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TNAP is encoded in humans by the ALPL (alkaline phosphatase, liver/bone/kidney) gene and by the Akp2 (alkaline phosphatase 2) gene in mice, both with 12 exons. In both species, two different transcripts derived from the same coding region have been described.

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ALPL - Wikipedia

Mutations in the ALPL gene lead to varying low activity of the enzyme tissue-nonspecific alkaline phosphatase (TNSALP) resulting in hypophosphatasia (HPP). There are different clinical forms of HPP which can be inherited by an autosomal recessive trait or autosomal dominant trait, [8] the former causing more severe forms of the disease.

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Indeed, parallel neuronal knockdown of P2X₇ 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 P2X₇ and TNAP (Figure 7G).