

**MESP2 rabbit pAb****Cat#: orb771926 (Manual)**

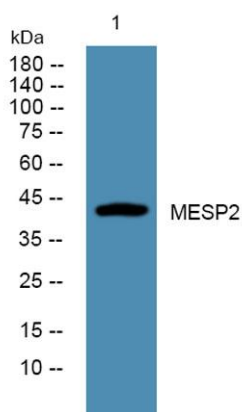
For research use only. Not intended for diagnostic use.

<b>Product Name</b>	MESP2 rabbit pAb
<b>Host species</b>	Rabbit
<b>Applications</b>	WB;ELISA
<b>Species Cross-Reactivity</b>	Human;Rat;Mouse;
<b>Recommended dilutions</b>	WB 1:500-2000 ELISA 1:5000-20000
<b>Immunogen</b>	Synthesized peptide derived from human protein . at AA range: 220-300
<b>Specificity</b>	MESP2 Polyclonal Antibody detects endogenous levels of protein.
<b>Formulation</b>	Liquid in PBS containing 50% glycerol, and 0.02% sodium azide..
<b>Storage</b>	Store at -20°C. Avoid repeated freeze-thaw cycles.
<b>Protein Name</b>	Mesoderm posterior protein 2 (Class C basic helix-loop-helix protein 6) (bHLHc6)
<b>Gene Name</b>	MESP2 BHLHC6 SCDO2
<b>Cellular localization</b>	Nucleus .
<b>Purification</b>	The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.
<b>Clonality</b>	Polyclonal

<b>Concentration</b>	1 mg/ml
<b>Observed band</b>	43kD
<b>Human Gene ID</b>	145873
<b>Human Swiss-Prot Number</b>	Q0VG99
<b>Alternative Names</b>	

### Background

This gene encodes a member of the bHLH family of transcription factors and plays a key role in defining the rostrocaudal patterning of somites via interactions with multiple Notch signaling pathways. This gene is expressed in the anterior presomitic mesoderm and is downregulated immediately after the formation of segmented somites. This gene also plays a role in the formation of epithelial somitic mesoderm and cardiac mesoderm. Mutations in the MESP2 gene cause autosomal recessive spondylocostal dysostosis 2 (SCD02). [provided by RefSeq, Oct 2008],



**Western blot analysis of lysates from A431 cells, primary antibody was diluted at 1:1000, 4° over night**