

## **Product Data Sheet**

### Anti-Tyrosine Hydroxylase Antibody

Catalog #	Source	Reactivity	Applications		
CPA3339	Rabbit	H, M, R, C, D	WB, IH, IF/IC		
Description	Rab	Rabbit polyclonal antibody to Tyrosine Hydroxylase			
Immunogen	KLH	-conjugated synthetic per	tide encompassing a sequence within the N-term		
	regi	on of human Tyrosine Hyd	lroxylase. The exact sequence is proprietary.		
Purification	The	antibody was purified by	immunogen affinity chromatography.		
Specificity	Rec	ognizes endogenous level	s of Tyrosine Hydroxylase protein.		
Clonality	Poly	vclonal			
Conjugation					
Form	Liqu	iid in 0.42% Potassium ph	osphate, 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), IF/IC (1/100 - 1/500)		
Gene Symbol	TH				
Alternative Na	ames TYH	; Tyrosine 3-monooxygen	ase; Tyrosine 3-hydroxylase; TH		
Entrez Gene	705	4 (Human); 21823 (Mouse	e); 25085 (Rat)		
SwissProt	P07	101 (Human); P24529 (M	ouse); P04177 (Rat)		
Storage/Stabi	lity Ship	oped at 4°C. Upon delivery	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

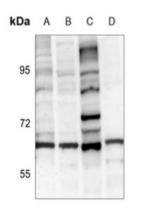
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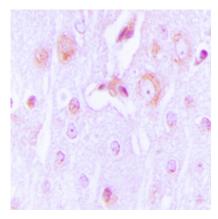


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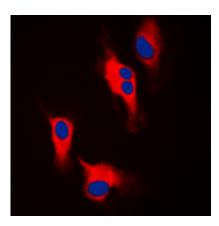
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Western blot analysis of Tyrosine Hydroxylase expression in HEK293T (A), Hela (B), SP20 (C), PC12 (D) whole cell lysates. (Predicted band size: 58 kD; Observed band size: 58 kD)



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 A549 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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