

Product Data Sheet

Anti-EPHX2 Antibody

| Catalog # | Source | Reactivity | Applications | | |
|-------------------|----------|---|--|--|--|
| CQA1088 | Rabbit | H, M, R | WB, IH | | |
| Description | Ra | bbit polyclonal antibody | to EPHX2 | | |
| Immunogen | Re | combinant full length pr | otein of human EPHX2 | | |
| Purification | | The antibody was purified by immunogen affinity chromatography. | | | |
| Specificity | | Recognizes endogenous levels of EPHX2 protein. | | | |
| Clonality | Ро | lyclonal | | | |
| Conjugation | | | | | |
| Form | Liq | Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, | | | |
| | an | d 0.01% sodium azide. | | | |
| Dilution | W | 3 (1/500 - 1/1000), IH (1/5 | 60 - 1/100) | | |
| Gene Symbol | EP | HX2 | | | |
| Alternative Names | | Bifunctional epoxide hydrolase 2 | | | |
| Entrez Gene | 20 | 53 (Human); 13850 (Mo | use); 65030 (Rat) | | |
| SwissProt | P3- | 4913 (Human); P34914 | (Mouse); P80299 (Rat) | | |
| Storage/Stabi | lity Shi | pped at 4°C. Upon deliv | ery aliquot and store at -20°C for one year. Avoid | | |
| | fre | eze/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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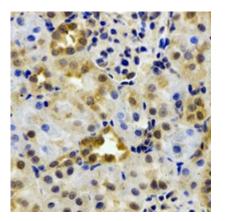
kDa A B

140

For research purposes only, not for human use

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Western blot analysis of EPHX2 expression in MCF7 (A), Jurkat (B) whole cell lysates. (Predicted band size: 55; 57; 62 kD; Observed band size: 63 kD)



Immunohistochemical analysis of EPHX2 staining in rat 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.

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