

## **Product Data Sheet**

### **Anti-LCN1** Antibody

Catalog #	Source	Reactiv	vity	Applications		
CQA3622	Rabbit	H, M, I	3	WB, IH		
Description	Ra	abbit polyclon	al antibody to LCN1			
Immunogen	Re	ecombinant fu	ull length protein of human I	LCN1		
Purification The antibody was purified by			as purified by immunogen a	affinity chromatography.		
Specificity	Re	ecognizes end	ogenous levels of LCN1 prot	ein.		
Clonality	Po	olyclonal				
Conjugation						
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,				
	ar	nd 0.01% sodi	um azide.			
Dilution	W	VB (1/500 - 1/20	000), IH (1/50 - 1/200)			
Gene Symbol	LC	CN1				
Alternative Na	ames V	'EGP; Lipocalin	-1; Tear lipocalin; Tlc; Tear p	realbumin; TP; Von Ebner gland protein;		
	V	'EG protein				
Entrez Gene	Entrez Gene 3933 (Human)					
SwissProt	P	P31025 (Human)				
Storage/Stabi	lity Sł	hipped at 4 $^\circ$ (	C. Upon delivery aliquot and	l store at -20 $^\circ~$ C for one year. Avoid		
	fr	reeze/thaw cyo	cles.			

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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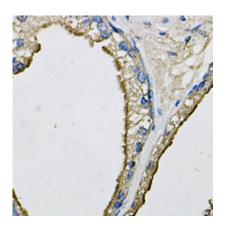
kDa A

70

For research purposes only, not for human use

## **Product Data Sheet**

Western blot analysis of LCN1 expression in HEK293T (A) whole cell lysates. (Predicted band size: 19 kD; Observed band size: 16 kD)



Immunohistochemical analysis of LCN1 staining in human prostate 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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