

## **Product Data Sheet**

## Anti-Cytochrome P450 2B6 Antibody

Catalog #	Source	Reactivity	Applications	
CPA1313	Rabbit	-	WB, IH	
CFAISIS	Nabbit			
Description		Rabbit polyclonal antibody t	o Cytochrome P450 2B6	
Immunogen		KLH-conjugated synthetic pe	ptide encompassing a sequence within the center	
		region of human Cytochrom	e P450 2B6. The exact sequence is proprietary.	
Purification		The antibody was purified by	<i>immunogen affinity chromatography.</i>	
Specificity		Recognizes endogenous leve	ls of Cytochrome P450 2B6 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 (1/50	- 1/200)	
Gene Symbol		CYP2B6		
Alternative Na	ames	Cytochrome P450 2B6; 1,4-c	ineole 2-exo-monooxygenase; CYPIIB6; Cytochrome	
		P450 IIB1		
Entrez Gene		1555 (Human)		
SwissProt		P20813 (Human)		
Storage/Stabi	lity	Shipped at 4°C. Upon deliver	y 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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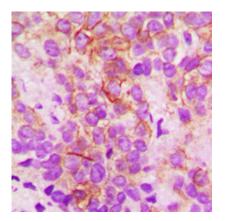
kDa A

250

For research purposes only, not for human use

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Western blot analysis of Cytochrome P450 2B6 expression in HepG2 (A) whole cell lysates. (Predicted band size: 33; 56 kD; Observed band size: 66 kD)



Immunohistochemical analysis of Cytochrome P450 2B6 staining in human breast 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.

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