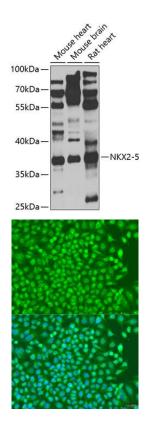
NKX2-5 Rabbit Polyclonal Antibody

CAB5651



roduct Information Size:	Protein Background This gene encodes a homeobox-containing transcription factor. This transcription facto
20uL, 50uL, 100uL, 200uL	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 hear malformation diseases. Mutations in this gene can also cause congenital hypothyroidism non goitrous type 5, a non-autoimmune condition. Alternative splicing results in multiple transcrip
Observed MW:	
37kDa	variants.
Calculated MW:	Immunogen information
11kDa/16kDa/34kDa	Gene ID:
Applications:	1482
WB IF	Uniprot
Reactivity:	P52952
Human, Mouse, Rat	Synonyms: NKX2-5; CHNG5; CSX; CSX1; HLHS2; NKX2.5; NKX2E; NKX4-1; VSD3
Antibody Information	
Recommended dilutions: WB 1:500 - 1:2000 IF 1:50 - 1:200	Immunogen: Recombinant fusion protein containing a sequence corresponding to amino acids 1-135 of human NKX2-5 (NP_004378.1).
Source: Rabbit	
	Storage: Store at -20°C. Avoid freeze / thaw cycles. Buffer: PBS with 0.02%
lsotype: lgG	sodium azide, 50% glycerol, pH7.3.

Purification: Affinity purification



Western blot analysis of extracts of various cell lines, using NKX2-5 antibody (CAB5651) at 1:1000 dilution._Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) (CABS014) at 1:10000 dilution._Lysates/proteins: 25ug per lane._Blocking buffer: 3% nonfat dry milk in TBST._Detection: ECL Enhanced Kit (CABM00021)._Exposure time: 60s.

Immunofluorescence analysis of U2OS cells using NKX2-5 antibody (CAB5651) at dilution of 1:100. Blue: DAPI for nuclear staining.