

9F, No. 108, Jhouzih St.,Taipei, Taiwan Tel: + 886-2-8751-1888 Fax: + 886-2-6602-1218 E-mail: sales@abnova.com

Datasheet

AGPAT2 MaxPab mouse polyclonal antibody (B01)

Catalog Number: H00010555-B01

Regulation Status: For research use only (RUO)

Product Description: Mouse polyclonal antibody raised

against a full-length human AGPAT2 protein.

Immunogen: AGPAT2 (NP_006403.2, 1 a.a. ~ 278 a.a)

full-length human protein.

Sequence:

MELWPCLAAALLLLLLLVQLSRAAEFYAKVALYCALCF TVSAVASLVCLLRHGGRTVENMSIIGWFVRSFKYFYGL RFEVRDPRRLQEARPCVIVSNHQSILDMMGLMEVLPE RCVQIAKRELLFLGPVGLIMYLGGVFFINRQRSSTAMT VMADLGERMVRENLKVWIYPEGTRNDNGDLLPFKKG AFYLAVQAQVPIVPVVYSSFSSFYNTKKKFFTSGTVTV QVLEAIPTSGLTAADVPALVDTCHRAMRTTFLHISKTP QENGATAGSGVQPAQ

Host: Mouse

Reactivity: Human

Applications: 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

Storage Buffer: No additive

Storage Instruction: Store at -20°C or lower. Aliquot to

avoid repeated freezing and thawing.

Entrez GenelD: 10555

Gene Symbol: AGPAT2

Gene Alias: 1-AGPAT2, BSCL, BSCL1, LPAAB,

LPAAT-beta

Gene Summary: This gene encodes a member of the 1-acylglycerol-3-phosphate O-acyltransferase family. The protein is located within the endoplasmic reticulum

membrane and converts lysophosphatidic acid to phosphatidic acid, the second step in de novo phospholipid biosynthesis. Mutations in this gene have associated with congenital generalized lipodystrophy (CGL), or Berardinelli-Seip syndrome, a disease characterized by a near absence of adipose and severe insulin resistance. Alternate transcriptional splice variants, encoding different isoforms, have been characterized. [provided by RefSeq]