

APT-1/LYPLA1 Protein, Human (His)

Cat. No.:	HY-P7539
Synonyms:	rHuAPT-1, His; LYPLA1; APT-1
Species:	Human
Source:	E. coli
Accession:	O75608 (M1-D230)
Gene ID:	10434
Molecular Weight:	Approximately 25.0 kDa

PROPERTIES

AA Sequence	<pre> MCGNNMSTPL PAIVPAARKA TAAVIFLHGL GDTGHGWAEA FAGIRSSHIAK YICPHAPVRP VTLNMNVAMP SWFDIIGLSP DSQEDESIGIK QAAENIKALI DQEVKNGIPS NRIILGGFSQ GGALSLYTAL TTQQKLAGVT ALSCWLPLRA SFPQGPIGGA NRDISILQCH GDCDPLVPLM FGSALTVEKLK TLVNPANVTF KTYEGMMHSS CQQEMMDVKQ FIDKLLPPID HHHHHH </pre>
Biological Activity	The enzyme activity of this recombinant protein is testing in progress, we cannot offer a guarantee yet.
Appearance	Solution.
Formulation	Supplied as a 0.2 µm filter solution of 20 mM Tris-HCl, 100 mM NaCl, 1 Mm DTT, 10% Glycerol, pH 8.0.
Endotoxin Level	<1 EU/µg, determined by LAL method.
Reconstitution	N/A
Storage & Stability	Stored at -80°C for 1 year. It is stable at -20°C for 3 months after opening. It is recommended to freeze aliquots at -80°C for extended storage. Avoid repeated freeze-thaw cycles.
Shipping	Shipping with dry ice.

DESCRIPTION

Background	<p>APT-1/LYPLA1 protein functions as an acyl-protein thioesterase, catalyzing the hydrolysis of fatty acids from S-acylated cysteine residues within proteins, including trimeric G alpha proteins and HRAS. Additionally, it exhibits depalmitoylating activity towards KCNMA1 and potentially ADRB2. Acting as a lysophospholipase, APT-1/LYPLA1 hydrolyzes lysophosphatidylcholine (lyso-PC) and other lysophospholipids such as lyso-PE, lyso-PI, and lyso-PS, with a higher affinity for thioesterase activity. This protein significantly contributes to blood coagulation by recognizing and cleaving plasma</p>
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phospholipids, generating lysophospholipids that serve as substrates for ENPP2, ultimately producing lysophosphatidic acid (LPA).

Caution: Product has not been fully validated for medical applications. For research use only.

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