neurons

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ABSTRACT Axonal growth is essential for establishing neuronal circuits during brain development and for regenerative processes in the adult brain. Unfortunately, the extracellular signals controlling axonal growth are poorly understood. Here we report that a reduction in extracellular ATP levels by tissue-nonspecific alkaline phosphatase (TNAP) is essential for the development of neuritic processes by cultured hippocampal neurons. Selective blockade of TNAP activity with levamisole or specific TNAP knockdown with short hairpin RNA interference inhibited the growth and branching of principal axons, whereas addition of alkaline phosphatase (ALP) promoted axonal growth. Neither activation nor inhibition of adenosine receptors affected the axonal growth, excluding the contribution of extracellular adenosine as a potential hydrolysis product of extracellular ATP to the TNAP-mediated effects. TNAP was colocalized at axonal growth cones with ionotropic ATP receptors (P2X7 receptor), whose activation inhibited axonal growth. Additional analyses suggested a close functional interrelation of TNAP and P2X7 receptors whereby TNAP prevents P2X7 receptor activation by hydrolyzing ATP in the immediate environment of the receptor. Furthermore inhibition of P2X7 receptor reduced TNAP expression, whereas addition of ALP enhanced P2X7 receptor expression. Our results demonstrate that TNAP, regulating both ligand availability and protein expression of P2X7 receptor, is essential for axonal development.

INTRODUCTION
Axonal outgrowth is central to the processes of brain development when neurons establish precise connectivity patterns as well as to the processes of regeneration and brain repair in the adult CNS.
of protein kinases such as calcium/calmodulin-dependent kinase II (CaMKII), p38 mitogen-activated protein kinase, or glycogen synthase kinase 3 (GSK-3) (Armstrong et al., 2002; Gomez-Villafuertes et al., 2009), and stimulates vesicular neurotransmitter release (Gomez-Villafuertes et al., 2001; Diaz-Hernandez et al., 2002b; Gualix et al., 2003). The availability and half-life of extracellular ATP is governed by ectonucleotidases that sequentially hydrolyze the nucleotide to adenosine (Robson et al., 2006; Zimmermann, 2006). Adenosine functions as a major negative neuromodulator in the CNS. It activates presynaptic metabotropic receptors to promote inhibition of neurotransmitter release and to reduce neuronal excitability (Fredholm et al., 2005).

Ectonucleotidases belong to several protein families that differ regarding their functional and molecular properties (Zimmermann, 2006). The alkaline phosphatase (ALP) family releases P; from a variety of organic compounds and is capable of degrading nucleoside 5’-tri-, -di-, and -monophosphates, eventually producing adenosine from extracellular ATP. The mammalian isoforms share an alkaline pH optimum and are anchored to the membrane via a glycosylphosphatidylinositol anchor (Millan, 2006). Tissue-nonspecific ALP (TNAP) is expressed in a multitude of tissues including liver, bone, kidney, and brain. In the adult mammalian CNS, TNAP represents the only isoform of ALPs and is associated with the blood vessel endothelium and with neuropil, including synaptic contacts (Langer et al., 2008).

By controlling extracellular levels of ATP and adenosine ectonucleotidases could play a pivotal role in the control of neurite development. In the present work we analyzed the role of TNAP and its interaction with a specific ATP receptor (P2X5 receptor) in axonal growth using cultured hippocampal neurons as a well established model system (Dotti et al., 1988).

RESULTS

TNAP regulates extracellular ATP levels in cultured hippocampal neurons

Hippocampal neurons initially plated in culture revealed no discernible neurites. Within the next 72 h, neurons started to grow extensive microtubule-associated protein 2 (MAP2)+-positive neurites (Figure 1, A1 and A2). One of these became the axon and could be identified with anti-Tau-1 antibodies (Figure 1, A3). Interestingly, the culture supernatant contained significant levels of ATP (2.28 ± 0.38 μM) that considerably declined with increasing postplating time. The fastest loss occurred during the first 24 h (Figure 1B), reaching a value of 0.33 ± 0.14 μM of ATP after 72 h. As shown by quantitative immunoblotting, the decrease in extracellular ATP levels was paralleled by an increase in neuronal expression of the ectonucleotidase TNAP (Figure 1C). In accordance with that, TNAP activity increased by a factor of 7.58 ± 0.8 during the first 72 h (Supplemental Figure S1).

We therefore investigated whether the extracellular ATP levels could be causally linked to TNAP activity. Addition of ALP (4 U/ml) to the culture media induced a rapid and persistent loss of extracellular ATP (Figure 1D). Moreover, specific inhibition of endogenous TNAP activity with levamisole (500 μM) (Kozlenkov et al., 2004; Langer et al., 2007) significantly elevated extracellular ATP concentrations at each time point (Figure 1D), suggesting that endogenous TNAP can control extracellular ATP levels.

