

Anti-cTnI (Phospho-S22/S23) Antibody

Catalog #	Source	Reactivity	Applications
CPA4534	Rabbit	H, M, R, P, Rb	WB, IH
Description	Rabbit polyclonal antibody to cTnI (Phospho-S22/S23)		
Immunogen	KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding S22/S23 of human cTnI protein. The exact sequence is proprietary.		
Purification	The antibody was purified by immunogen affinity chromatography.		
Specificity	Recognizes endogenous levels of cTnI protein only when phosphorylated at S22/S23.		
Clonality	Polyclonal		
Conjugation			
Form	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, and 0.01% sodium azide.		
Dilution	WB (1/500 - 1/1000), IH (1/100 - 1/200)		
Gene Symbol	TNNI3		
Alternative Names	TNNI3; TNNC1; Troponin I, cardiac muscle; Cardiac troponin I		
Entrez Gene	7137 (Human); 21954 (Mouse); 29248 (Rat)		
SwissProt	P19429 (Human); P48787 (Mouse); P23693 (Rat)		
Storage/Stability	Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid freeze/thaw cycles.		

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, CHIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference

Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb- Rabbit, S- Sheep, Z- Zebrafish

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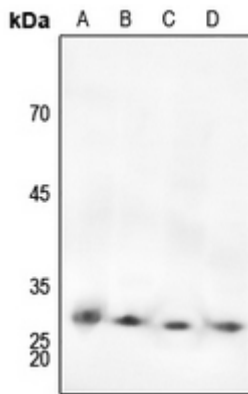
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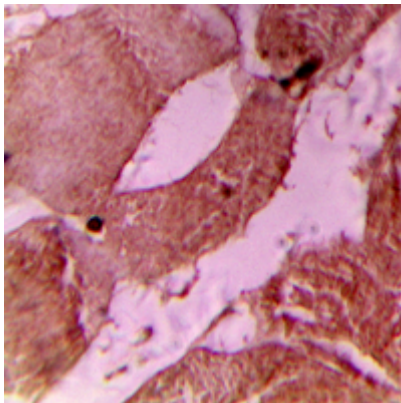
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Product Data Sheet



Western blot analysis of cTnI (Phospho-S22/S23) expression in mouse heart (A), mouse kidney (B), rat heart (C), rat kidney (D) whole cell lysates. (Predicted band size: 24 kD; Observed band size: 28 kD)



Immunohistochemical analysis of cTnI (Phospho-S22/S23) staining in human heart formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

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