

### **Product Data Sheet**

## Anti-Cytochrome P450 17A1 Antibody

Catalog #	Source	e Reactivity	Applications		
CPA1326	Rabbit	H, M, R	WB, IH, IF/IC		
Description		Rabbit polyclonal antibody to Cytochrome P450 17A1			
Immunogen		KLH-conjugated synthetic peptide encompassing a sequence within the center			
	ne P450 17A1. The exact sequence is proprietary.				
Purification		The antibody was purified by immunogen affinity chromatography.			
Specificity		Recognizes endogenous levels of Cytochrome P450 17A1 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 (1/5	0 - 1/100), IF/IC (1/50 - 1/200)		
Gene Symbol		CYP17A1			
Alternative Names		CYP17; S17AH; Steroid 17-alpha-hydroxylase/17,20 lyase;			
		17-alpha-hydroxyprogester	one aldolase; CYPXVII; Cytochrome P450 17A1;		
		Cytochrome P450-C17; Cyt	ochrome P450c17; Steroid 17-alpha-monooxygenase		
Entrez Gene		1586 (Human); 13074 (Mouse); 25146 (Rat)			
SwissProt		P05093 (Human); P27786 (Mouse); P11715 (Rat)			
Storage/Stabi	bility 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

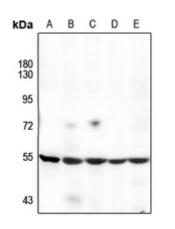
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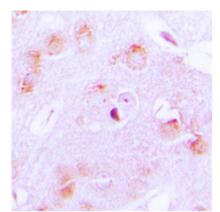


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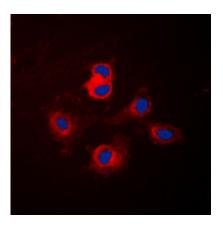
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Western blot analysis of Cytochrome P450 17A1 expression in PC3 (A), U87MG (B), A2780 (C), PC12 (D), AML12 (E) whole cell lysates. (Predicted band size: 57 kD; Observed band size: 55 kD)



Immunohistochemical analysis of Cytochrome P450 17A1 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 Cytochrome P450 17A1 staining in NIH3T3 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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