



SHOX2 mouse mAb

| Catalog No | BYmab-10999 | | | |
|---|--|--|--|--|
| lsotype | lgG | | | |
| Reactivity | Human; Mouse;Rat | | | |
| Applications | WB | | | |
| Gene Name | SHOX2 OG12X SHOT | | | |
| Protein Name | SHOX2 | | | |
| Immunogen | Synthesized peptide derived from human SHOX2 AA range: 37-87 | | | |
| Specificity | This antibody detects endogenous levels of SHOX2 at Human/Mouse/Rat | | | |
| FormulationLiquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium | | | | |
| Source | Monoclonal, Mouse,IgG | | | |
| PurificationThe antibody was affinity-purified from mouse antiserum by affinity-chromatography using epitope-specific immunogen. | | | | |
| Dilution WB 1:500-2000 | | | | |
| Concentration | 1 mg/ml | | | |
| Purity | ≥90% | | | |
| Storage Stability -20°C/1 year | | | | |
| Synonyms | | | | |
| Observed Band | | | | |
| Cell Pathway | Nucleus. | | | |
| Tissue Specificity Expressed in heart, skeletal muscle, liver, lung, bone marrow fibro and placenta. | | | | |
| Function | developmental stage:Expressed during cranofacial development as well as in heart.,function:May be a growth regulator and have a role in specifying neural systems involved in processing somatosensory information, as well as in face and body structure formation.,similarity:Belongs to the paired homeobox family. Bicoid subfamily.,similarity:Contains 1 homeobox DNA-binding domain.,similarity:Contains 1 OAR domain.,tissue specificity:Expressed in heart, skeletal muscle, liver, lung, bone marrow fibroblast, pancreas and placenta., | | | |
| Background | This gene is a member of the homeobox family of genes that encode proteins containing a 60-amino acid residue motif that represents a DNA binding domain. Homeobox genes have been characterized extensively as transcriptional regulators involved in pattern formation in both invertebrate and vertebrate | | | |
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| | | homeobox of is thought to short stature be a candid | veral human genetic disorders are caused by aberrations in human genes. This locus represents a pseudoautosomal homeobox gene that b be responsible for idiopathic short stature, and it is implicated in the e phenotype of Turner syndrome patients. This gene is considered to ate gene for Cornelia de Lange syndrome. Alternative splicing results ranscript variants. [provided by RefSeq, Jul 2009], | | |
|--|---|--|--|--|--|
| matters needing attention | | Avoid repea | ted freezing and thawing! | | |
| Usage suggestions | | | t can be used in immunological reaction related experiments. For ation, please consult technical personnel. | | |
| Products Images | | | | | |
| kDa 180 140 100 75 60 45 35 | 1 | SHOX2 | Western Blot analysis of various cells using SHOX2 mouse mAb | | |
| 25 15 | | | | | |

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