

Ihh Rabbit mAb

Catalog # AP76546

Product Information

| | |
|--------------------------|------------------------|
| Application | WB, IHC-P |
| Primary Accession | Q14623 |
| Reactivity | Rat, Human, Mouse |
| Host | Rabbit |
| Clonality | Monoclonal Antibody |
| Isotype | IgG |
| Conjugate | Unconjugated |
| Purification | Affinity Purified |
| Calculated MW | 45251 |

Additional Information

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|--------------------|---|
| Gene ID | 3549 |
| Other Names | IHH |
| Dilution | WB~~1:1000 IHC-P~~N/A |
| Format | Liquid in 50mM Tris-Glycine(pH 7.4), 0.15M NaCl, 40%Glycerol, 0.01% sodium azide and 0.05% BSA. |
| Storage | Store at 4°C short term. Aliquot and store at -20°C long term. Avoid freeze/thaw cycles. |

Protein Information

| | |
|--------------------------|---|
| Name | IHH (HGNC:5956) |
| Function | Plays a role in embryonic morphogenesis; it is involved in the regulation of endochondral skeleton formation, and the development of retinal pigment epithelium (RPE), photoreceptors and periocular tissues (By similarity). |
| Cellular Location | [Indian hedgehog protein N-product]: Cell membrane; Lipid-anchor {ECO:0000250 UniProtKB:Q62226}. Note=The N-product remains associated with the cell surface. {ECO:0000250 UniProtKB:Q15465} |
| Tissue Location | Expressed in embryonic lung, and in adult kidney and liver |

Background

This gene encodes a member of the hedgehog family of secreted signaling molecules. Hedgehog proteins

are essential regulators of a variety of developmental processes including growth, patterning and morphogenesis. The encoded protein specifically plays a role in bone growth and differentiation. Mutations in this gene are the cause of brachydactyly type A1 which is characterized by shortening or malformation of the phalanges. Mutations in this gene are also the cause of acrocapitofemoral dysplasia.

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