

Product Data Sheet

Anti-AMACR Antibody

Catalog # Source Reactivity Applications

CQA1817 Rabbit H, M, R WB, IH, IF/IC

Description Rabbit polyclonal antibody to AMACR

Immunogen Recombinant full length protein of human AMACR

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of AMACR 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), IF/IC (1/50 - 1/200)

Gene Symbol AMACR

Alternative Names Alpha-methylacyl-CoA racemase; 2-methylacyl-CoA racemase

Entrez Gene 23600 (Human); 17117 (Mouse)

SwissProt Q9UHK6 (Human); O09174 (Mouse); P70473 (Rat)

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

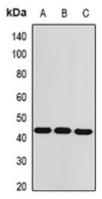
freeze/thaw cycles.

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

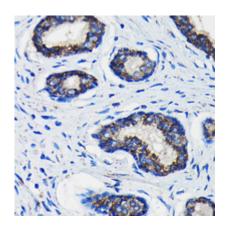
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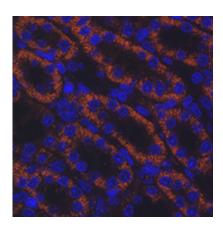
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Western blot analysis of AMACR expression in SW480 (A), Hela (B), mouse liver (C) whole cell lysates. (Predicted band size: 22; 25; 42; 43 kD; Observed band size: 42 kD)



Immunohistochemical analysis of AMACR staining in human prostate 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.



Immunofluorescent analysis of AMACR staining in rat kidney. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a humidified chamber. Cells were washed with PBST and incubated with a AF594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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