TNAP promotes axonal but not dendritic elongation

Double immunolabeling of neurons for TNAP and βIII-tubulin or F-actin staining (Alexa Fluor 594 phalloidin) revealed a strong somatic and axonal and a weaker dendritic localization of TNAP (Figure 2A). TNAP immunostaining was most intense at the most distal axon regions. Furthermore TNAP immunoreactivity typically revealed a dotted pattern both in the principal axonal growth cone and in most of the actin-rich branching points of the primary axon (Figure 2, A and B). TNAP catalytic activity was identified in the identical locations by enzyme histochemical staining using the 5-bromo-4-chloro-3-indolyl phosphate/nitroblue tetrazolium (BCIP/NBT) reaction. It was present in the entire neuron and was strongest in the soma and at the axonal growth cone (Figure 2C).

We therefore investigated whether TNAP activity is related to axonal or dendritic growth, in both hippocampal and cortical neurons. Neurons were stained with anti-Tau-1 and anti-MAP2 antibodies to evaluate the specific effects of TNAP on axons and dendrites, respectively (Figure 3A and Supplemental Figure S2A). When grown

![Image](https://via.placeholder.com/150)

**FIGURE 1:** Increased TNAP expression in differentiating cultured hippocampal neurons reduces extracellular ATP levels. (A) Representative images of hippocampal neurons, stained with anti–Tau-1 (green) and anti-MAP2 (red) antibodies to illustrate their morphology during the first 72 h of culture. (A1) Emerging neurites (at 24 h). (A2) One of the neurites takes the lead (at 48 h). (A3) Axon as identified by anti–Tau-1 immunofluorescence and dendrites further grow and branch (at 72 h). Scale bar, 50 μm. (B) Extracellular ATP levels in the medium of hippocampal neurons cultured for 1, 2, and 3 d in vitro (DIV). Values represent mean ± SEM of six experiments in triplicate. (C) Western blots of TNAP in cultured hippocampal neurons at 1, 2, and 3 DIV. α-Tubulin served as a loading control. Representative immunoblots probed for TNAP are shown at the bottom of the graph. Values represent mean ± SEM (n = 3). (D) Effects of ALP (4 U/ml) and levamisole (500 μM) on extracellular ATP levels in the medium of hippocampal neurons cultured for 1, 2, and 3 DIV. ATP levels represent mean ± SEM of five experiments in triplicate. *p < 0.05, **p < 0.01, and ****p < 0.0001 vs. control (unpaired t test).
in the presence of levamisole (500 μM), neurons developed significantly shorter axons (Figure 3, A and B; Supplemental Figure S2, A and B). The inhibitory effect of levamisole on total axonal length was concentration dependent with a half-maximal inhibitory concentration value of 523.1 ± 162.8 μM (Figure 3C). In contrast, levamisole (500 μM) had no effect on dendrite length (Figure 3B). Addition of ALP (500 μM) had no effect on dendrite length (Figure 3B). In addition, at a concentration of 5 mM levamisole entirely abolished neuritogenesis and triggered neuronal death. In accordance with the inhibitory effect of levamisole, addition of ALP to the culture media induced a considerable increase in total axonal length (Figure 3, A and B; Supplemental Figure S2, A and B). This concerned a significant elongation of both the principal axon and secondary axon branches. But in hippocampal neurons, the number of secondary ramifications was unaltered (Figure 3, A and B). The ALP-induced axonal elongation was concentration dependent with a half-maximal effective concentration value of 1.62 ± 0.34 U/ml, and maximal axonal growth was obtained at 4 U/ml (Figure 3D). As for levamisole, exogenous ALP (4 U/ml) had no significant effect on dendrite length (Figure 3B).

We further analyzed the effect of TNAP over expression and knockdown on axonal growth by cell transfection using plasmids encoding green fluorescent protein (GFP), TNAP, and TNAP-targeting short-hairpin RNA (shRNA). The efficiency of the commercial shRNA TNAP for TNAP knockdown was verified using a human cell line (human embryonic kidney [HEK] 293T) transfected to overexpress the mouse TNAP enzyme (Supplemental Figure S3). The specificity of commercial shRNA TNAP used was confirmed by checking that it was able to prevent the protein expression of murine wild-type TNAP full-length cDNA, but not the protein resistant to the shRNA sequence generated by a site-directed mutated TNAP cDNA (TNAPmut) (Supplemental Figure S3). Subsequently hippocampal and cortical neurons were transfected with the TNAP-, TNAPmut-, and shRNA TNAP–containing vector at 1 d in vitro (DIV) and fixed at 3 DIV to examine the length and ramifications of their axons. As compared with transfection with GFP alone, TNAP knockdown induced shortening of the principal axon and of the axonal branches (Figure 4, A–C) while the expression of TNAP and TNAPmut induced axonal elongation and branches either in hippocampal neurons (Figure 4B) or cortical neurons (Supplemental Figure S2, C and D). In addition, the concomitant transfection of neurons with TNAP and shRNA TNAP abrogated the negative effect of TNAP knockdown on axonal growth. However, when neurons were cotransfected with TNAPmut and shRNA TNAP, the axons grew in a similar range to that observed on neurons transfected with TNAP or TNAPmut (Figure 4, A and B). These data suggest that endogenous neuronal TNAP activity is essential for proper axon extension.

