

## Anti-Cytochrome P450 Reductase Antibody (A16086)

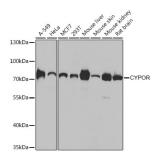
## Specifications:

Name:	Anti-Cytochrome P450 Reductase Antibody
Description:	Rabbit polyclonal antibody to Cytochrome P450 Reductase.
Applications:	WB, IHC, ICC/IF
Recommended Dilutions:	WB: 1:500-1:2,000, IHC: 1:50-1:100, ICC/IF: 1:50-1:200
Reactivity:	Human, Mouse, Rat
Immunogen:	Recombinant fusion protein containing a sequence corresponding to amino acids 1-250 of human CYPOR (NP_000932.3).
Sequence:	MINMGDSHVDTSSTVSEAVAEEVSLFSMTDMILFSLIVGLLTYWFLFRKKKEEVPEFT KIQTLTSSVRESSFVEKMKKTGRNIIVFYGSQTGTAEEFANRLSKDAHRYGMRGMSAD PEEYDLADLSSLPEIDNALVVFCMATYGEGDPTDNAQDFYDWLQETDVDLSGVKFAVF GLGNKTYEHFNAMGKYVDKRLEQLGAQRIFELGLGDDDGNLEEDFITWREQFWPAVCE HFGVEATGEESSIRQYEL
Host:	Rabbit
Clonality:	Polyclonal
Isotype:	lgG
Conjugate:	Unconjugated
Purification:	Affinity purification.
Molecular Weight:	77 kDa
Molecular Weight: Product Form:	77 kDa Liquid
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Product Form:	Liquid Supplied in Phosphate Buffered Saline, pH 7.3, with 50% Glycerol and 0.02% Sodium

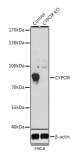
## antibodies

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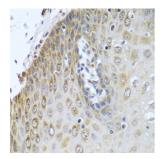
Images:



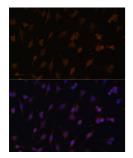
Western blot analysis of extracts of various cell lines, using Anti-Cytochrome P450 Reductase Antibody (A16086) at 1:1,000 dilution. The secondary antibody was Goat Anti-Rabbit IgG H&L Antibody (HRP) at 1:10,000 dilution. Lysates/proteins were present at 25µg per lane. The blocking buffer used was 3% non-fat dry milk in TBST. Detection was with a ECL Basic Kit. Exposure time: 5s.



Western blot analysis of extracts from normal (control) and CYPOR knockout (KO) HeLa cells, using Anti-Cytochrome P450 Reductase Antibody (A16086) at 1:1,000 dilution. The secondary antibody was Goat Anti-Rabbit IgG H&L Antibody (HRP) at 1:10,000 dilution. Lysates/proteins were present at 25µg per lane. The blocking buffer used was 3% non-fat dry milk in TBST. Detection was with a ECL Basic Kit. Exposure time: 1s.



Immunohistochemistry analysis of paraffin-embedded human esophagus using Anti-Cytochrome P450 Reductase Antibody (A16086) at a dilution of 1:100 (40x lens). Perform microwave antigen retrieval with 10 mM PBS buffer pH 7.2 before commencing with IHC staining protocol.



Immunofluorescence analysis of C6 cells using Anti-Cytochrome P450 Reductase Antibody (A16086) at a dilution of 1:100 (40x lens). DAPI was used to stain the cell nuclei (blue).