

Product Data Sheet

Anti-STIP1 Antibody

Catalog #	Source	Reactivity	Applications
CQA2258	Rabbit	H, M, R	WB, IH, IF/IC, IP
Description	Ra	bbit polyclonal antibody	to STIP1
Immunogen	Re	combinant full length pr	otein of human STIP1
Purification	The	e antibody was purified	oy immunogen affinity chromatography.
Specificity	Re	cognizes endogenous lev	rels of STIP1 protein.
Clonality	Po	lyclonal	
Conjugation			
Form	Liq	uid in 0.42% Potassium	phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	an	d 0.01% sodium azide.	
Dilution	WE	3 (1/500 - 1/2000), IH (1/5	0 - 1/200), IF/IC (1/50 - 1/200), IP (1/20 - 1/50)
Gene Symbol	STI	IP1	
Alternative Na	ames Str	ess-induced-phosphopro	otein 1; STI1; Hsc70/Hsp90-organizing protein; Hop; Renal
	car	cinoma antigen NY-REN	11; Transformation-sensitive protein IEF SSP 3521
Entrez Gene	10	963 (Human); 20867 (M	ouse); 192277 (Rat)
SwissProt	P3	1948 (Human); Q60864	(Mouse); O35814 (Rat)
Storage/Stabi	lity Shi	ipped at 4°C. Upon deliv	ery aliquot and store at -20°C for one year. Avoid
	fre	eze/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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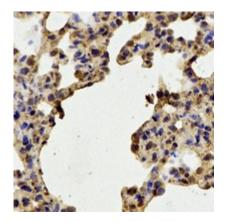
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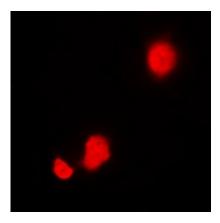


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Western blot analysis of STIP1 expression in Hela (A), COS7 (B), mouse brain (C) whole cell lysates. (Predicted band size: 59; 62; 68 kD; Observed band size: 63 kD)



Immunohistochemical analysis of STIP1 staining in rat lung 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 STIP1 staining in Hela 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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