

Anti-cTnI (Phospho-S43) Antibody

| Catalog # | Source | Reactivity | Applications |
|--------------------------|---|------------------|--------------|
| CPA4850 | Rabbit | H, M, R, B, D, P | WB, IH |
| Description | Rabbit polyclonal antibody to cTnI (Phospho-S43) | | |
| Immunogen | KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding S43 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 S43. | | |
| 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 | 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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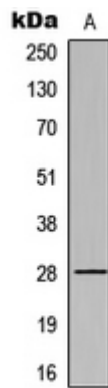
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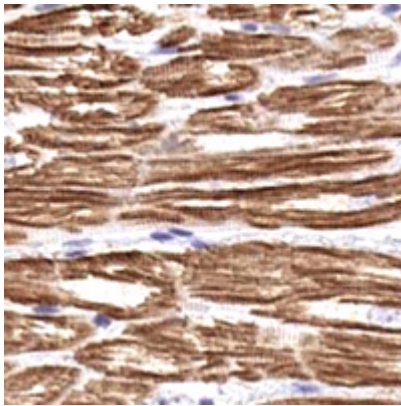
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Product Data Sheet



Western blot analysis of cTnI (Phospho-S43) expression in human heart (A) whole cell lysates. (Predicted band size: 24 kD; Observed band size: 28 kD)



Immunohistochemical analysis of cTnI (Phospho-S43) 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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