

## Datasheet

### PSAP monoclonal antibody (M01), clone 1D1-C12

**Catalog Number:** H00005660-M01

**Regulatory Status:** For research use only (RUO)

**Product Description:** Mouse monoclonal antibody raised against a full length recombinant PSAP.

**Clone Name:** 1D1-C12

**Immunogen:** PSAP (AAH01503, 18 a.a. ~ 524 a.a) full-length recombinant protein with GST tag. MW of the GST tag alone is 26 KDa.

**Sequence:**

PVLGLKECTRGSVWCQNVKTASDCGAVKHCLQTVW  
NKPTVKSLPCDICKDVVTAAGDMLKDNATEEEILVYLE  
KTCDWLPKPNMSASCKEIVDSYLPVILDIIKGEMSRPG  
EVCSALNLCESLQKHLAELNHQKQLESNKIPELDMTEV  
VAPFMANIPLLLYPQDGRSKPQPKDNGDVCQDCIQM  
VTDIQTAVRTNSTFVQALVEHVKEECDRLGPGMADICK  
NYISQYSEIAIQMMMHHMQPKEICALVGFCDVEKEMPM  
QTLVPAKVASKNVIPALELVEPIKKHEVPAKSDVYCEV  
CEFLVKEVTKLIDNNKTEKEILDADFDMCSKLPKSLSEE  
CQEVVDTYGSSILSILLEEVSPPELVCMSLHLCSGTRLPA  
LTVHVTQPKDGGFCEVCKKLVGYLDRNLEKNSTKQEIL  
AALEKGCFLPDYQKQCDQFVAEYEPVLIIEILVEVMD  
PSFVCLKIGACPSAHKPLLGTCKIWGPSYWCQNTET  
AAQCNAVEHCKRHVWN

**Host:** Mouse

**Reactivity:** Human

**Applications:** ELISA, IHC-P, IP, S-ELISA, WB-Re,  
WB-Tr

(See our web site product page for detailed applications information)

**Protocols:** See our web site at  
<http://www.abnova.com/support/protocols.asp> or product page for detailed protocols

**Isotype:** IgG2a kappa

**Storage Buffer:** In 1x PBS, pH 7.4

**Storage Instruction:** Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

**Entrez GeneID:** 5660

**Gene Symbol:** PSAP

**Gene Alias:** FLJ00245, GLBA, MGC110993, SAP1

**Gene Summary:** This gene encodes a highly conserved glycoprotein which is a precursor for 4 cleavage products: saposins A, B, C, and D. Each domain of the precursor protein is approximately 80 amino acid residues long with nearly identical placement of cysteine residues and glycosylation sites. Saposins A-D localize primarily to the lysosomal compartment where they facilitate the catabolism of glycosphingolipids with short oligosaccharide groups. The precursor protein exists both as a secretory protein and as an integral membrane protein and has neurotrophic activities. Mutations in this gene have been associated with Gaucher disease, Tay-Sachs disease, and metachromatic leukodystrophy. Alternative splicing results in multiple transcript variants encoding different isoforms. [provided by RefSeq]

**References:**

1. Serum prosaposin levels are increased in patients with advanced prostate cancer. Koochekpour S, Hu S, Vellasco-Gonzalez C, Bernardo R, Azabdaftari G, Zhu G, Zhou HE, Chung LW, Vessella RL. Prostate. 2011 May 31. doi: 10.1002/pros.21427. [Epub ahead of print]