**Adenosine does not affect the first steps of axonal growth**

As an ectonucleotidase, TNAP hydrolyzes extracellular ATP or ADP, thus limiting their effects on P2 receptors, and simultaneously produces extracellular adenosine that acts via adenosine-specific receptors (Zimmermann, 2006). We thus investigated the possibility that the effects of TNAP on axonal growth are mediated by extracellular adenosine. Treatment of neurons with 10 nM adenosine to activate A₁ receptors or with the selective A₁ receptor agonist, N°-cyclopentyladenosine (CPA), induced no significant changes in axonal growth (Figure 5). Similar results were obtained when neurons were treated with 10 μM adenosine to activate A₃ receptors (Figure 5).

To fully discard a participation of adenosine in TNAP-mediated axonal growth, we added to the culture medium adenosine deaminase (ADA, 0.2 U/ml), an enzyme that converts adenosine to its inactive metabolite, inosine (Cristalli et al., 2001). ADA itself had no significant effect on the length of principal and secondary axons (Figure 5, D and G). Moreover, when coapplied with ALP, ADA could not reverse the axonal growth induced by ALP (Figure 5, F and G).

**Coordinated action of TNAP and P2X₇R on axonal growth**

We therefore investigated the possibility that TNAP exerts its effects by eliminating extracellular ATP. This notion was supported...
TNAP induces axonal elongation

by our recent work describing ATP acting via P2X7 receptors as a negative modulator of axonal growth and branching (Díaz-Hernández et al., 2008). Double immunostaining with antibodies against TNAP and P2X7 revealed that the two proteins were colocalized in microdomains of the growth cones (Figure 6, A–C). Furthermore, axonal growth induced by inhibition of P2X7 receptors with 1 μM brilliant blue G (BBG) (Michel et al., 2007) (Figure 6, F, J, and K) or 1 μM A438079 (Honore et al., 2006; McGarvaughy et al., 2007) (Figure 6, J and K) was reverted when TNAP activity was inhibited by levamisole (Figure 6, G, J, and K), suggesting that increased extracellular ATP levels reduced the effect of the inhibitors. Similarly, when neurons were cultured in the presence of BBG or A438079, levamisole no longer inhibited axonal growth (Figure 6, D, G, J, and K).

Coapplication of ALP (4 U/ml) and BBG (1 μM) was no more effective on axonal growth than application of either ALP or BBG alone (Figure 6, D, F, and H–K). Because our previous work had shown that overexpression of P2X7 receptors reduces axonal length and branching (Díaz-Hernández et al., 2008) (Figure 7, A and B), we investigated whether this effect could be reverted by TNAP. Neurons

FIGURE 3: Effect of TNAP on axonal growth. (A) Representative images of hippocampal neurons (immunostained for Tau-1 [green] and MAP2 [red] to differentiate between axons and dendrites) cultured for 3 DIV in the absence (1) or presence of ALP (4 U/ml, 2) and of levamisole (500 μM, 3). 4′,6-Diamino-2-phenylindole–stained nuclei are in blue. Scale bar, 50 μm. (B) Quantification of axonal length and the number of ramifications from experiments shown in (A). The total axonal length (corresponding to the principal axon plus ramifications), length of the principal axon, length of the axonal ramifications, number of axonal ramifications, and dendritic length were analyzed. The 100% values in control neurons for total axonal length, principal axon length, and length of ramifications correspond to 214.5 ± 13.4 μm, 158.6 ± 10.9 μm, and 43.1 ± 7.1 μm, respectively. The number of axonal ramifications in control neurons was 1.8 ± 0.3. The total length of the dendritic processes includes its ramifications. The 100% value of the total dendritic length in control neurons corresponds to 31.1 ± 2.6 μm. Values represent the mean ± SEM (n = 4) with at least 20 neurons analyzed in each experiment. *p < 0.05, **p < 0.01, and ***p < 0.001 vs. control (one-way analysis of variance [ANOVA]). (C and D) Concentration-response curves for the effect of levamisole (C) and ALP (D) on axonal growth of hippocampal neurons maintained for 3 DIV. Values represent mean ± SEM (n = 3) with at least 30 neurons analyzed in each experiment.
overexpressing P2X$_7$-GFP and treated with exogenous ALP developed axons of similar length as neurons transfected with a plasmid expressing GFP alone (Figure 7G). Similar results were obtained when neurons transfected with the P2X$_7$-ires-GFP plasmid, which independently encodes the P2X$_7$ receptor and GFP, were treated with ALP (Figure 7, B, C, and G).

