

## **Product Data Sheet**

## Anti-Serpin A10 Antibody

Catalog #	Source	Reactivity	Applications
CQA2225	Rabbit	H <i>,</i> R	WB, IH, IF/IC
Description	Ra	abbit polyclonal antibody t	o Serpin A10
Immunogen	Re	ecombinant full length pro	tein of human Serpin A10
Purification	Tł	he antibody was purified b	y immunogen affinity chromatography.
Specificity	Re	ecognizes endogenous leve	els of Serpin A10 protein.
Clonality	Ро	olyclonal	
Conjugation			
Form	Li	iquid in 0.42% Potassium p	hosphate, 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 - 1/100)
Gene Symbol	SE	ERPINA10	
Alternative Na	ames ZF	PI; Protein Z-dependent pr	otease inhibitor; PZ-dependent protease inhibitor; PZI;
	Se	erpin A10	
Entrez Gene	52	1156 (Human); 100909524	, 171154 (Rat)
SwissProt	Q	9UK55 (Human); Q62975 (	Rat)
Storage/Stabi	<b>lity</b> Sł	hipped at 4 $^\circ$ C. Upon deliv	ery aliquot and store at -20 $^\circ~$ 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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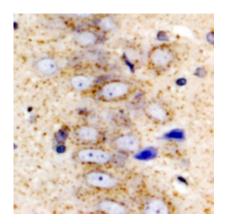
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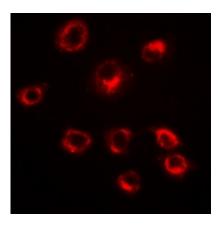
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Western blot analysis of Serpin A10 expression in HepG2 (A), mouse liver (B), rat liver (C) whole cell lysates. (Predicted band size: 50 kD; Observed band size: 51 kD)



Immunohistochemical analysis of Serpin A10 staining in mouse brain 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 Serpin A10 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  $^{\circ}$  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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