

Recombinant Human VAP-1/AOC3 Protein (Fc Tag)

Catalog Number:PKSH033590



Note: Centrifuge before opening to ensure complete recovery of vial contents.

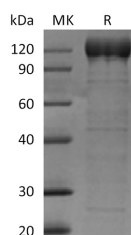
Description

Synonyms	Membrane primary amine oxidase;Copper amine oxidase;Semicarbazide-sensitive amine oxidase;Vascular adhesion protein 1;AOC3;VAP-1;SSAO;HPAO;VAP1
Species	Human
Expression Host	HEK293 Cells
Sequence	Arg28-Asn763
Accession	Q16853
Calculated Molecular Weight	108.5 kDa
Observed molecular weight	120 kDa
Tag	C-Fc

Properties

Purity	> 95 % as determined by reducing SDS-PAGE.
Endotoxin	< 1.0 EU per µg of the protein as determined by the LAL method.
Storage	Generally, lyophilized proteins are stable for up to 12 months when stored at -20 to -80°C. Reconstituted protein solution can be stored at 4-8°C for 2-7 days. Aliquots of reconstituted samples are stable at < -20°C for 3 months.
Shipping	This product is provided as lyophilized powder which is shipped with ice packs.
Formulation	Lyophilized from a 0.2 µm filtered solution of 20mM Tris-HCl, 500mM NaCl, pH 8.0. Normally 5 % - 8 % trehalose, mannitol and 0.01% Tween80 are added as protectants before lyophilization. Please refer to the specific buffer information in the print
Reconstitution	Please refer to the printed manual for detailed information.

Data



> 95 % as determined by reducing SDS-PAGE.

Background

Membrane primary amine oxidase(AOC3); also known as vascular adhesion protein (VAP-1) and HPAO; this protein is a member of the semicarbazide-sensitive amine oxidase (SSAO) family. VAP-1 is a type 1 membrane-bound glycoprotein that has a distal adhesion domain and an enzymatically active amine oxidase site outside of the membrane; VAP-1 has adhesive properties; functional monoamine oxidase activity; and possibly plays a role in glucose handling; leukocyte trafficking; and migration during inflammation. This rise in metabolic products contributes to generating advanced glycation end-products and oxidative stress along with the monoamine detoxification in the organism. It is highly

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expressed on the endothelium of the lung and trachea; and absent from leukocytes and epithelial cells. Membrane-bound VAP-1 releases an active; soluble form of the protein; which may be conducive to increased inflammation and the progression of many vascular disorders. In particular; elevation of VAP-1 activity and the increased enzymatic-mediated deamination is proposed to play a role in renal and vascular disease; oxidative stress; acute and chronic hyperglycemia; and diabetes complications.

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