

Product Data Sheet

Anti-Cytochrome P450 2D6 Antibody

Catalog #	Source	Reactivity	Applications		
-		-			
CPA1316	Rabbit	H, M, R, Mk	WB, IH		
Description	Ra	abbit polyclonal antibody to	Cytochrome P450 2D6		
Immunogen	KL	H-conjugated synthetic pe	otide encompassing a sequence within the C-term		
	re	gion of human Cytochrome	P450 2D6. The exact sequence is proprietary.		
Purification	Th	ne antibody was purified by	immunogen affinity chromatography.		
Specificity	Re	ecognizes endogenous leve	s of Cytochrome P450 2D6 protein.		
Clonality Polyclonal					
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	an	nd 0.01% sodium azide.			
Dilution	W	/B (1/500 - 1/1000), IH (1/50	- 1/100)		
Gene Symbol	CY	(P2D6			
Alternative Na	ames CY	(P2DL1; Cytochrome P450	2D6; CYPIID6; Cytochrome P450-DB1; Debrisoquine		
	4-	hydroxylase			
Entrez Gene	15	565 (Human)			
SwissProt	P1	10635 (Human)			
Storage/Stabi	lity Sh	nipped at 4°C. Upon deliver	/ aliquot and store at -20°C for one year. Avoid		
	fre	eeze/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

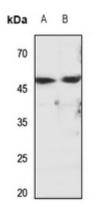
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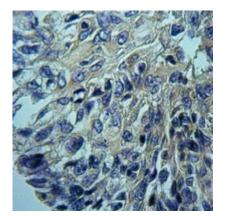


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Western blot analysis of Cytochrome P450 2D6 expression in mouse liver (A), rat liver (B) whole cell lysates. (Predicted band size: 55 kD; Observed band size: 55 kD)



Immunohistochemical analysis of Cytochrome P450 2D6 staining in rat heart 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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