

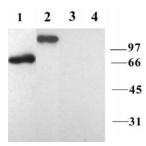
RABBIT ANTI-P2Y1 RECEPTOR (P2RY1) AFFINITY PURIFIED, POLYCLONAL ANTIBODY

CATALOG NUMBER:	AB5814-50UL
LOT NUMBER:	
QUANTITY:	50 μL
CONCENTRATION:	0.8 mg/mL (after reconstitution)
SPECIFICITY:	Recognizes P2Y1 Receptor. The epitope does not share homology with any other known proteins.
IMMUNOGEN:	Synthetic peptide corresponding to amino acids 242-258 of rat or human P2Y1 (SwissProt accession numbers P49651, P47900).
APPLICATIONS:	Western blot: 1:200-1:400 using ECL on rat brain membranes or human platelets. Immunohistochemistry on rat brain sections. Dilutions should be made using a carrier protein such as BSA (1-3%) Optimal working dilutions must be determined by the end user.
CONTROL ANTIGEN:	Included free of charge with the antibody is XX μ g of control antigen (lyophilized powder). The stock solution of the antigen can be made up using 100 μ L of sterile distilled water. For negative control, preincubate 1 μ g of protein with 1 μ g of antibody for one hour at room temperature. Optimal concentrations must be determined by the end user.
SPECIES REACTIVITIES:	Human and rat. It is expected that the antibody may also react with bovine (17/17), chicken and turkey (16/17) and possibly mouse (15/17) due to sequence homology. Other species have not been tested.
FORMAT:	Affinity purified immunoglobulin.
PRESENTATION:	Lyophilized from phosphate buffered saline, pH 7.4, containing 1% BSA and 0.05% sodium azide as a preservative. Reconstitute with 50 μ L of sterile deionized water. Centrifuge antibody preparation before use (10,000 xg for 5 min).
STORAGE/HANDLING:	Maintain lyophilized material at -20°C for up to 12 months after date of receipt. After reconstitution maintain at -20°C in undiluted aliquots for up to 6 months. Avoid repeated freeze/thaw cycles.



SUGGESTED WESTERN BLOT PROTOCOL

- 1. Mix the samples (organ membranes: 50 μg/lane; transfected cells: 500,000 cells/lane) with samplebuffer X 2, and heat 10 min at 70°C.
- 5-50 μL applied to Minigel lane (0.75-1.5 mm width) and run at standard conditions. (60 mA for 2 1.5 mm Minigel gels, 1.4 h). It is suggested that you run 5-15% acrylamide (37.5:1 acrylamide:bisacrysmide) minigel (1.5 mm width) at 30 mA/gel ~1-1.5 hours.
- Transfer in semi-dry system under standard conditions (3 h 100 mA for two minigel gels)
- 4. Stain the transferred bands with Millipore BLOT-*FastStain* (Catalog Number 2076).
- 5. Destain with deionized water.
- Block with 5% non-fat milk (Marvel or Carnation) in PBS, and 0.025 % sodium azide, overnight at 2-8°C. The non-fat milk should be dissolved freshly, centrifuged 10,000 rpm for 10 min, and filtered through glass filter (Gelman Acrodisc).
- Incubation with first antibody 2 h at room temperature or overnight at 4°C in blocking solution. The antibody preparation should be centrifuged before use (10,000 g 5 min.). Optimal working dilutions and incubation time will need to be determined by the end user.
- 8. Wash 4 x 10 min. with PBS-0.1% tween 20. From this stage, azide should be omitted.
- 9. Incubation with the secondary antibody (HRP-conjugated goat anti-rabbit antibody, for example Millipore Catalog Number AP132P, diluted appropriately) 1 h at room temperature.
- 10. Wash 4 x 10 min. with PBS-0.1% tween 20.
- 11. Perform ECL with commercial kits (ChemiLUCENT, Millipore Catalog Number 2600).



Western blotting of rat brain membranes (lanes 1,3) or human platelets lysate (lanes 2,4).

1, 2. AB5814 (1:200)

3, 4. AB5814, preincubated with the control peptide antigen.

Important Note: During shipment, small volumes of product will occasionally become entrapped in the seal of the product vial. For products with volumes of 200 μ L or less, we recommend gently tapping the vial on a hard surface or briefly centrifuging the vial in a tabletop centrifuge to dislodge any liquid in the container's cap.

FOR RESEARCH USE ONLY; NOT FOR USE IN DIAGNOSTIC PROCEDURES. NOT FOR HUMAN OR ANIMAL CONSUMPTION

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