

Product datasheet for TP727231

PLA2G1B Human Recombinant Protein

Product data:

Product Type: Recombinant Proteins

Description: Recombinant Human PLA2G1B/PLA2/PLA2A (C-6His)

Species: Human

Expression cDNA Clone

or AA Sequence:

Ala23-Ser148

Tag: C-His

Buffer: Supplied as a 0.2 um filtered solution of 20mM Tris-HCl, 150mM NaCl, 10% Glycerol, pH 8.0.

Note: Recombinant Human Phospholipase A2 Group IB is produced by our Mammalian expression

system and the target gene encoding Ala23-Ser148 is expressed with a 6His tag at the C-

terminus.

Storage: Store at < -20°C, stable for 6 months after receipt. Please minimize freeze-thaw cycles.

Stability: 12 months from date of despatch

Locus ID: 5319 **UniProt ID:** P04054

Synonyms: PLA2G1B;Phospholipase A2;Group IB phospholipase A2;PLA2A;PPLA2

Summary: Phospholipase A2(PLA2G1B) is a secreted protein which belongs to the phospholipase A2

family. It catalyzes the release of fatty acids from glycero-3-phosphocholines. It catalyzes the calcium-dependent hydrolysis of the 2-acyl groups in 3-sn-phosphoglycerides. This releases glycerophospholipids and arachidonic acid that serve as the precursors of signal molecules. Sequences of pancreatic PLA2G1B enzymes from a variety of mammals have been reported. One striking feature of these enzymes is their close homology to venom phospholipases of snakes. Mice lacking in PLA2G1B are resistant to obesity and diabetes induced by feeding a diabetogenic high-fat/high-carbohydrate diet. Oral supplementation of a diabetogenic diet with the PLA2G1B inhibitor methyl indoxam effectively suppresses diet-induced obesity and diabetes. PLA2G1B inhibition may be a potentially effective oral therapeutic option for

treatment of obesity and diabetes.

Protein Families: Druggable Genome, Secreted Protein



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Protein Pathways:

alpha-Linolenic acid metabolism, Arachidonic acid metabolism, Ether lipid metabolism, Fc epsilon RI signaling pathway, Glycerophospholipid metabolism, GnRH signaling pathway, Linoleic acid metabolism, Long-term depression, MAPK signaling pathway, Metabolic pathways, Vascular smooth muscle contraction, VEGF signaling pathway