FIGURE 4: Expression of TNAP control axonal growth. (A) Hippocampal neurons transfected at 1 DIV with pEGFP (1), TNAP IRES (2), shRNA TNAP (3), or TNAPmut (4) and fixed at 3 DIV. Scale bar, 50 μm. (B) Effect of overexpression of wild-type or mutant TNAP and silencing of TNAP on axon length and the number of ramifications in hippocampal neurons. The total axonal length (corresponding to the principal axon plus ramifications), length of the principal axon, and length and number of the axonal ramifications were analyzed. The 100% values in control neurons for total axonal length of EGFP-transfected neurons, principal axon length of EGFP-transfected neurons, and length of ramifications correspond to 310.8 ± 28.0 μm, 242.4 ± 28.7 μm, and 126.9 ± 24.2 μm, respectively. The number of axonal ramifications in EGFP neurons was 2.6 ± 0.9. Values represent mean ± SEM (n = 5) with at least 20 neurons analyzed in each experiment. *p < 0.05, **p < 0.01, ***p < 0.001 (one-way ANOVA). (C) Representative images of electroporated neurons with shRNA TNAP and its neighboring nontransfected neurons (immunostained with βIII-tubulin [red]). Scale bar, 50 μm.

In an additional series of experiments, we tested the possibility that axonal growth and branching induced by knockdown of P2X$_7$ receptors (Diaz-Hernandez et al., 2008) could be counterbalanced by silencing TNAP expression. Indeed, parallel neuronal knockdown of P2X$_7$ receptors and TNAP permitted normal neuronal axon development and reversed the axonal growth impairment induced by shRNA TNAP (Figure 7, D–F and H). A higher neuronal axon development was observed when the hippocampal neurons were cotransfected with P2X$_7$ and TNAP (Figure 7G).

These data suggested that TNAP-induced stimulation of axonal growth is mediated by enzymatic degradation of ATP, the growth-inhibiting P2X$_7$ receptor agonist. We therefore searched for possible interrelations in the expression of the two proteins. We quantified the levels of TNAP mRNA in the presence of the P2X$_7$ receptor antagonist BBG and the levels of P2X$_7$R mRNA in the presence of exogenously added ALP (3 d each). Treatment of hippocampal neurons with BBG induced a significant decrease in TNAP mRNA levels (35.4 ± 9.3%), and treatment with exogenous ALP increased the levels of P2X$_7$ mRNA (93.3 ± 40.8%) (Figure 7, I and J).

DISCUSSION

In this study we provide evidence that the ectonucleotidase TNAP in concert with the ionotropic P2X$_7$ ATP receptor governs the axonal growth in neurons. ATP is a short-lived and short-ranged extracellular signaling molecule whose lifetime is controlled by cell surface–located ectonucleotidases. TNAP is colocalized at axonal growth cones with the P2X$_7$ receptor. By eliminating extracellular ATP, TNAP prevents the activation of P2X$_7$ receptors, which are inhibitory to axonal growth. This novel purinergic signaling mechanism involving TNAP and the P2X$_7$ receptor could be critically involved in the control of axonal growth during development but also during axonal regeneration in the adult brain.

P2X$_7$ receptors are ATP-gated, non-selective ion channels permeable to Na$^+$, K$^+$, and Ca$^{2+}$. This confers to them the ability to act as direct conduits for Ca$^{2+}$ influx in the absence of membrane depolarization (Khakh and North, 2006). Intracellular Ca$^{2+}$ dynamics play a central role in the control of neuronal motility including axonal growth and guidance (Gomez and Zheng, 2006). In a previous study we demonstrated that axonal growth of cultured hippocampal neurons is inhibited by extracellular ATP via the P2X$_7$ receptor. This was correlated with the CaMKII-dependent regulation of actin-associated proteins (Diaz-Hernandez et al., 2008). Similar to our findings in the
neuronal culture supernatant, extracellular ATP is present in the brain extracellular space. It may be released via exocytosis or through membrane channels, or it may be derived from damaged cells (Abbracchio et al., 2009). Although ATP levels measured in our experimental conditions are in micromolar range, it is necessary to take into account that this value is the result of cellular release that dilutes a long time in the culture media, and therefore higher concentration of the nucleotide at the cellular environment can be deduced. This conclusion is supported by several experimental evidences; first, both addition of catalytically active

Ca\textsuperscript{2+} concentrations could mediate the direct interaction of Ca\textsuperscript{2+}-sensitive proteins with regulatory elements within the DNA sequence.

TNAP is expressed in a variety of mammalian tissues, but little is known concerning its functional role. The best investigated function concerns its involvement in bone mineralization and remodeling (Kaunitz and Yamaguchi, 2008). Mice deficient in the TNAP gene reveal defective bone mineralization and mimic a severe form of human hypophosphatasia. But TNAP knockout mice also suffer from epileptiform seizures and die from apnea before weaning. They

ALP and TNAP overexpression are able by themselves to promote axonal growth as both remove the basal extracellular ATP. Second, electroporation data showed that effects induced by TNAP are restricted to neurons that express this protein, without affecting neighboring cells. Finally, the ATP levels reached at the growth cone are high enough to activate the P2X\(_7\) receptor, which is the P2 receptor with lower affinity for ATP. We further show that the ATP hydrolysis product adenosine is not involved in the control of neural axonal outgrowth.

