Anti-Human GAL Antibody [K24019_15H8]

Catalog No. K24019M15H08C

Overview

Product name: Anti-Human GAL Antibody [K24019_15H8]
Antibody specificity: Galanin and GMAP prepropeptide
Species reactivity: Human
Clonality: Monoclonal
Clone number: K24019_15H8
Host / isotype: Mouse / IgG1
Immunogen: Recombinant Human GAL
Cross reactivity: Not tested
Kinetic characterization by BLI (biolayer interferometry): Not tested
Purification: Protein A purified from cell culture supernatants
Form: Liquid
Concentration: 1 mg/mL in phosphate buffered saline containing 0.09% preservative
Conjugation: Unconjugated

Shipping, storage and shelf life
* 3 months when stored at 2 to 8 °C
* 1 year when aliquoted and stored at -20 °C
* 3 years when aliquoted and stored at -80 °C

Applications

<table>
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<tr>
<th>Application</th>
<th>Recommended concentration</th>
<th>Note</th>
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<tbody>
<tr>
<td>Indirect ELISA</td>
<td>1 µg/mL</td>
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<tr>
<td>Immunohistochemistry-paraffin (IHC-P)</td>
<td>5 µg/mL</td>
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Note:
* The applications above have already been verified. The antibody may be suitable for additional applications.
* Optimal antibody concentrations for each application should be determined by the user.

Additional information

Target antigen
Protein name: Galanin and GMAP prepropeptide
Gene name: GAL
UniProt Accession: P22466
Organism: Homo sapiens (Human)

Paired antibody information
K24019_15H8 may pair with K24019_4D11 and K24019_11A9 for sandwich based immune assays.
Immunohistochemistry

IHC-P analysis of human adrenal gland tissue by anti-human GAL antibody (K24019_15H8).

IHC-P was performed using sections of the formalin-fixed paraffin-embedded human adrenal gland tissue. Antigen was retrieved through addition of boiling Tris/EDTA buffer pH 9 in a pressure cooker for 3 min. Endogenous peroxidase activity was quenched by incubating the sections with 3% H₂O₂ for 30 min at room temperature. The sections were then incubated with anti-human GAL primary antibody (K24019_15H8) at 5 µg/mL at room temperature for 1 h. Poly-peroxidase conjugated goat anti-mouse IgG was used as the secondary antibody. Diaminobenzidine was used as the chromogen. The section was counterstained with hematoxylin. A tissue section incubated with phosphate-buffered saline followed by incubation with the secondary antibody was used as the background control.

Result: Glandular cells are positively stained at the cytoplasm.