

## CAB21992

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### Product Information

<b>Product SKU:</b> CAB21992	<b>Gene ID:</b> 1482	<b>Size:</b> 20uL, 100uL
<b>Clone No:</b> -	<b>Host Species:</b> Rabbit	<b>Reactivity:</b> Mouse,Rat

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### Additional Information

<b>Observed MW:</b> 30-42kDa	<b>Conjugate:</b> Unconjugated
<b>Calculated MW:</b> 35kDa	<b>Isotype:</b> IgG

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### Immunogen Information

**Background:** This gene encodes a homeobox-containing transcription factor. This transcription factor functions in heart formation and development. Mutations in this gene cause atrial septal defect with atrioventricular conduction defect, and also tetralogy of Fallot, which are both heart malformation diseases. Mutations in this gene can also cause congenital hypothyroidism non-goitrous type 5, a non-autoimmune condition. Alternative splicing results in multiple transcript variants.

**Recommended Dilution:** WB,1:100 - 1:500

**Synonyms:** CSX; CSX1; VSD3; CHNG5; HLHS2; NKX2E; NKX2.5; NKX4-1; NKX2-5

**Purification Method:** Affinity purification

**Immunogen:** Recombinant fusion protein containing a sequence corresponding to amino acids 24-137 of human NKX2-5 (NP\_004378.1).

**Storage:** Store at -20°C. Avoid freeze / thaw cycles.Buffer: PBS with 0.05% proclin300,50% glycerol,pH7.3.