The data presented reveal that TNAP and the P2X\(_7\) receptor exhibit a dotted distribution on the axonal growth cones and that both proteins are also very closely functionally interrelated. Knockdown of the P2X\(_7\) receptor increases axon length and has the opposite effect as TNAP knockdown—whereby knockdown of both proteins counterbalances the effect. Similarly, overexpression of the P2X\(_7\) receptor reduces axonal length, and it was reverted by overexpression of TNAP or addition of ALP. However, while addition of exogenous ALP only compensates to return to control levels, the overexpression of TNAP was able to significantly increase axonal length. This discrepancy can be explained by considering that overexpression of TNAP specifically hydrolyzes ATP at the environment of the P2X\(_7\) receptor on the axonal growth cone. On the other hand, the fact that ALP treatment of hippocampal neurons increased P2X\(_7\) mRNA levels and P2X\(_7\) antagonism decreased TNAP mRNA levels indicates a coordinated regulation of these two proteins at the transcriptional level. At present little is known regarding the transcriptional regulation of the P2X\(_7\) receptor and TNAP. TNAP expression is enhanced by retinoic acid (Zhou et al., 1994). In previous studies we demonstrated that inhibiting the P2X\(_7\) receptor induces phosphorylation of GSK-3 and thus inhibits its catalytic activity (Diaz-Hernandez et al., 2008; Gomez-Villafuertes et al., 2009). GSK-3 activity in turn can regulate transcription factors (Grimes and Jope, 2001; Ortega et al., 2010), potentially modulating TNAP mRNA levels. Moreover, enhanced P2X\(_7\) receptor activation and increased intracellular

FIGURE 5: Lack of effects of adenosine on ALP-induced axonal growth. (A–F) Hippocampal neurons were cultured for 3 DIV in the absence (A) or presence of adenosine (Ado, 10 μM) (C) or the A\(_3\) adenosine receptor agonist CPA (50 nM) (E) and in the presence of exogenous ALP (4 U/ml) (B). Neurons were also treated with ADA (0.2 U/ml) in the absence (D) or presence (F) of exogenous ALP (4 U/ml). They were fixed and immunostained for βIII-tubulin (green) for analysis of axonal length. Scale bar, 50 μm. (G) Analysis of axonal length corresponding to the experiments shown in A–F. The 100% values in control neurons for total axonal length, principal axonal length, and length of ramifications correspond to 198.9 ± 9.8 μm, 148.2 ± 7.7 μm, and 31.3 ± 4.4 μm, respectively. Values represent mean ± SEM (n = 3) with at least 25 neurons analyzed in each experiment. *p < 0.05 and **p < 0.001 vs. control; NS, nonsignificant (one-way ANOVA).
display neural defects revealing aberrant development of the lumbar nerve roots and disturbances in intestinal physiology (Narisawa et al., 1997). Our demonstration of an association of TNAP with developing axons of cultured hippocampal neurons matches previous findings of an axonal localization of TNAP during neural development in situ, where the enzyme may similarly be involved in the control of axon elongation. TNAP activity was detected in association with nerve fibers emerging from the myelencephalon and spinal cord of the developing mouse brain (Narisawa et al., 1994), and an electron microscopic analysis of the developing marmoset brain revealed an association of ALP activity with the axonal surface (Fonta et al., 2005).

Furthermore TNAP is associated with the brain neurogenic regions during early mouse embryonic development. From E8.5 onward, it is highly expressed at the plasma membrane of the neuroepithelial stem cells of the neural tube (Narisawa et al., 1994). During later development of the CNS, TNAP activity becomes down-regulated, but it remains highly expressed in the ventricular neurogenic regions up to adult stage (Langer et al., 2007). The functional role of TNAP during neural development has not been investigated, but it is possible that—by controlling the extracellular availability of purinergic signaling molecules—it plays a role in immature neuron mobility and migration. In a variety of nonmammalian species, ALPs have been implicated in cell guidance and migration (Thibaudeau et al., 1993).

The purinergic mechanisms described here may in addition have implications for regenerative processes in the adult brain and for neuronal plasticity. In the injured CNS severed axons can start axonal outgrowth at the site of lesion, but the growth cones soon adopt a dystrophic morphology typical of growth cone inhibition (Schnorrer and Dickson, 2004; Liu et al., 2006). Under some pathological conditions like traumatic injury, hypoxia, or ischemia, ATP is released from damaged cells, resulting in elevated extracellular concentrations (Phillis et al., 1993; Juranyi et al., 1999) that would be inhibitory to axonal growth. Elevated ATP levels were shown to cause even neuronal cell death in a variety of models that reproduce these pathological conditions in vitro (Amandio et al., 2002) or in vivo (Ryu et al., 2002). This can involve activation of P2X7 receptors (Delarasse et al., 2009).

