# MRVI1 (M-17): sc-10962



The Power to Question

#### **BACKGROUND**

The integration of the murine leukemia virus (MuLV) into the mammalian genome is frequently associated with insertional mutagenesis of cellular proto-oncogenes and tumor suppressor genes leading to cellular transformation and leukemias. Several proto-oncogenes were initially identified as sites of viral integration, including the tumor-suppressors Myc, Myb, and Hox. A related murine virus, MRV, also induces leukemia through viral integration and the disruption of the MRVI1 encoding gene. MRVI1 is specifically expressed in megakaryocytes and various myeloid leukemias, and its expression is downregulated during monocytic differentiation. The human and murine homologs of MRVI1 share substantial sequence similarity and similar expression patterns and are most closely related to the lymphoid specific protein Jaw1. The transcripts generated from MRVI1 are alternatively spliced and initiated from two distinct promoters to produce a longer isoform, MRVI1a, which contains an N-terminal 84 amino acid extension that is not present in the otherwise identical, shorter isoform, MRVI1b. These two isoforms have distinct subcellular localization patterns as MRVI1a contains an additional transmembrane domain and localizes to the endoplasmic recticulum, while MRVI1b is diffusely distributed throughout the cell.

## **REFERENCES**

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- Buchberg, A.M., et al. 1990. Evi-2, a common integration site involved in murine myeloid leukemogenesis. Mol. Cell. Biol. 10: 4658-4666.
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#### CHROMOSOMAL LOCATION

Genetic locus: Mrvi1 (mouse) mapping to 7 F1.

# SOURCE

MRVI1 (M-17) is an affinity purified goat polyclonal antibody raised against a peptide mapping near the C-terminus of MRVI1 of mouse origin.

### **RESEARCH USE**

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

#### **PRODUCT**

Each vial contains 200  $\mu g$  IgG in 1.0 ml of PBS with <0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-10962 P, (100  $\mu$ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

#### **APPLICATIONS**

MRVI1 (M-17) is recommended for detection of MRVI1 of mouse 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).

MRVI1 (M-17) is also recommended for detection of MRVI1 in additional species, including equine and canine.

Suitable for use as control antibody for MRVI1 siRNA (m): sc-42929, MRVI1 shRNA Plasmid (m): sc-42929-SH and MRVI1 shRNA (m) Lentiviral Particles: sc-42929-V.

#### **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) 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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