

Product Data Sheet

Anti-MERTK/TYRO3 Antibody

Catalog # Source Reactivity Applications

CPA5918 Rabbit H, M, R WB, IH

Description Rabbit polyclonal antibody to MERTK/TYRO3

Immunogen KLH-conjugated synthetic peptide encompassing a sequence within the C-term

region of human MERTK/TYRO3. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of MERTK/TYRO3 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 - 1/1000), IH (1/50 - 1/200)

Gene Symbol MERTK; TYRO3

Alternative Names MERTK; MER; Tyrosine-protein kinase Mer; Proto-oncogene c-Mer; Receptor

tyrosine kinase MerTK; TYRO3; BYK; DTK; RSE; SKY; Tyrosine-protein kinase receptor

TYRO3; Tyrosine-protein kinase DTK; Tyrosine-protein kinase RSE; Tyrosine-protein

kinase SKY; Tyrosine-protein kinase byk

Entrez Gene 10461, 7301 (Human); 17289, 22174 (Mouse); 65037, 25232 (Rat)

SwissProt Q12866, Q06418 (Human); Q60805, P55144 (Mouse); P57097, P55146 (Rat)

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

freeze/thaw cycles.

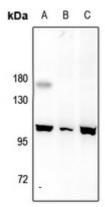
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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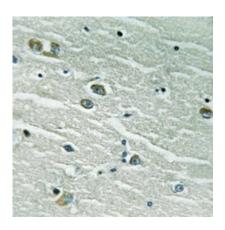
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Western blot analysis of MERTK/TYRO3 expression in A375 (A), mouse brain (B), rat brain (C) whole cell lysates. (Predicted band size: 110; 96 kD; Observed band size: 110 kD)



Immunohistochemical analysis of MERTK/TYRO3 staining in human brain 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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