Moreover, similar mechanisms may apply to axonal sprouting and to presynaptic structural and functional plasticity. Axonal sprouting occurs as a restorative process emerging from neurons other than the lesioned ones and can result in functional synaptic reinnervation (Frotscher et al., 1997). It is interesting to note that TNAP has also been identified in association with the neuropil of the adult rodent and primate brain, where it was located around synaptic contacts (Fonta et al., 2005; Langer et al., 2008). Synaptic TNAP expression coincided with the time course of high-rate synaptogenesis, implicating a function in synaptic maturation and later in mature synaptic activity (Fonta et al., 2005). While synaptic TNAP may be involved in the control of purinergic-mediated neurotransmission or cotransmission (Díaz-Hernandez et al., 2000, 2002a; North and Verkhratsky, 2006), one may speculate that TNAP and ATP could also play an antagonistic role in the structural remodeling of synapses induced by neural activity (Bourne and Harris, 2008).

Taken together our results suggest that purinergic signaling mechanisms involving TNAP and the P2X7 receptor critically control axonal extension and axonal branching in hippocampal neurons, whereby TNAP controls the local availability of growth-inhibiting extracellular ATP. These mechanisms may be of

FIGURE 6: Involvement of TNAP and P2X7 in axonal growth. (A–C) Double immunofluorescence analysis of hippocampal neurons fixed at 3 DIV and stained with antibodies against TNAP (green) and P2X7 (red). (B and C) Images of the axonal growth cones at higher magnification. The axonal growth cone depicted in B belongs to the neuron shown in A (box). Scale bars, A, 50 μm; B and C, 20 μm. (D–I) Effect of P2X7 receptor antagonism and inhibition of TNAP on cultured hippocampal neurons; representative images of cultured hippocampal neurons; cultured hippocampal neurons; representative images of cultured hippocampal neurons cultured for 3 DIV in the absence (D) or presence (E) of levamisole (500 μM), BBG (1 μM) (F), levamisole (500 μM) and BBG (1 μM) (G), ALP (4 U/ml) (H), and BBG (1 μM) and ALP (4 U/ml) (I) and immunostained for βIII-tubulin (green). (J and K) Analysis of axonal length corresponding to the experiments shown in D–I (Leva, levamisole). The 100% values in control neurons for total axonal length and principal axon length correspond to 198.9 ± 9.8 μm and 148.2 ± 7.7 μm, respectively. Values represent mean ± SEM (n = 3) with at least 25 neurons analyzed in each experiment. *p < 0.05, **p < 0.01, and ***p < 0.001 vs. control; NS, nonsignificant (one-way ANOVA).
relevance for the developing nervous system but also for neuronal regeneration and possibly even synaptic plasticity in the adult brain.

**MATERIALS AND METHODS**

**Reagents**

The following reagents were used in this study: levasimole (A4341; AppliChem, Darmstadt, Germany); BCIP/NBT ALP substrate (B5655), bovine intestinal ALP (P6772), adenosine (A9251), and BBG (B5133) (all from Sigma-Aldrich, St. Louis, MO); A-438079 hydrochloride (2972; Tocris Bioscience, Bristol, UK); CPA (691763; Boehringer Ingelheim, Ingelheim am Rhein, Germany); and ADA (10901221; Roche Diagnostics, Berlin, Germany).

**Cell culture**

Primary cultures of hippocampal and cortical neurons were prepared as previously described (Banker and Goslin, 1988). Briefly, the hippocampus and the cortex were dissected and dissociated from E18 mouse embryos using the Papain Dissociation System (Worthington Biochemical, Lakewood, NJ). For axonal growth experiments and electroporation, neurons were plated at a density of 10,000 cells/cm² on poly-l-lysine-coated coverslips (1 mg/ml; Sigma-Aldrich), and for measurement of ATP levels or cell transfection, neurons were plated at 100,000 cells/cm² on coverslips or 35-mm plates coated with polylysine (10 μg/ml; Biochrom, Berlin, Germany) and laminin (3 μg/ml; Sigma-Aldrich). After plating, neurons were cultured for 3 d in Neurobasal medium (Life Technologies, Gaithersburg, MD) supplemented with 1% B-27, 0.5 mM glutamine, 1 mM pyruvate, 100 U/ml penicillin, and 100 mg/ml streptomycin. Levamisole, an inhibitor of TNAP (Kozlenkov et al., 2004), the P2X7 receptor antagonists BBG and A-438079 (McGaraughty et al., 2007), ADA (Cristalli et al., 2001), and adenosine at different concentrations or the selective A1 adenosine receptor agonist CPA were added to the cultured neurons 3 h after plating, at the concentrations indicated. HEK293T cells were maintained in DMEM (Life Technologies, Barcelona, Spain) supplemented with 10% (vol/vol) fetal calf serum (FCS). Cells were reseeded at 10² cells/cm² 1 d before transfection, after which FCS was reduced to 0.5% (vol/vol).

**ATP measurement**

Wells of 35 mm containing 10⁵ cells and 2 ml culture media were used to measure extracellular ATP. The nucleotide concentration in the neuronal culture supernatant was determined using the ENLITEN ATP Assay System Bioluminescence Detection Kit for ATP Measurement (Promega, Madison, WI) according to the manufacturer’s protocol. Briefly, 20 μl cell culture medium from the upper layer of the well was mixed with 100 μl luciferin/luciferase reagent.

