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# WBSCR22

**Reactivity:**Human Rat

**Tested applications:**WB IHC IF

**Recommended Dilution:**WB 1:500 - 1:2000 IHC 1:50 - 1:200 IF 1:50 - 1:100

**Calculated MW:**32kDa

**Observed MW:**Refer to figures

**Immunogen:**

Recombinant protein of human WBSCR22

**Storage Buffer:**

Store at -20. Avoid freeze / thaw cycles. Buffer: PBS with 0.02% sodium azide, 50% glycerol, pH7.3.

**Synonym:**

WBMT; MERM1; PP3381; HUSSY-3; HASJ4442;

**Catalog #:**A7317

**Antibody Type:**

Polyclonal Antibody

**Species:**Rabbit

**Gene ID:**114049

**Isotype:**IgG

**Swiss Prot:**O43709

**Purity:**Affinity purification

For research use only.

**Background:**

This gene encodes a protein containing a nuclear localization signal and an S-adenosyl-L-methionine binding motif typical of methyltransferases, suggesting that the encoded protein may act on DNA methylation. This gene is deleted in Williams syndrome, a multisystem developmental disorder caused by the deletion of contiguous genes at 7q11.23. Alternatively spliced transcript variants have been found.

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