



Anti-FKTN (aa 29-138) polyclonal antibody (DPAB-DC970)

This product is for research use only and is not intended for diagnostic use.

PRODUCT INFORMATION

Antigen Description	The protein encoded by this gene is a putative transmembrane protein that is localized to the cis-Golgi compartment, where it may be involved in the glycosylation of alpha-dystroglycan in skeletal muscle. The encoded protein is thought to be a glycosyltransferase and could play a role in brain development. Defects in this gene are a cause of Fukuyama-type congenital muscular dystrophy (FCMD), Walker-Warburg syndrome (WWS), limb-girdle muscular dystrophy type 2M (LGMD2M), and dilated cardiomyopathy type 1X (CMD1X). Alternatively spliced transcript variants have been found for this gene.
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Immunogen	FCMD (NP_006722, 29 a.a. ~ 138 a.a) partial recombinant protein with GST tag. The sequence is KHYLSTKNGAGLSKSKGSRIGFDSTQWRAVKKFIMLTSNQNVFVLIDPLILELINKNFE QVKNTSHGSTSQCKFFCVPRDFTAFALQYHLWKNEEGWFRIAENMGFQCL
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Source/Host	Mouse
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Species Reactivity	Human
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Conjugate	Unconjugated
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Applications	WB (Tissue lysate), WB (Recombinant protein), ELISA,
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Size	50 µl
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Buffer	50 % glycerol
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Preservative	None
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Storage	Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.
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GENE INFORMATION

Gene Name	FKTN fukutin [Homo sapiens (human)]
Official Symbol	FKTN
Synonyms	FKTN; fukutin; FCMD; CMD1X; LGMD2M; MDDGA4; MDDGB4; MDDGC4; patient fukutin; Fukuyama type congenital muscular dystrophy protein;
Entrez Gene ID	2218
Protein Refseq	NP_001073270
UniProt ID	O75072
Chromosome Location	9q31-q33
Function	transferase activity;