**FIGURE 7:** Modulation of axonal growth by expression of TNAP and P2X7. Hippocampal neurons were transfected at 1 DIV and fixed after 3 DIV. (A–F) GFP fluorescence images of representative neurons transfected with pEGFP (A), P2X7 IRES (B, C), shRNA TNAP (D), shRNA P2X7 (E), or P2X7 shRNA plus TNAP shRNA (F) in the presence (C) or absence of exogenous ALP. Scale bar, 50 μm. (G and H) Analysis of axonal length corresponding to the experiments shown in A–F and P2X7 IRES plus TNAP-IRES. The 100% value for total axonal length of EGFP-transfected neurons corresponds to 310.8 ± 28 μm. Values represent mean ± SEM (n = 5) with at least 12 neurons analyzed in each experiment. *p < 0.05 and **p < 0.01 vs. control (one-way ANOVA). (I and J) Quantification of the mRNA levels of TNAP and P2X7 by quantitative RT-PCR in hippocampal neurons cultured in the presence or absence of 1 μM BBG (I) and 4 U/ml ALP (J). Results were normalized to values obtained for β-actin mRNA and expressed as mean ± SEM (n = 3). *p < 0.05 (unpaired t test).
Plasmid constructs and the design of shRNAs for the P2X<sub>R</sub>

Plasmid construction and shRNA plasmid design were as described previously (Diaz-Hernandez et al., 2008). Murine TNAP cDNA (Accession BC065175) was purchased from Thermo Scientific (Huntsville, AL) in a nonexpression vector. For expression in eukaryotic cells, the TNAP sequence was obtained from the original vector using EcoRI and NotI and cloned into EcoRI–XmaI sites of pIRRES2-GFP vector (Clontech, Mountain View, CA). Positive clones were sequenced to confirm the correct insertion nucleotide sequence. Mouse P2X<sub>R</sub> cDNA was isolated using a commercial plasmid (Geneservice, Nottingham, UK) and cloned into the bicistronic plasmid pIRRES2-enhanced GFP (EGFP) (Clontech) for expression in mammalian cells. To construct the P2X<sub>R</sub>–GFP plasmid, P2X<sub>R</sub> was cloned into the pd2EGFP-N1 vector (Clontech), and the correctness of the ligation product was confirmed by sequencing. P2X<sub>R</sub> receptor knockdown was achieved by RNA interference using a vector-based shRNA approach. The shRNA target sequence 5′-GTTTTGACATCCTGGTTTT-3′ for the P2X<sub>R</sub> receptor was selected according to a previously reported rational design protocol. As a control (shRNA Luc), the firefly luciferase-targeted oligonucleotide 5′-CTGACGGAGATCTTGCGAGG-3′ was applied. The specificity of the sequence was confirmed by a BLAST analysis of the human, mouse, and rat P2X<sub>R</sub> receptors. Synthetic forward and reverse 64-nucleotide oligonucleotides (Sigma Genosys, Dorset, UK) were designed, annealed, and inserted into the BglII/HindIII sites of the pSUPER.neo.GFP vector (OligoEngine, Seattle, WA) following the manufacturer’s instructions. These constructs express 19-base pair 9-nucleotide stem-loop shRNAs targeted against P2X<sub>R</sub> receptor or luciferase (control shRNA) mRNAs. shRNA TNAP plasmids were purchased from SABiosciences (Frederick, MD). The concomitant expression of GFP by this vector allowed the identification of transfected cells by fluorescence.

Site-directed mutagenesis

To assess whether murine TNAP shRNA was specific, wild-type (TNAP) or mutant (TNAPmut) forms of murine TNAP full-length cDNA were used. TNAPmut was obtained by PCR site-directed mutagenesis using PCR forward and reverse 61-nucleotide oligonucleotides carried si- lent third codon–base point mutations within the murine cDNA of corresponding TNAP shRNA target sequence 5′-TGGACTAGAACTAATAAGCAT-3′ (mutated nucleotides underlined).

Cell transfection

HEK293T cell transfections were performed with the pSUPERneo-GFP–derived plasmid constructs, pIRES2-TNAP-GFP, and shRNA TNAP using Lipofectamine 2000 (Invitrogen, San Francisco, CA) according to the manufacturer’s instructions. After 6 h, the medium was removed and the cells were further incubated for 24 h in the presence of culture medium.

Neuronal transfection was performed with Lipofectamine 2000 or electroporation (Amaza Biosystems, Gaithersburg, MD) as indicated.

Lipofectamine transfection was carried out 24 h after plating using Lipofectamine 2000 (9 μl), 3 μg control shRNA Luc or shRNA P2X<sub>R</sub>, shRNA TNAP, P2X<sub>R</sub>–GFP, and pIRE2-P2X<sub>R</sub>–EGFP vectors. The transfection mix was removed after 4 h, and the neurons were washed and maintained for 3 DIV after transfection.

