

Product Data Sheet

Anti-Cytochrome P450 2D6 Antibody

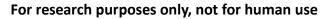
Catalog #	Source	Reactivity	Applications
CQA8184	Rabbit	н	WB, IH, IF/IC
Description	R	abbit monoclonal antibo	dy to Cytochrome P450 2D6
Immunogen	R	Recombinant protein of h	uman CYP2D6
Purification	Т	he antibody was purified	by immunogen affinity chromatography.
Specificity	R	ecognizes endogenous le	vels of Cytochrome P450 2D6 protein.
Clonality	N	Monoclonal	
Conjugation			
Form	L	iquid in 50mM Tris-Glycir	ne (pH 7.4), 0.15M NaCl, 50% Glycerol, 0.01% Sodium azide
	а	ind 0.05% BSA.	
Dilution	V	VB (1/500 - 1/1000), IH (1/	50 - 1/100), IF/IC (1/50 - 1/100)
Gene Symbol	C	CYP2D6	
Alternative Na	ames C	YP2DL1; Cytochrome P4	50 2D6; CYPIID6; Cytochrome P450-DB1; Debrisoquine
	4	l-hydroxylase	
Entrez Gene	1	.565 (Human)	
SwissProt	Р	210635 (Human)	
Storage/Stabi	lity S	hipped at 4°C. Upon deliv	very aliquot and store at -20°C for one year. Avoid
	fı	reeze/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

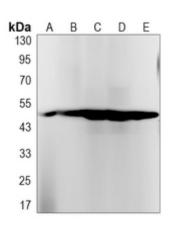
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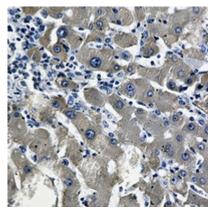




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Western blot analysis of Cytochrome P450 2D6 expression in Hela (A), A549 (B), HL60 (C), U251 (D), U87MG (E) whole cell lysates. (Predicted band size: 56 kD; Observed band size: 50 kD)



Immunohistochemical analysis of Cytochrome P450 2D6 staining in human liver cancer 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 2D6 staining in HeLa 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 hidified chamber. Cells were washed with PBST and incubated with a AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark.

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