

# **Product Data Sheet**

## **Anti-TAP2** Antibody

Catalog #	Source	e Reactivity	Applications	
CQA1062	Rabbit	H, M, R	WB, IH, IF/IC	
Description		Rabbit polyclonal antibody	o TAP2	
Immunogen		Recombinant full length pro	tein of human TAP2	
Purification		The antibody was purified b	y immunogen affinity chromatography.	
Specificity		Recognizes endogenous leve	els of TAP2 protein.	
Clonality		Polyclonal		
Conjugation				
Form		Liquid in 0.42% Potassium p	hosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,	
		and 0.01% sodium azide.		
Dilution		WB (1/500 - 1/2000), IH (1/50	- 1/200), IF/IC (1/50 - 1/200)	
Gene Symbol		TAP2		
Alternative Names		ABCB3; PSF2; RING11; Y1; Antigen peptide transporter 2; APT2; ATP-binding cassette		
		sub-family B member 3; Per	tide supply factor 2; Peptide transporter PSF2; PSF-2;	
		Peptide transporter TAP2; P	eptide transporter involved in antigen processing 2;	
		Really interesting new gene	11 protein	
Entrez Gene		6891 (Human); 21355 (Mou	se); 24812 (Rat)	
SwissProt		Q03519 (Human); P36371 (I	Mouse); P36372 (Rat)	
Storage/Stabi	lity	Shipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid	
		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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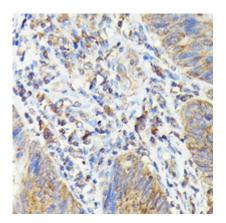
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Western blot analysis of TAP2 expression in MCF7 (A), mouse spleen (B), mouse lung (C) whole cell lysates. (Predicted band size: 72; 75 kD; Observed band size: 75 kD)



Immunohistochemical analysis of TAP2 staining in human colon 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 TAP2 staining in U2OS 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 hidified 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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