

MNX1/HB9/HLXB9 Rabbit pAb

货号: **AYP22898**

产品信息

| | |
|-------|--|
| 反应 | Human,Mouse,Rat |
| 宿主 | Rabbit |
| 克隆性 | Polyclonal |
| 预测反应 | |
| 应用 | WB |
| 推荐浓度 | WB: 1:500 - 1:1000 |
| 理论分子量 | 41kDa |
| 实测分子量 | 55KDa |
| 形式 | Liquid |
| 保存条件 | Store at -20°C. Avoid freeze / thaw cycles. Buffer: PBS with 0.05% proclin300,50% glycerol,pH7.3. |
| 偶联物 | Unconjugated |
| 阳性对照 | HeLa,Mouse thymus,Rat liver |
| 细胞定位 | cytosol,nucleolus,nucleoplasm,nucleus |
| 纯化 | Affinity purification |

抗原信息

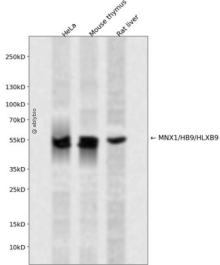
| | |
|------|---|
| 抗原信息 | Recombinant fusion protein containing a sequence corresponding to amino acids 201-315 of human MNX 1/HB9/HLXB9 (NP_005506.3). |
| 序列 | IKLGAGTFQLDQWLRASTAGMILPKMPDFNSQAQSNLLGKCRPRPTAFTSQLLLELHQFKLNKYLSRPKRFEVATSLMLT ETQVKIWFQNRMRKWKRSKKAKEQAAQAEKQKG |

靶点信息

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| 研究背景 | This gene encodes a nuclear protein, which contains a homeobox domain and is a transcription factor. Mutations in this gene result in Currarino syndrome, an autosomic dominant congenital malformation. Alternately spliced transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, Sep 2009] |
| 基因ID | 3110 |

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|-------|---------------------------|
| 基因名 | MNX1 |
| Swiss | P50219 |
| 别名 | HB9; HLXB9; HOXHB9; SCRA1 |

产品验证



Western blot analysis of MNX1/HLXB9 expressed in HeLa, Mouse thymus, Rat liver using MNX1/HLXB9 Rabbit pAb at 1:1000. Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) at 1:5000. Lysates/proteins: 30ug per lane. Blocking buffer: 5% non-fat dry milk in TBST. Detection: ECL Enhanced Kit. Exposure time: 120s.

实验步骤

访问官网浏览详情: www.ablybio.cn