

Product Data Sheet

Anti-Actin Sarcomeric Muscle Antibody

Catalog # Source Reactivity Applications

CPA9787 Mouse H IH

Description Mouse monoclonal antibody to Actin Sarcomeric Muscle

Immunogen KLH-conjugated synthetic peptide encompassing a sequence within human Actin

Sarcomeric Muscle. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of Actin Sarcomeric Muscle protein.

Clonality Monoclonal

Conjugation

Form Mouse IgG1. Liquid in PBS containing 50% glycerol, 0.2% BSA and 0.01% sodium

azide.

Dilution IH (1/100 - 1/300)

Gene Symbol ACTA1; ACTC1

Alternative Names ACTA; Actin alpha skeletal muscle; Alpha-actin-1; ACTC; Actin alpha cardiac muscle

1; Alpha-cardiac actin

Entrez Gene 58, 70 (Human)

SwissProt P68133, P68032 (Human)

Storage/Stability Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid

freeze/thaw cycles.

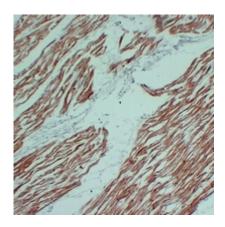
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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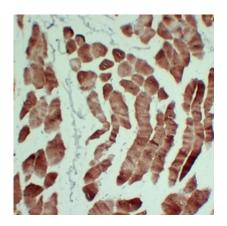
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Immunohistochemical analysis of Actin Sarcomeric Muscle staining in human myocardium formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was th



Immunohistochemical analysis of Actin Sarcomeric Muscle staining in human skeletal muscle 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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