Electroporation was performed with Nucleofector II Kit (Amaza Biosystems) according to the manufacturer’s instructions. Briefly, dissociated hippocampal cells were incubated in 100 ml neuron nucleofection solution for primary mammalian neural cells, mixed with plasmids (1.5 μg), and electroporated using the fixed program (O-05) for optimal mouse neuronal transfection. Cells were then quickly centrifuged, resuspended, and plated.

Enzyme cytochemistry

Activity of ALP on hippocampal neurons was detected as previously described (Langer et al., 2008) using BCIP/NBT (0.35 mM BCIP, 0.37 mM NBT, 5 mM MgCl<sub>2</sub>, 100 mM Tris buffer, pH 9.5, 45 min) as a precipitating substrate according to the manufacturer’s protocol (Sigma-Aldrich). In brief, prefixed cells were rinsed with 0.1 M Tris-HCl buffer (pH 7.6) and incubated with substrate to obtain optimal staining intensity. After washing with Tris-HCl buffer, coverslips were embedded in antifading solution (glycerol gelatin; Sigma-Aldrich).

TNAP activity assay

Cultured neurons were washed with phosphate-buffered saline (PBS), lysed, and homogenized in 10 mM Tris-HCl buffer (pH 8.0) supplemented with 0.25 M sucrose and Complete Protease Inhibitor Cock-tail Tablets (Roche Diagnostics). Aliquots of the homogenates were assayed at 25°C in the following reaction mix: 0.2 M diethanolamine buffer (Sigma-Aldrich) at pH 9.8, 1 mM MgCl<sub>2</sub>, and 5 mM p-nitrophenyl phosphate (Merck, Whitehouse Station, NJ) in the presence or in the absence of 5 mM levamisole. Reactions were stopped after 20 min with 0.1 M NaOH. Specific ALP activity was determined as the absorbance at 405 nm of the liberated p-nitrophenol minus the absorbance of the p-nitrophenol formed in the presence of levamisole (arbitrary units) and normalized to cellular protein content.

Antibodies and fluorescent reagents

The following antibodies were applied: rabbit polyclonal anti-MAP2 (Sigma-Aldrich), mouse monoclonal anti–Tau-1 (Chemicon, Temecula, CA), rabbit polyclonal anti-TNAP (Abcam, Cambridge, UK), rabbit polyclonal anti-P2X<sub>R</sub> (Alomone, Jerusalem, Israel, and Chemi-con), and mouse monoclonal anti–βIII-tubulin (Promega). Phalloidin Alexa Fluor 594–conjugated phalloidin, goat anti–mouse Alexa Fluor 594 conjugate, and goat anti–rabbit Alexa Fluor 488 conjugate were from Invitrogen.

Immunofluorescence analysis

Immunofluorescence was performed on hippocampal neurons cultured for 3 DIV fixed with 4% paraformaldehyde. Nonspecific binding was blocked with 1% bovine serum albumin (BSA), 5% fetal bovine serum and 0.2% Triton X-100 in PBS. The cells were then incubated with primary antibodies; anti–TNAP (1/50); anti–P2X<sub>R</sub> (1/100), anti–Tau-1 (1/500), anti MAP2 (1/200) and anti–βIII-tubulin (1/1000) for 1 h at room temperature. Coverslips were washed three times with 1% BSA in PBS and incubated with Alexa-Fluor-488 or Alexa-Fluor-594-conjugated secondary antibodies (1/400) and Alexa-Fluor-594-conjugated phalloidin (1/400). Coverslips were mounted using FluorSave (Calbiochem, Nottingham, UK) and images were acquired using a Nikon TE-200 fluorescence microscope coupled to a Kappa DX2 camera. Analysis of axon length and ramifications were carried out using the Image J software v.1.41o. Images were processed and presented using Adobe Photoshop and Illustrator CS3.

Immunoblotting

Cultured neurons were washed with PBS, lysed, and homogenized in a radioimmunoprecipitation assay lysis buffer (50 mM Tris-HCl, pH
Quantitative PCR experiments
Total DNA was extracted from cultured neurons and HEK-transfected cells using an RNeasy Plus Mini Kit (Qiagen, Hilden, Germany), following the manufacturer’s instructions. RNA isolation and reverse transcription (RT) reactions were performed as described previously (Diaz-Hernandez et al., 2009). Quantitative real-time PCRs were performed using gene-specific primers and TaqMan MGB probes for the mouse P2X receptor, TNAP, glyceraldehyde-3-phosphate dehydrogenase (GAPDH), and β-actin (all from Applied Biosystems, Nutley, NJ). Fast thermal cycling was performed using a StepOnePlus Real-Time PCR System (Applied Biosystems) as follows: denaturation, one cycle of 95°C for 20 s, followed by 40 cycles each of 95°C for 1 s and 60°C for 20 s. The results were normalized as indicated by parallel amplification of the endogenous controls β-actin or GAPDH.

Statistics
All experiments were repeated at least three times, and the results are presented as mean ± standard error of the mean (SEM). Statistical differences were analyzed as indicated using GraphPad Prism 5 software and the unpaired t test or the one-way analysis of variance (ANOVA) test followed by the Newman–Keuls test correction.

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