



IHC image of neurons in rat striatum.

## NK1R (Neurokinin 1 Receptor) Antibody

<b>Catalog #</b>	20060	<b>Product type</b>	Primary antibodies
<b>Lot #</b>	902001	<b>Clonality</b>	Polyclonal
<b>Form</b>	Lyophilized Whole Serum (100 µL)	<b>Isotype</b>	N/A
<b>Host</b>	Rabbit	<b>Preservative</b>	≤ 0.09% sodium azide
<b>Reacts With</b>	Rat	<b>Antigen</b>	Synthetic peptide corresponding to rat NK1R (393–407) coupled to carrier protein.

### INSTRUCTIONS

<b>Preparation</b>	<p>Do not reconstitute until ready to use since the product is most stable when lyophilized. The product does not need to be cooled during shipping; however, for long-term storage, store lyophilized antibody until ready to use at -15°C or lower. Reconstitute with 100 µL of distilled or deionized water. After reconstitution, use immediately or refrigerate at 2°–8° C. To avoid freeze/thaw cycles, dilute unused antibody with PBS or Tris buffer at a dilution no higher than 1/10, then aliquot and freeze at -15°C or lower.</p> <p>Refer to the Instruction Manual available online at <a href="http://www.immunostar.com">www.immunostar.com</a> for information on tissue preparation, immunostaining techniques, troubleshooting, and formulas.</p>
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### APPLICATION

<b>Quality Control</b>	The ImmunoStar NK1R antiserum was quality control tested using standard immunohistochemical methods in rat brain using biotin/avidin-HRP techniques. Specificity of the antiserum was demonstrated by soluble preadsorption and western blot. Tissue staining is completely eliminated by pretreatment of the diluted antibody with 25 µg of rat NK1R peptide residues (393–407). Western blot analysis of crude rat brain homogenate demonstrates two immunoreactive bands of approximately 70 and 110 kD.
<b>Tissue</b>	Rat brain - striatum and cortex
<b>Absorption Control</b>	Rat NK1R (393–407) 10 µg/mL diluted antibody completely eliminates immunolabeling
<b>Perfusion Fixation</b>	<ul style="list-style-type: none"> <li>Fixative: 4% paraformaldehyde in 0.1 M Phosphate buffer, pH 7.4; 500 mL over 20 min.</li> <li>Post Fixation: 1.5 hour at 4°C in 4% paraformaldehyde in 0.1 M phosphate buffer, pH 7.4.</li> <li>Note: Paraformaldehyde is a necessary component in fixation. If needed, low levels of glutaraldehyde (0.1–0.3%) may be used in conjunction with paraformaldehyde.</li> </ul>
<b>Sections</b>	10 µm cryostat or 50 µm vibratome
<b>Tissue Incubation</b>	48 hours at 2°–8°C
<b>Detection System</b>	Bn/AV-HRP at dilutions recommended by the manufacturers.
<b>Suggested Dilution</b>	1/3,000–1/5,000 in PBS/0.3% Triton X-100 - Bn/AV-HRP immunohistochemistry

### NOTES

<b>Special Instructions</b>	It is recommended that users perform a primary antibody dilution series using the dilution recommendations above as a guideline. Note that a change in the fixation or buffering system from our protocol may change the configuration of the protein which could alter the reactivity with the tissue tested.
<b>Storage</b>	After reconstitution, use immediately or refrigerate at 2°–8°C up to 2 days. For long-term storage, aliquot antibody and freeze at -15°C. or lower. Avoid repeated freeze/thaw cycles.
<b>Concentration</b>	Not applicable. Antibody concentration is only relevant for purified antibodies.
<b>Journal Articles</b>	<a href="http://www.immunostar.com/publications">www.immunostar.com/publications</a>

*For Laboratory Reagent Use Only. Analytical and performance characteristics are not established.*

**ALL PRODUCTS ARE FOR RESEARCH USE ONLY AND ARE NOT INTENDED FOR DIAGNOSTIC OR THERAPEUTIC USE**