

Product Data Sheet

Anti-E2F4 Antibody

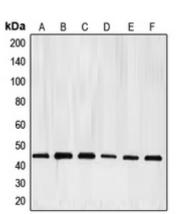
Catalog #	Source	Reactivity	Applications		
CPA1360	Rabbit	H, M, R, C	WB, IH		
Description	ļ	Rabbit polyclonal antibody to E2F4			
Immunogen	ļ	KLH-conjugated synthetic pe	otide encompassing a sequence within the center		
	I	region of human E2F4. The exact sequence is proprietary.			
Purification	-	The antibody was purified by	immunogen affinity chromatography.		
Specificity	I	Recognizes endogenous leve	s of E2F4 protein.		
Clonality	l	Polyclonal			
Conjugation					
Form	l	Liquid in 0.42% Potassium pł	osphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	i	and 0.01% sodium azide.			
Dilution	,	WB (1/500 - 1/1000), IH (1/10) - 1/200)		
Gene Symbol	I	E2F4; E2F5			
Alternative Names E2F4; Transcription factor E2F4; E2F-4; E2F5;		E2F4; Transcription factor E2	F4; E2F-4; E2F5; Transcription factor E2F5; E2F-5		
Entrez Gene	:	1874, 1875 (Human); 104394	, 13559 (Mouse)		
SwissProt		Q16254, Q15329 (Human); C	8R0K9, Q61502 (Mouse); Q62814 (Rat)		
Storage/Stabi	lity	Shipped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid		
	t	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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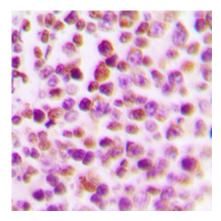




For research purposes only, not for human use

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Western blot analysis of E2F4 expression in HeLa (A), K562 (B), Jurkat (C), Raji (D), A431 (E), mouse kidney (F) whole cell lysates. (Predicted band size: 43; 37 kD; Observed band size: 44 kD)



Immunohistochemical analysis of E2F4 staining in human tonsil 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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