

## **Product Data Sheet**

## Anti-ASCT1 Antibody

Catalog #	Source	Reactivity	Applications		
CQA1160	Rabbit	Н, М	WB, IH		
Description	F	Rabbit polyclonal antibody t	o ASCT1		
Immunogen	k	KLH-conjugated synthetic pe	ptide of human ASCT1		
Purification	Т	The antibody was purified b	y immunogen affinity chromatography.		
Specificity	F	Recognizes endogenous leve	els of ASCT1 protein.		
Clonality	F	Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	а	and 0.01% sodium azide.			
Dilution	V	WB (1/500 - 1/2000), IH (1/50	- 1/200)		
Gene Symbol	S	SLC1A4			
Alternative Na	ames A	ASCT1; SATT; Neutral amino	acid transporter A; Alanine/serine/cysteine/threonine		
	t	transporter 1; ASCT-1; SATT;	Solute carrier family 1 member 4		
Entrez Gene	6	6509 (Human); 55963 (Mouse)			
SwissProt	F	P43007 (Human); O35874 (Mouse)			
Storage/Stabi	lity S	Shipped at 4 $^\circ$ C. Upon deliv	ery aliquot and store at -20 $^\circ~$ C for one year. Avoid		
	f	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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kDa A

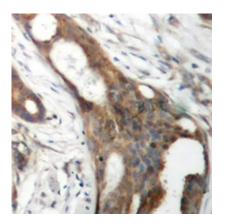
200

140

For research purposes only, not for human use

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Western blot analysis of ASCT1 expression in HepG2 (A) whole cell lysates. (Predicted band size: 55 kD; Observed band size: 55 kD)



Immunohistochemical analysis of ASCT1 staining in human ovarian 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.

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