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Product Data Sheet

Anti-GlnRS Antibody

Catalog #	Source	Reactivity	Applications
CQA1502	Rabbit	H, M	WB, IH, IF/IC

Description	Rabbit polyclonal antibody to GlnRS
Immunogen	Recombinant full length protein of human GlnRS
Purification	The antibody was purified by immunogen affinity chromatography.
Specificity	Recognizes endogenous levels of GlnRS 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 - 1/100)
Gene Symbol	QARS
Alternative Names	Glutamine--tRNA ligase; Glutaminyl-tRNA synthetase; GlnRS
Entrez Gene	5859 (Human)
SwissProt	P47897 (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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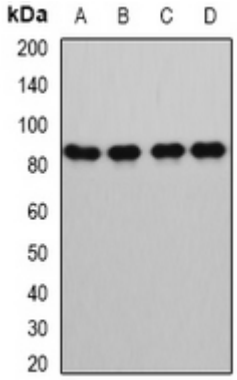
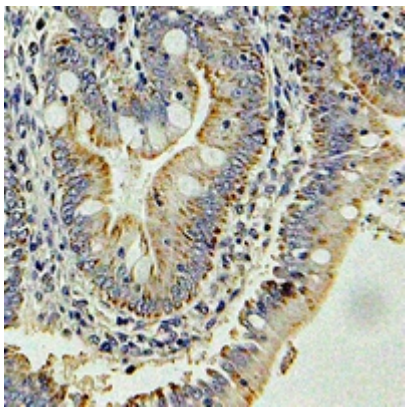
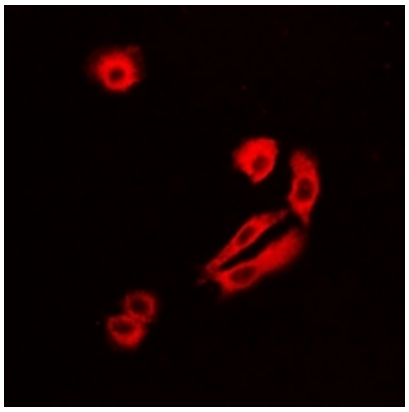


Western blot analysis of GlnRS expression in HL60 (A), Hela (B), mouse spleen (C), mouse kidney (D) whole cell lysates. (Predicted band size: 86; 87 kD; Observed band size: 88 kD)



Immunohistochemical analysis of GlnRS staining in human colon 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 GlnRS staining in U2OS 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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