

# **Product Data Sheet**

# **Anti-SFPQ Antibody**

Catalog #	Source	Reactivity	Applications	
CQA2361	Rabbit	H, M, R	WB, IH, IF/IC	
Description	F	Rabbit polyclonal antibody t	o SFPQ	
Immunogen	F	Recombinant full length pro	tein of human SFPQ	
Purification	Т	The antibody was purified b	y immunogen affinity chromatography.	
Specificity	F	Recognizes endogenous leve	els of SFPQ protein.	
Clonality	F	Polyclonal		
Conjugation				
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	a	and 0.01% sodium azide.		
Dilution	V	NB (1/500 - 1/2000), IH (1/50	- 1/200), IF/IC (1/50 - 1/200)	
Gene Symbol	S	SFPQ		
Alternative N	ames P	SF; Splicing factor, proline-	and glutamine-rich; 100 kDa DNA-pairing protein;	
	ł	POMp100; DNA-binding p	52/p100 complex, 100 kDa subunit; Polypyrimidine	
	t	ract-binding protein-associa	ated-splicing factor; PSF; PTB-associated-splicing factor	
Entrez Gene	6	5421 (Human); 71514 (Mou	se)	
SwissProt	F	23246 (Human); Q8VIJ6 (N	louse)	
Storage/Stabi	l <b>ity</b> S	Shipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid	
	f	reeze/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

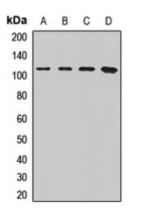
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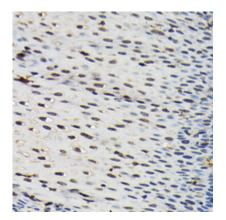


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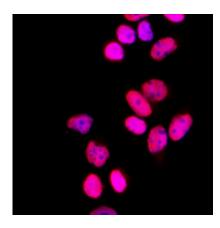
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Western blot analysis of SFPQ expression in Jurkat (A), HepG2 (B), mouse liver (C), rat brain (D) whole cell lysates. (Predicted band size: 72; 76 kD; Observed band size: 115 kD)



Immunohistochemical analysis of SFPQ staining in human esophageal 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 SFPQ 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 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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