SHFM3 (D-16): sc-163369



The Power to Question

BACKGROUND

SHFM3 (split-hand/foot malformation type 3), also known as FBXW4 (F-box and WD-40 domain-containing protein 4) or dactylin, is a 412 amino acid protein that is primarily involved in signaling pathways that are essential for normal limb development. Specifically, SHFM3 promotes ubiquination and degradation of certain phosphorylated proteins. Defects in the gene encoding SHFM3 are the cause of split-hand/foot malformation type 3, an autosomal dominant disorder that is characterized by hypoplasia/aplasia of the central digits, causing variable fusion with the remaining digits. The disease is not a result of a point mutation, but rather a genomic rearrangement resulting in a tandem duplication that contains a disrupted copy of the gene encoding SHFM3, as well as extra copies of two other genes that are linked to limb development. SHFM3 is typically expressed in kidney, brain, lung and liver, with highest expression in fetal brain.

REFERENCES

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- Sidow, A., et al. 1999. A novel member of the F-box/WD40 gene family, encoding dactylin, is disrupted in the mouse dactylaplasia mutant. Nat. Genet. 23: 104-107.
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- 4. Basel, D., et al. 2003. Split hand foot malformation is associated with a reduced level of dactylin gene expression. Clin. Genet. 64: 350-354.
- de Mollerat, X.J., et al. 2003. A genomic rearrangement resulting in a tandem duplication is associated with split hand-split foot malformation 3 (SHFM3) at 10q24. Hum. Mol. Genet. 12: 1959-1971.
- 6. Kano, H., et al. 2005. Genomic rearrangement at 10q24 in non-syndromic split-hand/split-foot malformation. Hum. Genet. 118: 477-483.

CHROMOSOMAL LOCATION

Genetic locus: FBXW4 (human) mapping to 10q24.32; Fbxw4 (mouse) mapping to 19 C3.

SOURCE

SHFM3 (D-16) is an affinity purified goat polyclonal antibody raised against a peptide mapping near the C-terminus of SHFM3 of human origin.

PRODUCT

Each vial contains 200 μg lgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-163369 P, (100 μ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

RESEARCH USE

For research use only, not for use in diagnostic procedures.

APPLICATIONS

SHFM3 (D-16) is recommended for detection of SHFM3 of mouse, rat and human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

SHFM3 (D-16) is also recommended for detection of SHFM3 in additional species, including equine, canine, bovine and porcine.

Suitable for use as control antibody for SHFM3 siRNA (h): sc-90449, SHFM3 siRNA (m): sc-153450, SHFM3 shRNA Plasmid (h): sc-90449-SH, SHFM3 shRNA Plasmid (m): sc-153450-SH, SHFM3 shRNA (h) Lentiviral Particles: sc-90449-V and SHFM3 shRNA (m) Lentiviral Particles: sc-153450-V.

Molecular Weight of SHFM3: 46 kDa.

RECOMMENDED SECONDARY REAGENTS

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use donkey anti-goat IgG-HRP: sc-2020 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible donkey anti-goat IgG-HRP: sc-2033 (dilution range: 1:2000-1:5000), Cruz Marker™ Molecular Weight Standards: sc-2035, TBS Blotto A Blocking Reagent: sc-2333 and Western Blotting Luminol Reagent: sc-2048. 2) Immunofluorescence: use donkey anti-goat IgG-FITC: sc-2024 (dilution range: 1:100-1:400) or donkey anti-goat IgG-TR: sc-2783 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941.

STORAGE

Store at 4° C, **DO NOT FREEZE**. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

PROTOCOLS

See our web site at www.scbt.com or our catalog for detailed protocols and support products.

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