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## MNX1/HB9/HLXB9 Rabbit mAb

Catalog No.: A19690 Recombinant 1 Publications

## **Basic Information**

#### **Observed MW**

46kDa

#### **Calculated MW**

41kDa

## **Category**

SMab Recombinant Monoclonal Antibody

## **Applications**

WB, ELISA

## **Cross-Reactivity**

Human, Mouse

#### CloneNo number

ARC2233

## Background

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. Alternatively spliced transcript variants encoding different isoforms have been found for this gene.

## **Recommended Dilutions**

WB 1:1000 - 1:5000

**ELISA** 

Recommended starting concentration is 1 µg/mL. Please optimize the concentration based on your specific assay requirements.

## **Immunogen Information**

**Gene ID**3110

Swiss Prot
P50219

### **Immunogen**

A synthetic peptide corresponding to a sequence within amino acids 250-350 of human MNX1/HB9/HLXB9 (P50219).

#### **Synonyms**

HB9; HLXB9; SCRA1; HOXHB9; MNX1/HB9/HLXB9

## Contact

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## **Product Information**

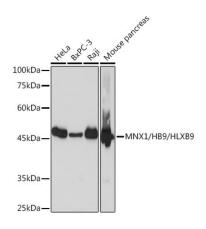
SourceIsotypePurificationRabbitIgGAffinity purification

#### Storage

Store at -20°C. Avoid freeze / thaw cycles.

Buffer: PBS with 0.02% sodium azide, 0.05% BSA, 50% glycerol, pH7.3.

## Validation Data



Western blot analysis of various lysates using MNX1/MNX1/HB9/HLXB9 Rabbit mAb (A19690) at 1:1000 dilution.

Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

Lysates/proteins: 25µg per lane.

Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

Exposure time: 1s.