

Product Data Sheet

Anti-UBE2F Antibody

Catalog #	Source	Reactivity	Applications
CQA3058	Rabbit	H, M, R	WB, IH, IF/IC
Description	Ra	abbit polyclonal antibody t	o UBE2F
Immunogen	Re	ecombinant full length pro	ein of human UBE2F
Purification	Tł	he antibody was purified b	y immunogen affinity chromatography.
Specificity	Re	ecognizes endogenous leve	ls of UBE2F protein.
Clonality	Po	olyclonal	
Conjugation			
Form	Lie	quid in 0.42% Potassium p	nosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	ar	nd 0.01% sodium azide.	
Dilution	W	/B (1/500 - 1/2000), IH (1/50	- 1/200), IF/IC (1/50 - 1/200)
Gene Symbol	U	BE2F	
Alternative N	ames N	CE2; NEDD8-conjugating e	nzyme UBE2F; NEDD8 carrier protein UBE2F; NEDD8
	pr	rotein ligase UBE2F; NEDD	8-conjugating enzyme 2; Ubiquitin-conjugating enzyme
	E2	2 F	
Entrez Gene	14	40739 (Human); 363284 (R	at)
SwissProt	Q	969M7 (Human); Q5U203	(Rat)
Storage/Stabi	ility Sł	hipped at 4°C. Upon delive	y 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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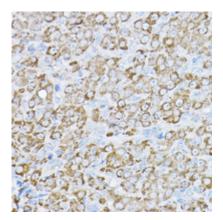


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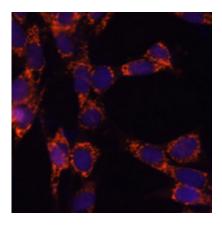
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Western blot analysis of UBE2F expression in K562 (A), mouse liver (B), mouse heart (C) whole cell lysates. (Predicted band size: 11; 13; 17; 18; 21; 23 kD; Observed band size: 21 kD)



Immunohistochemical analysis of UBE2F staining in rat ovary 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 UBE2F staining in NIH3T3 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 AF594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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