

## InnoCyto Inc.

15375 Barranca Pkwy, Suite I-103 Irvine, CA 92618

## **Technical Data Sheet**

Human GM2A Protein (C-His)

Catalog Number: 601401, 601402

Size: 25 ug, 100 ug

Target Name: GM2A, Ganglioside GM2 activator

Regulatory Status: RUO

#### **Product Details**

Application: ELISA, BLI Format: Liquid, Purified Expression Host: HEK293

Species: Human

Accession Number: P17900

Sources: Recombinant human GM2A ( (His24-Ile193) with C-terminus His tag was expressed in 293

Cells

Molecular Weight: This protein has a predicted molecular weight of 20 kDa. Under DTT-reducing

conditions, the protein migrates at approximately 24 kDa on SDS-PAGE.

Affinity Tag: C-His

**Purity:** >95% based on SDS-PAGE under reducing condition **Formulation:** 1xPBS with 300mM NaCl, pH7.4, 0.22 μm filtered

Endotoxin level: Not tested

Protein Concentration: 25µg size is bottled at 0.2mg/mL concentration. 100 µg size is supplied at a

lot-specific concentration.

**Storage and Handling:** Briefly centrifuge the vial upon receipt. An unopened vial can be stored at  $4^{\circ}$ C for up to 2 weeks, or at -20°C or below for up to six months. The protein may be further diluted to 0.1 mg/mL using 0.22 µm-filtered PBS buffer (pH 7.4). For long-term storage, the diluted stock solution should be aliquoted and stored at <=  $-70^{\circ}$ C to minimize freeze-thaw cycles. If additional dilution is required, carrier proteins such as FBS or BSA should be added to maintain protein stability.

#### **Background Information**

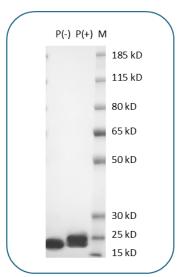
GM2 ganglioside activator (GM2A) is a lipid transfer protein of the ML domain family that facilitates the degradation of ganglioside GM2 by acting as a substrate-specific cofactor for ?-hexosaminidase A. GM2A extracts GM2 from membranes and presents it in soluble form to ?-hexosaminidase A for cleavage, enabling conversion to GM3. It can bind several single-chain phospholipids and fatty acids and shows some calcium-independent phospholipase activity. Mutations in GM2A cause GM2-gangliosidosis type AB, a Tay-Sachs disease variant characterized by impaired GM2 degradation and lipid accumulation.

### **Product Data**



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Purified Human GM2A (His24-Ile193, with C-terminus His tag, 293 expressed) final product on SDS-PAGE under reducing (P+) conditions. The purity of Human GM2A appears to be greater than 95%.