

Product Data Sheet

Anti-Tyrosine Hydroxylase Antibody

| Catalog # | Source | Reactivity | Applications |
|---------------|-----------|----------------------------|--|
| CPA3338 | Rabbit | H, M, R, B, D | WB, IH, IF/IC, IP |
| Description | Rab | bit polyclonal antibody t | o Tyrosine Hydroxylase |
| Immunogen | KLH | -conjugated synthetic pe | eptide encompassing a sequence within the N-term |
| | regi | on of human Tyrosine H | ydroxylase. The exact sequence is proprietary. |
| Purification | The | antibody was purified b | y immunogen affinity chromatography. |
| Specificity | Rec | ognizes endogenous leve | els of Tyrosine Hydroxylase protein. |
| Clonality | Poly | yclonal | |
| Form | Liqu | uid in 0.42% Potassium p | hosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, |
| | and | 0.01% sodium azide. | |
| Dilution | WB | (1/500 - 1/1000), IH (1/10 | 00 - 1/200), IF/IC (1/100 - 1/500), IP (1/10 - 1/100) |
| Gene Symbol | ТН | | |
| Alternative N | ames TYH | ; Tyrosine 3-monooxyge | nase; Tyrosine 3-hydroxylase; TH |
| Entrez Gene | 705 | 4 (Human); 21823 (Mou | se); 25085 (Rat) |
| SwissProt | P07 | 101 (Human); P24529 (N | Nouse); P04177 (Rat) |
| Storage/Stabi | lity Ship | oped at 4°C. Upon delive | ry aliquot and store at -20°C for one year. Avoid |
| | free | ze/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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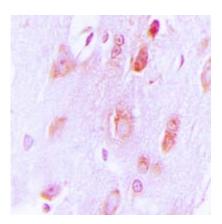


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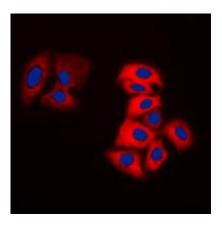
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Western blot analysis of Tyrosine Hydroxylase expression in DLD (A), mouse liver (B), PC12 (C) whole cell lysates.



Immunohistochemical analysis of Tyrosine Hydroxylase 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.



Immunofluorescent analysis of Tyrosine Hydroxylase staining in DLD 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 humidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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