



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

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| 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. |
| Immunogen | FCMD (NP_006722, 29 a.a. ~ 138 a.a) partial recombinant protein with GST tag. The sequence is KHYLSTKNGAGLSKSKGSRIGFDSTQWRAVKKFIMLTSNQNVPVFLIDPLILELINKNF QVKNTSHGSTSQCKFFCVPRDFTAFLQYHLWKNEEGWFRIAENMGFQCL |
| Source/Host | Mouse |
| Species Reactivity | Human |
| Conjugate | Unconjugated |
| Applications | WB (Tissue lysate), WB (Recombinant protein), ELISA, |
| Size | 50 µl |
| Buffer | 50 % glycerol |
| Preservative | None |
| Storage | Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing. |

GENE INFORMATION

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| 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; |
