

Product Data Sheet

Anti-Cytochrome P450 1A1 Antibody

Catalog #	Source	React	ivity	Applications		
CQA1125	Rabbit	Н, М		WB, IH		
Description	R	Rabbit polyclonal antibody to Cytochrome P450 1A1				
Immunogen	R	Recombinant f	full length protein of hum	an Cytochrome P450 1A1		
Purification	т	The antibody was purified by immunogen affinity chromatography.				
Specificity	R	Recognizes en	dogenous levels of Cytoch	hrome P450 1A1 protein.		
Clonality	Р	Polyclonal				
Conjugation						
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,				
	а	and 0.01% sod	lium azide.			
Dilution	V	NB (1/500 - 1/2	2000), IH (1/50 - 1/200)			
Gene Symbol	C	CYP1A1				
Alternative Na	ames C	Cytochrome P	450 1A1; CYPIA1; Cytochr	rome P450 form 6; Cytochrome P450-C;		
	С	Cytochrome P	450-P1			
Entrez Gene	1	L543 (Human)	; 13076 (Mouse)			
SwissProt	Р	P04798 (Human); P00184 (Mouse)				
Storage/Stabi	lity S	Shipped at 4 $^\circ$	C. Upon delivery aliquot	and store at -20 $^\circ$ C for one year. Avoid		
	fı	reeze/thaw cy	/cles.			

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

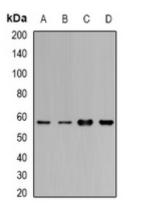
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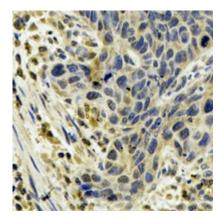


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Western blot analysis of Cytochrome P450 1A1 expression in MCF7 (A), BT474 (B), mouse lung (C), mouse liver (D) whole cell lysates. (Predicted band size: 58 kD; Observed band size: 55 kD)



Immunohistochemical analysis of Cytochrome P450 1A1 staining in human esophageal 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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