NEUROMICS

ACTH

Data Sheet

Catalog Number:	RA20070	Host:	Rabbit
Product Type:	Whole Serum	Species Reactivity:	Dove, Eel, Goldfish, Hamster, Human, Monkey, Mouse, Ram, Rat, Trout
Immunogen Sequence:	Adrenocorticotropic hormone (ACTH) corresponding to amino acids (1-39) generated against porcine pituitary	Format:	100ul Lyophilized, ≤ 0.09% sodium azide
Applications:	Immunohistochemistry: 1:2,000–1:4,000 in PBS/0.3% Triton X-100 – Bn-AV/HRP		
Storage and Preparation:	Storage: Dilute with phosphate buffer or Tris buffer at dilutions no higher than $1/10$, aliquot and freeze at -15° C or lower. Antibody can be stored for up to six months if handled as described above.		
	It is strongly recommended that the customer perform a primary antibody dilution series using our dilution recommendations as a guideline. Note that a change in the fixation or buffering system as used in our protocol may change the configuration of the protein and, therefore, may alter the reactivity with the tissue tested.		

Application Notes for Immunohistochemistry

Tissue: Rat anterior and intermediate pituitary

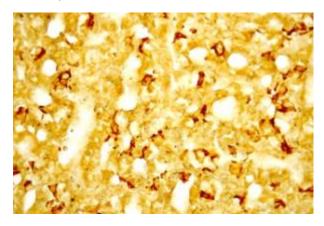
Perfusion Fixation • Fixation: 4% paraformaldehyde in 0.1M phosphate buffer, pH 7.4; 500 mL over 20 min. • Post Fixation: 1.5 hour at 4°C in 4% paraformaldehyde in 0.1M phosphate buffer, pH 7.4. • Note: Paraformaldehyde is a necessary component of fixation for this antiserum. If needed for other applications, glutaraldehyde may be used at low levels (0.1–0.3%) in conjunction with paraformaldehyde.

Sections 10 µm cryostat or 50 µm vibratome

Tissue Incubation 18-24 hours at 2°-8°C.

Detection System Bn-Av/HRP reagents at dilutions recommended by the manufacturer

Image: IHC staining for ACTH in the rat anterior pituitary gland. The tissue was fixed with 4% formaldehyde in 0.1 M phosphate buffer, before being removed and prepared for frozen sectioning. Sections were incubated at RT in 10% goat serum in PBS, before standard IHC procedure. Primary antibody was incubated at 1:1000 for 48 hours, goat anti-rabbit secondary was subsequently added for 1 hour after washing with PBS. Light microscopy staining was achieved with standard biotinstreptavidin/HRP procedure and DAB chromogen



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