ANTIBODIES ONLINE

Datasheet for ABIN7317442 UBA1 Protein (GST tag,His tag)



Overview	
Quantity:	50 µg
Target:	UBA1
Origin:	Human
Source:	Baculovirus infected Insect Cells
Protein Type:	Recombinant
Purification tag / Conjugate:	This UBA1 protein is labelled with GST tag,His tag.
Product Details	
Purpose:	Recombinant Human UBE1/UBA1 Protein (His & GST Tag)
Sequence:	Ser 2-Arg 1058
Characteristics:	A DNA sequence encoding the human UBA1 (NP_003325.2) (Ser 2-Arg 1058) was fused with the N-terminal polyhistidine-tagged GST tag at the N-terminus.
Purity:	> 96 % as determined by reducing SDS-PAGE.
Endotoxin Level:	< 1.0 EU per µg as determined by the LAL method.
Target Details	

Target:	UBA1
Alternative Name:	UBE1/UBA1 (UBA1 Products)
Background:	Background: UBE1, also known as UBA1, belongs to the ubiquitin-activating E1 family. UBE1 gene complements an X-linked mouse temperature-sensitive defect in DNA synthesis, and thus may function in DNA repair. It is part of a gene cluster on chromosome Xp11.23. UBE1

Order at www.antibodies-online.com | www.antikoerper-online.de | www.anticorps-enligne.fr | www.antibodies-online.cn International: +49 (0)241 95 163 153 | USA & Canada: +1 877 302 8632 | support@antibodies-online.com Page 1/2 | Product datasheet for ABIN7317442 | 07/25/2024 | Copyright antibodies-online. All rights reserved.

	catalyzes the first step in ubiquitin conjugation to mark cellular proteins for degradation. It also
	catalyzes the first step in ubiquitin conjugation to mark cellular proteins for degradation by first
	adenylating its C-terminal glycine residue with ATP, and thereafter linking this residue to the
	side chain of a cysteine residue in E1, yielding an ubiquitin-E1 thioester and free AMP. Defects
	in UBA1 can cause spinal muscular atrophy X-linked type 2 (SMAX2), also known as X-linked
	lethal infantile spinal muscular atrophy, distal X-linked arthrogryposis multiplex congenita or X-
	linked arthrogryposis type 1 (AMCX1). Spinal muscular atrophy refers to a group of
	neuromuscular disorders characterized by degeneration of the anterior horn cells of the spinal
	cord, leading to symmetrical muscle weakness and atrophy. SMAX2 is a lethal infantile form
	presenting with hypotonia, areflexia, and multiple congenital contractures.
	Synonym: A1S9,A1S9T,A1ST,AMCX1,CFAP124,CTD-
	2522E6.1,GXP1,POC20,SMAX2,UBA1A,UBE1,UBE1X
Molecular Weight:	146 kDa
NCBI Accession:	NP_003325
Application Details	
Restrictions:	For Research Use only
Handling	
Format:	Lyophilized
Reconstitution:	Please refer to the printed manual for detailed information.
Buffer:	Lyophilized from sterile 50 mM Tris, 100 mM NaCl, pH 7.4, 10 % glycerol, 0.5 mM GSH
Storage:	4 °C,-20 °C,-80 °C
Storage Comment:	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.

Order at www.antibodies-online.com | www.antikoerper-online.de | www.anticorps-enligne.fr | www.antibodies-online.cn International: +49 (0)241 95 163 153 | USA & Canada: +1 877 302 8632 | support@antibodies-online.com Page 2/2 | Product datasheet for ABIN7317442 | 07/25/2024 | Copyright antibodies-online. All rights reserved.