

Product Data Sheet

Anti-Tyrosine Hydroxylase (pS71) Antibody

Catalog # Source Reactivity Applications

CPA3337 Rabbit H, M, R, C, D WB, IH

Description Rabbit polyclonal antibody to Tyrosine Hydroxylase (pS71)

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

region of human Tyrosine Hydroxylase (pS71). The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of Tyrosine Hydroxylase (pS71) protein.

Clonality Polyclonal

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 (1/100 - 1/200)

Gene Symbol TH

Alternative Names TYH; Tyrosine 3-monooxygenase; Tyrosine 3-hydroxylase; TH

Entrez Gene 7054 (Human); 21823 (Mouse); 25085 (Rat)

SwissProt P07101 (Human); P24529 (Mouse); P04177 (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

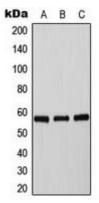
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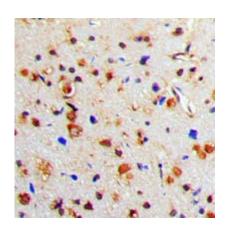




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Western blot analysis of Tyrosine Hydroxylase (pS71) expression in HepG2 UV-treated (A), mouse brain (B), PC12 (C) whole cell lysates.



Immunohistochemical analysis of Tyrosine Hydroxylase (pS71) 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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