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Datasheet for ABIN533790 anti-EPM2A antibody (AA 243-331)

3 Publications



Overview

Quantity:	100 μL
Target:	EPM2A
Binding Specificity:	AA 243-331
Reactivity:	Human
Host:	Mouse
Clonality:	Monoclonal
Conjugate:	This EPM2A antibody is un-conjugated
Application:	Western Blotting (WB), ELISA, Immunofluorescence (IF), Immunocytochemistry (ICC)

Product Details

Purpose:	Mouse monoclonal antibody raised against partial recombinant EPM2A.
Immunogen:	Recombinant protein corresponding to amino acids 243-331 of human EPM2A.
Clone:	K2A3
lsotype:	lgG1
Cross-Reactivity:	Human
Characteristics:	Antibody Reactive Against Recombinant Protein.

Target Details

Target:	EPM2A
Alternative Name:	Laforin (EPM2A Products)

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Target Details

Gene ID:	7957
Pathways:	Cellular Glucan Metabolic Process

Application Details

Application Notes:	The optimal working dilution should be determined by the end user.
Restrictions:	For Research Use only

Handling

Format:	Liquid
Buffer:	In PBS, pH 7.4 (10 % glycerol, 0.02 % sodium azide).
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	-20 °C,-80 °C
Storage Comment:	Store at 2°C to 8°C for 1 week. For long term storage, aliquot and store at -20°C to -80°C. Aliquot to avoid repeated freezing and thawing.
Publications	
Product cited in:	Tagliabracci, Turnbull, Wang, Girard, Zhao, Skurat, Delgado-Escueta, Minassian, Depaoli-Roach,
	Roach: "Laforin is a glycogen phosphatase, deficiency of which leads to elevated
	phosphorylation of glycogen in vivo." in: Proceedings of the National Academy of Sciences of
	the United States of America, Vol. 104, Issue 49, pp. 19262-6, (2007) (PubMed).
	Wang, Parker, Skurat, Raben, DePaoli-Roach, Roach: "Relationship between glycogen
	accumulation and the laforin dual specificity phosphatase." in: Biochemical and biophysical
	research communications, Vol. 350, Issue 3, pp. 588-92, (2006) (PubMed).
	Ganesh, Delgado-Escueta, Suzuki, Francheschetti, Riggio, Avanzini, Rabinowicz, Bohlega, Bailey
	Alonso, Rasmussen, Thomson, Ochoa, Prado, Medina, Yamakawa: "Genotype-phenotype
	correlations for EPM2A mutations in Lafora's progressive myoclonus epilepsy: exon 1
	mutations associate with an early-onset cognitive deficit subphenotype." in: Human molecular
	genetics, Vol. 11, Issue 11, pp. 1263-71, (2002) (PubMed).

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Images

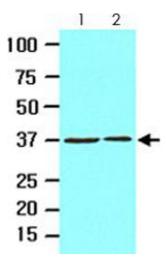


Image 1. Cell lysates of HeLa (lane 1) and 293T (lane 2) (20 ug) were resolved by SDS-PAGE and probed with EPM2A monoclonal antibody, clone k2A3 (1:1000). Proteins were visualized using a goat anti-mouse secondary antibody conjugated to HRP and an ECL detection system.

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