

# Mouse GFRα-like Antibody

Antigen Affinity-purified Polyclonal Sheep IgG Catalog Number: AF5728

DESCRIPTION		
Species Reactivity	Mouse	
Specificity	Detects mouse GFRα-like in direct ELISAs and Western blots.	
Source	Polyclonal Sheep IgG	
Purification	Antigen Affinity-purified	
Immunogen	Chinese hamster ovary cell line CHO-derived recombinant mouse GFRα-like isoform A Gln20-Gly349 Accession # Q6SJE0	
Formulation	Lyophilized from a 0.2 µm filtered solution in PBS with Trehalose. See Certificate of Analysis for details. *Small pack size (-SP) is supplied either lyophilized or as a 0.2 µm filtered solution in PBS.	

# APPLICATIONS

Please Note: Optimal dilutions should be determined by each laboratory for each application. General Protocols are available in the Technical Information section on our website.

	Recommended Concentration	Sample
Western Blot	2 μg/mL	See Below
Immunohistochemistry	5-15 μg/mL	See Below

# DATA

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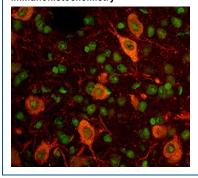
# Detection of Mouse GFRα-like by Western Blot. Western blot shows lysates of Neuro-2A mouse neuroblastoma cell line, mouse brain and mouse brain stem. PVDF membrane was probed with 2 μg/mL of Sheep Anti-Mouse GFRa-like Antigen Affinity-purified Polyclonal Antibody (Catalog # AF5728) followed by HRP-conjugated Anti-Sheep IgG Secondary Antibody (Catalog # Catalog # HAF016). Specific bands were detected for GFRa-like at approximately 25 and 50 kDa (as indicated). This experiment was conducted under reducing conditions and using Immunoblot Buffer Group 1.

#### Immunohistochemistry



GFRα-like in Mouse Spinal Cord. GFRαlike was detected in immersion fixed paraffinembedded sections of mouse spinal cord using Sheep Anti-Mouse GFRα-like Antigen Affinity-purified Polyclonal Antibody (Catalog # AF5728) at 5 µg/mL for 1 hour at room temperature followed by incubation with the Anti-Sheep IgG VisUCyte™ HRP Polymer Antibody (Catalog # VC006). Before incubation with the primary antibody, tissue was subjected to heat-induced epitope retrieval using Antigen Retrieval Reagent-Basic (Catalog # CTS013). Tissue was stained using DAB (brown) and counterstained with DAPI (blue). Specific staining was localized to dorsal and ventral horns. Staining was performed using our protocol for IHC Staining with VisUCyte HRP Polymer Detection Reagents.

# Immunohistochemistry



GFRα-like in Mouse Brain. GFRa-like was detected in immersion fixed frozen sections of mouse brain using Sheep Anti-Mouse GFRa-like Antigen Affinity-purified Polyclonal Antibody (Catalog # AF5728) at 15 µg/mL overnight at 4 °C. Tissue was stained using the NorthernLights ™ 557-conjugated Anti-Sheep IgG Secondary Antibody (red; Catalog # Catalog # NL010) and counterstained with Fluoro NissI Green (green). View our protocol for Fluorescent IHC Staining of Frozen Tissue Sections.

# PREPARATION AND STORAGE

**Reconstitution** Reconstitute at 0.2 mg/mL in sterile PBS.

Shipping The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below.

\*Small pack size (-SP) is shipped with polar packs. Upon receipt, store it immediately at -20 to -70 °C

# Stability & Storage

# Use a manual defrost freezer and avoid repeated freeze-thaw cycles.

- 12 months from date of receipt, -20 to -70 °C as supplied.
- 1 month, 2 to 8 °C under sterile conditions after reconstitution.
   6 months, -20 to -70 °C under sterile conditions after reconstitution.

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Global bio-techne.com info@bio-techne.com techsupport@bio-techne.com TEL +1 612 379 2956 USA TEL 800 343 7475 Canada TEL 855 668 8722 China TEL +86 (21) 52380373 Europe | Middle East | Africa TEL +44 (0)1235 529449



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#### BACKGROUND

GFRα-like (GDNF receptor-alpha-like) is a predicted 26 and 44 kDa glycoprotein (1) and distant member of the GDNFR family of proteins. It is expressed on both fetal and adult neurons of the CNS, and would appear to function as an anti-apoptotic molecule during neuronal stress. Mature mouse GFRα-like is a 374 amino acid (aa) type I transmembrane protein. It contains a 330 aa extracellular region that shows a Gas1 domain (aa 131-210), and a 23 aa cytoplasmic tail. There is one splice variant that is likely soluble and shows a four aa substitution for aa 235-393. Over aa 20-349, mouse GFRα-like shares 72% aa identity with human GFRα-like.

### References:

1. Li, Z. et al. (2005) J. Neurochem. 95:361.

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