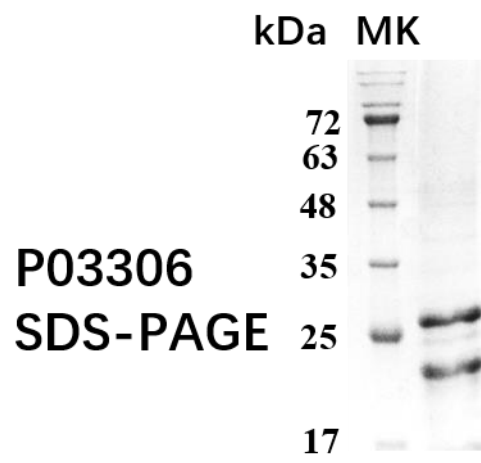


## Recombinant human GM2A protein

Catalog#:P03306    Derived from E.coli

|                        |   |
|------------------------|---|
| <b>DESCRIPTION</b>     | Recombinant Human GM2A protein is produced by our E.coli expression system with His tag.<br>Uniprot: P17900<br>Gene ID: 2760  |
| <b>Size</b>            | 21 kDa  |
| <b>FORMULATION</b>     | Lyophilized from a 0.2 µm filtered solution of Tris-HCl, PH 8.0.  |
| <b>SHIPPING</b>        | The product is shipped at -20°C temperature. Upon receipt, store it immediately at the temperature listed below.  |
| <b>STORAGE</b>         | Reconstituted protein solution can be stored at 4-7°C for 2-3 months, stable at < -20°C for 1-2 years.  |
| <b>RECONSTITUTION</b>  | Reconstituted protein solution can be diluted with distilled PBS.<br>Please aliquot the reconstituted solution to minimize freeze-thaw cycles.  |
| <b>QUALITY CONTROL</b> | (Liquid)Concentration: 1 mg/mL as determined by BCA.<br>Purity: 85% as determined by reducing SDS-PAGE.   |
| <b>AMINOACID</b>       | Recombinant protein is produced by our E.coli expression system and the target gene encoding 1-193aa is expressed with a 6*His tag at the N-terminus.   |
| <b>BACKGROUND</b>      | This gene encodes a small glycolipid transport protein which acts as a substrate specific co-factor for the lysosomal enzyme beta-hexosaminidase A. Beta-hexosaminidase A, together with GM2 ganglioside activator, catalyzes the degradation of the ganglioside GM2, and other molecules containing terminal N-acetyl hexosamines. Mutations in this gene result in GM2-gangliosidosis type AB or the AB variant of Tay-Sachs disease. Alternative splicing results in multiple transcript variants. |



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