

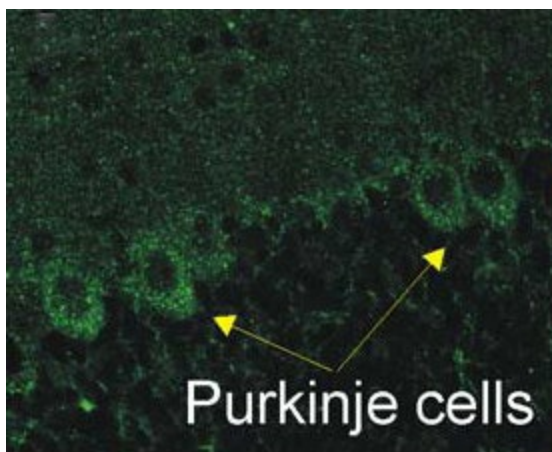
## Product datasheet for **GP14106-150**

### **P2X2 (P2RX2) Guinea Pig Polyclonal Antibody**

#### **Product data:**

Product Type:	Primary Antibodies
Applications:	IF, IHC, WB
Reactivity:	Human, Monkey, Rat
Host:	Guinea Pig
Clonality:	Polyclonal
Formulation:	State: Serum
Gene Name:	Homo sapiens purinergic receptor P2X 2 (P2RX2), transcript variant 6
Database Link:	<a href="#">Entrez Gene 114115 Rat</a> <a href="#">Entrez Gene 22953 Human</a>
Synonyms:	P2X purinoceptor 2, ATP receptor, Purinergic receptor
Protein Families:	Druggable Genome, Ion Channels: ATP Receptors, Transmembrane
Protein Pathways:	Calcium signaling pathway, Neuroactive ligand-receptor interaction

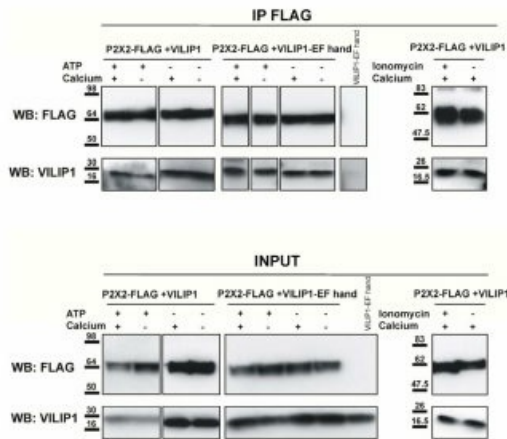
#### **Product images:**



P2x2 staining of Cerebellar Purkinje cell layer. ~50  $\mu$ m sections from 3 brains from P15 rats.  
<http://stke.sciencemag.org/cgi/data/sigtrans;1/41/ra8/DC1/1>

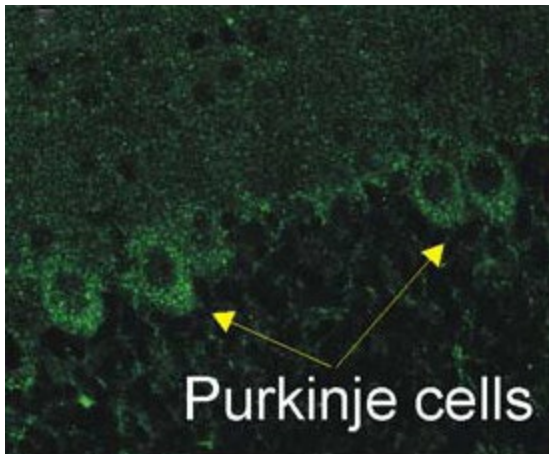


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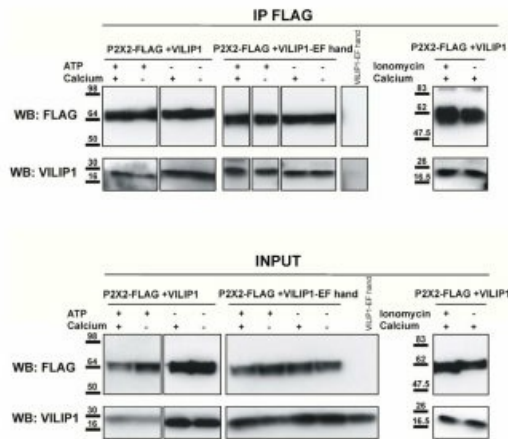


The constitutive interaction between P2X2 receptors and VILIP1 is calcium independent. Calcium dependence of the interaction between P2X2 receptors and VILIP1 was assessed by coimmunoprecipitation from HEK cells expressing P2X2-FLAG receptors and either VILIP1 or VILIP1 EF-hand mutants. The effect of calcium on the interaction was tested before cell lysis by increasing intracellular calcium with ATP application or ionomycin treatment, and also during the immunoprecipitation steps. In the latter case, experiments were carried in the absence of calcium and in the presence of 5 mM EDTA or in the absence of EDTA and in the presence of 500µM calcium. P2X2 receptors were immunoprecipitated using a FLAG antibody conjugated to agarose beads. Bound proteins were eluted by competition using a FLAG peptide. As shown in the top panel, no difference in P2X2-FLAG-VILIP1 interaction was observed in the presence or the absence of calcium in the immunoprecipitation buffer. In addition, stimulation of transfected cells with ATP (100µM, 5 minutes) or with ionomycin (2µM, 5 minutes) prior to lysis did not affect the interaction between the two proteins. A similar observation was made when the VILIP1 EF hand mutant was co-immunoprecipitated. No immunoprecipitation was observed when VILIP1 was transfected alone. The bottom panel shows the total protein input used for immunoprecipitation.

[www.sciencesignaling.org/cgi/content/full/1/41/ra8/DC1](http://www.sciencesignaling.org/cgi/content/full/1/41/ra8/DC1)



P2x2 staining of Cerebellar Purkinje cell layer. ~50 µm sections from 3 brains from P15 rats.  
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[www.sciencesignaling.org/cgi/content/full/1/41/ra8/DC1](http://www.sciencesignaling.org/cgi/content/full/1/41/ra8/DC1)