



MESP2 Polyclonal Antibody

ImmunogenSynthesized peptide derived from human protein . at AA range: 220-300SpecificityMESP2 Polyclonal Antibody detects endogenous levels of protein.FormulationLiquid in PBS containing 50% glycerol, and 0.02% sodium azide.SourcePolyclonal, Rabbit,IgGPurificationThe antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.DilutionWB 1:500-2000 ELISA 1:5000-20000Concentration1 mg/mlPurity≥90%Storage Stability-20°C/1 yearSynonymsCollege and the severity associated with vertebraeCell PathwayNucleus .Tissue Specificitydisease:Defects in MESP2 are the cause of spondylocostal dysostosis autoson recessive type 2 (SCDO2) (IMI of variability. Automations include fusion of vertebrae, fusion of certain ribs, and other rib malformations. Deformity of the chest and spine (severe scolosis, kyphoscoliosis and lordsis) a ratural consequence of the malformations include fusion of vertebrae, fusion of certain ribs, and other rib malformations. Deformity of the chest and spine (severe rescilosis, kyphoscoliosis and lordsis) a ratural consequence of the malformation and leads to a dwarf-like appearanc As the thorax is small, infants frequently have respiratory insufficiency and repeated respiratory infections resulting in life-threatening complications in the first year of life, disease:Defects in MESP2 may be a cause of spondylocotal dysostosis and lordsis) a ratural consequence of the malformation and leads to a dwarf-like appearanc As the thorax is small, infants frequently have respiratory instriction of setting complications in the first year of life, disease:Defects in MESP2 may be a cause of spon		
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Nanjing BYabscience technology Co.,Ltd

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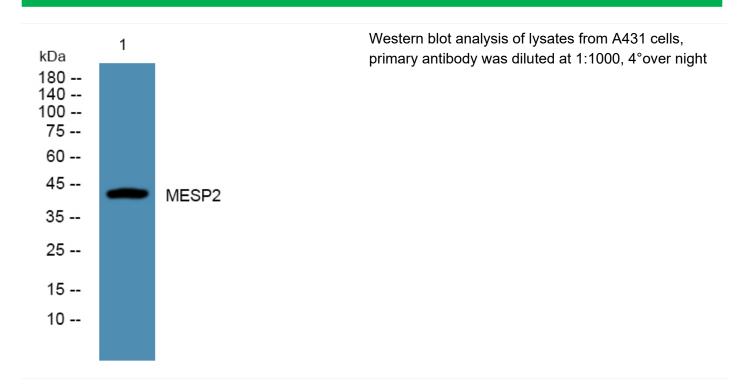


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Background	This gene encodes a member of the bHLH family of transcription factors and plays a key role in defining the rostrocaudal patterning of somites via interactions with multiple Notch signaling pathways. This gene is expressed in the anterior 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 dystosis 2 (SCD02). [provided by RefSeq, Oct 2008],
matters needing attention	Avoid repeated freezing and thawing!
Usage suggestions	This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.

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