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SHFM1 Rabbit pAb

Catalog No.: A18152

Basic Information

Observed MW
Refer to figures

Calculated MW
8kDa

Category
Primary antibody

Applications
ELISA,WB

Cross-Reactivity
Human

Background

The product of this gene has been localized within the split hand/split foot malformation locus SHFM1 at chromosome 7. It has been proposed to be a candidate gene for the autosomal dominant form of the heterogeneous limb developmental disorder split hand/split foot malformation type 1. In addition, it has been shown to directly interact with BRCA2. It also may play a role in the completion of the cell cycle.

Recommended Dilutions

WB	1:500 - 1:2000

Immunogen Information

Gene ID
7979

Swiss Prot
P60896

Immunogen
A synthetic peptide corresponding to a sequence within amino acids 1-70 of human SHFM1 (NP_006295.1).

Synonyms
ECD; DSS1; SHFD1; SHFM1; SHSF1; PSMD15; Shfdg1; C7orf76

Contact

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Product Information

Source
Rabbit

Isotype
IgG

Purification
Affinity purification

Storage
Store at -20℃. Avoid freeze / thaw cycles.
Buffer: PBS with 0.01% thimerosal,50% glycerol,pH7.3.