



IHC image of neurons in rat hypothalamus.

## Alpha-MSH (Melanocyte Stimulating Hormone) Antibody

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| <b>Catalog #</b>   | 20074  | <b>Product type</b> | Primary antibodies   |
| <b>Lot #</b>       | 1340001  | <b>Clonality</b>    | Polyclonal   |
| <b>Form</b>        | Lyophilized whole serum (100 µL)   | <b>Isotype</b>      | IgG  |
| <b>Host</b>        | Rabbit   | <b>Preservative</b> | ≤ 0.09% sodium azide   |
| <b>Reacts With</b> | Cat, Frog, Guinea Pig, Hamster, Human, Monkey, Mouse, Rabbit, Rat, Trout | <b>Antigen</b>      | Synthetic human α-MSH coupled to bovine thyroglobulin with glutaraldehyde. |

### INSTRUCTIONS

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| <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 kept 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

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| <b>IHC Quality Control</b> | The ImmunoStar alpha melanocyte stimulating hormone antiserum was quality control tested using standard immunohistochemical methods. The antiserum demonstrates significant labeling of rat pituitary using indirect immunofluorescent and biotin/avidin-HRP techniques. Staining is completely eliminated by pretreatment of the diluted antibody with 100 µg/mL of α-MSH. |
| <b>Absorption Control</b>  | α-MSH 100 µg/mL diluted serum   |
| <b>Tissue</b>              | Rat pituitary   |
| <b>Perfusion Fixation</b>  | <ul style="list-style-type: none"> <li>Fixative: 4% paraformaldehyde in 0.1M Phosphate buffer, pH 7.4; 500 mL over 20 min.</li> <li>Post Fixation: 1.5 hour at 4°C in 4% paraformaldehyde in 0.1M phosphate buffer, pH 7.4.</li> <li>Note: 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  |
| <b>Tissue Incubation</b>   | 18–24 hours at 2°–8°C.  |
| <b>Detection System</b>    | Use IF or Bn/AV-HRP according to manufacturers' directions.   |
| <b>Suggested Dilution</b>  | 1/4,000–1/6,000 in PBS/0.3% Triton X-100 – Bn/AV-HRP immunohistochemistry   |

### NOTES

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| <b>Special Instructions</b> | It is recommended that the researcher perform a primary antibody dilution series using our dilution recommendations 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. Please read the instruction booklet carefully before beginning the procedure. |
| <b>Storage</b>              | After reconstitution, use immediately or refrigerate at 2°–8°C up to 2 days. For long-term storage, aliquot 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 References</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.*

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