

Product Data Sheet

Anti-HCCR-1 Antibody

Catalog # Source Reactivity Applications

CQA1121 Rabbit H, M, R WB, IH, IF/IC

Description Rabbit polyclonal antibody to HCCR-1

Immunogen Recombinant full length protein of human HCCR-1

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of HCCR-1 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 LETMD1

Alternative Names LETM1 domain-containing protein 1; Cervical cancer 1 proto-oncogene protein p40;

Cervical cancer proto-oncogene 2 protein; HCCR-1; HCRR-2

Entrez Gene 25875 (Human); 68614 (Mouse)

SwissProt Q6P1Q0 (Human); Q924L1 (Mouse)

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

freeze/thaw cycles.

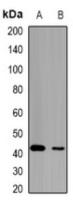
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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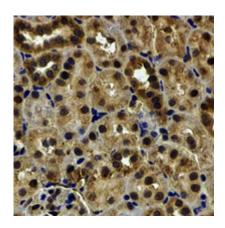
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Western blot analysis of HCCR-1 expression in mouse kidney (A), mouse liver (B) whole cell lysates. (Predicted band size: 11; 17; 23; 35; 41; 43 kD; Observed band size: 41 kD)



Immunohistochemical analysis of HCCR-1 staining in human kidney 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 HCCR-1 staining in MCF7 cells. 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 hidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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