

RABBIT ANTI-Kv3.4, VOLTAGE-GATED POTASSIUM CHANNEL **AFFINITY PURIFIED** POLYCLONAL ANTIBODY

CATALOG NUMBER: AB5192-50UL

LOT NUMBER:

QUANTITY: 50 μL

CONCENTRATION: 0.3 mg/mL (after reconstitution)

SPECIFICITY: Recognizes a full length Kv3.4 protein. Does not cross react with any other potassium

channel antigens tested so far.

IMMUNOGEN: Purified peptide from rat Kv3.4 (amino acids 177-195) (Accession Q63734).

Western blot: 1:200 using ECL on rat brain membranes. **APPLICATIONS:**

Immunohistochemistry on rat brain tissue 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 µg of control antigen (lyophilized

> powder). The stock solution of the antigen can be made up using 100 μL of sterile deionized water. For negative control, preincubate 1 µg of peptide with 1 µg of antibody for one hour at room temperature. Optimal concentrations must be determined by the

end user.

SPECIES REACTIVITIES: Rat. The immunogen sequence is identical in human. Other species have not been

FORMAT: Affinity purified immunoglobulin.

Lyophilized from phosphate buffered saline, pH 7.4, containing 1% BSA, 5% sucrose as PRESENTATION:

> a stabilizer and 0.025% sodium azide as a preservative. Reconstitute with 50 µL 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.

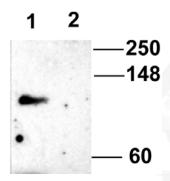
REFERENCES: 1) Gopel, S.O. et al. (2000) J. Physiol. **528**:497.



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SUGGESTED WESTERN BLOT PROTOCOL

- 1. Mix the samples with sample-buffer X 2, and heat 10 min at 70°C.
- 2. 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 aels. 1.4 h)
- Transfer in semi-dry system under standard conditions (3 h 200 mA for two minigel gels)
- 4. Stain the transferred bands with Chemicon BLOT-FastStain (Catalog Number 2076).
- 5. Destain with deionized water.
- 6. Block with 5% non-fat milk (Marvel or Carnation) in PBS, and 0.025 % sodium azide, overnight at 4°C. The non-fat milk should be dissolved freshly, centrifuged 10,000 rpm for 10 min, and filtered through glass filter (Gelman Acrodisc).
- 7. 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.
- Wash 4 x 15 min. with PBS-0.1% tween 20. From this stage, azide should be omitted.
- Incubation with the secondary antibody (HRP-conjugated goat anti-rabbit antibody, for example Chemicon Catalog Number AP132P, diluted appropriately) 1 h at room temperature.
- 10. Wash 4 x 15 min. with PBS-0.1% tween 20.
- 11. Perform ECL with commercial kits (ChemiLUCENT, Chemicon Catalog Number 2600).



Western blotting of rat brain membrane proteins (25 µg)

- AB5192 1:200
- AB5192 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 µL 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

Unless otherwise stated in our catalog or other company documentation accompanying the product(s), our products are intended for research use only and are not to be used for any other purpose, which includes but is not limited to, unauthorized commercial uses, in vitro diagnostic uses, ex vivo or in vivo therapeutic uses or any type of consumption or application to humans or animals





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