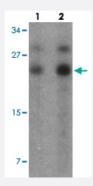


EPM2A polyclonal antibody

Catalog # PAB25630 Size 100 ug

Applications



Western Blot (Cell lysate)

Western blot analysis of EPM2A in SW480 cell lysate with EPM2A polyclonal antibody (Cat # PAB25630) at (1) 1 and (2) 2 ug/mL.

Specification	
Product Description	Rabbit polyclonal antibody raised against synthetic peptide of EPM2A.
Immunogen	A synthetic peptide corresponding to 17 amino acids at C-terminus of human EPM2A.
Host	Rabbit
Theoretical MW (kDa)	36
Reactivity	Human, Mouse, Rat
Specificity	At least four isoforms of EPM2A are known to exist; this antibody will detect all but the shortest isofor m.
Form	Liquid
Purification	Peptide affinity purification
Concentration	1 mg/mL
Isotype	lgG
Recommend Usage	Western Blot (1-2 ug/mL) The optimal working dilution should be determined by the end user.



Product Information

Storage Buffer	In PBS (0.05% sodium azide)
Storage Instruction	Store at 4°C for three months. For long term storage store at -20°C. Aliquot to avoid repeated freezing and thawing.
Note	This product contains sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which shoul d be handled by trained staff only.

Applications

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Enzyme-linked Immunoabsorbent Assay

Gene Info — EPM2A	
Entrez GenelD	<u>7957</u>
Protein Accession#	NP_005661
Gene Name	EPM2A
Gene Alias	EPM2, MELF
Gene Description	epilepsy, progressive myoclonus type 2A, Lafora disease (laforin)
Omim ID	<u>254780</u> <u>607566</u>
Gene Ontology	<u>Hyperlink</u>
Gene Summary	This gene encodes a dual-specificity phosphatase that associates with polyribosomes. The enco ded protein may be involved in the regulation of glycogen metabolism. Mutations in this gene have been associated with myoclonic epilepsy of Lafora. Alternative splicing results in multiple transcript variants. [provided by RefSeq
Other Designations	OTTHUMP00000017360 epilepsy, progressive myoclonus type 2, Lafora disease (laforin) laforin

Disease

Epilepsy



Lafora Disease