

## CAB19690

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### Product Information

<b>Product SKU:</b>	CAB19690	<b>Gene ID:</b>	3110	<b>Size:</b>	20uL, 100uL
<b>Clone No:</b>	ARC2233	<b>Host Species:</b>	Rabbit	<b>Reactivity:</b>	Human,Mouse

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### Additional Information

<b>Observed MW:</b>	41kDa	<b>Conjugate:</b>	Unconjugated
<b>Calculated MW:</b>	41kDa	<b>Isotype:</b>	IgG

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### Immunogen Information

**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 Dilution:** WB,1:1000 - 1:5000

**Synonyms:** HB9; HLXB9; SCRA1; HOXHB9; MNX1/HB9/HLXB9

**Purification Method:** Affinity purification

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

**Storage:** Store at -20°C. Avoid freeze / thaw cycles.Buffer: PBS with 0.02% sodium azide,0.05% BSA,50% glycerol,pH7.3.