

Product Data Sheet

Anti-FOXO1/3 Antibody

Catalog # Source Reactivity Applications

CPA5311 Rabbit H, M, R, B, P WB, IH

Description Rabbit polyclonal antibody to FOXO1/3

Immunogen KLH-conjugated synthetic peptide encompassing a sequence within the center

region of human FOXO1/3. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of FOXO1/3 protein.

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 - 1/2000), IH (1/50 - 1/200)

Gene Symbol FOXO1; FOXO3

Alternative Names FOXO1; FKHR; FOXO1A; Forkhead box protein O1; Forkhead box protein O1A;

Forkhead in rhabdomyosarcoma; FOXO3; FKHRL1; FOXO3A; Forkhead box protein

O3; AF6g21 protein; Forkhead in rhabdomyosarcoma-like 1

Entrez Gene 2308 (Human); 56458, 56484 (Mouse); 84482 (Rat)

SwissProt Q12778, O43524 (Human); Q9R1E0, Q9WVH4 (Mouse); G3V7R4 (Rat)

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

freeze/thaw cycles.

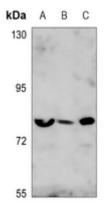
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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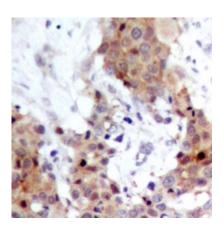
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Western blot analysis of FOXO1/3 expression in mouse heart (A), mouse muscle (B), rat heart (C) whole cell lysates. (Predicted band size: 69; 71 kD; Observed band size: 78 kD)



Immunohistochemical analysis of FOXO1/3 staining in human breast cancer 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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