

Product Data Sheet

Anti-NR2F6 Antibody

Catalog #	Source	Reactivity	Applications
CPA1395	Rabbit	H, M, R, B, D, P	WB, IH, IF/IC
Description	Ra	abbit polyclonal antibody to	NR2F6
Immunogen	K	LH-conjugated synthetic pept	ide encompassing a sequence within the N-term
	re	egion of human NR2F6. The e	xact sequence is proprietary.
Purification	TI	he antibody was purified by i	mmunogen affinity chromatography.
Specificity	R	ecognizes endogenous levels	of NR2F6 protein.
Clonality	Po	olyclonal	
Conjugation			
Form	Li	iquid in 0.42% Potassium phc	sphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	aı	nd 0.01% sodium azide.	
Dilution	W	/B (1/500 - 1/1000), IH (1/50 -	1/100), IF/IC (1/50 - 1/200)
Gene Symbol	Ν	R2F6	
Alternative Na	ames E	AR2; ERBAL2; Nuclear recept	or subfamily 2 group F member 6; V-erbA-related
	р	rotein 2; EAR-2	
Entrez Gene	20	063 (Human); 13864 (Mouse	; 245980 (Rat)
SwissProt	P	10588 (Human); P43136 (Mo	use); O09017 (Rat)
Storage/Stabi	lity Sł	hipped at 4°C. Upon delivery	aliquot and store at -20°C for one year. Avoid
	fr	eeze/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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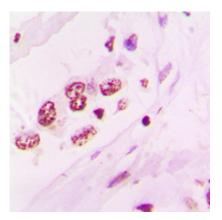
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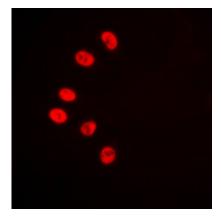
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Western blot analysis of NR2F6 expression in HEK293T (A) whole cell lysates. (Predicted band size: 42 kD; Observed band size: 42 kD)



Immunohistochemical analysis of NR2F6 staining in human lung 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 NR2F6 staining in HepG2 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 humidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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