

## **Product Data Sheet**

## Anti-Defensin alpha 3 Antibody

Catalog #	Source	Reactivity	Applications		
CPA6424	Rabbit	н	WB, IH		
Description	R	Rabbit polyclonal antibody to Defensin alpha 3			
Immunogen	К	H-conjugated synthetic peptide encompassing a sequence within the center			
	re	region of human Defensin alpha 3. The exact sequence is proprietary.			
Purification	Т	he antibody was purified	by immunogen affinity chromatography.		
Specificity	R	ecognizes endogenous le	vels of Defensin alpha 3 protein.		
Clonality	Р	olyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	а	nd 0.01% sodium azide.			
Dilution	V	VB (1/500 - 1/1000), IH (1/	50 - 1/200)		
Gene Symbol	D	DEFA3			
Alternative Na	a <mark>mes</mark> D	EF3; Neutrophil defensir	3; Defensin alpha 3; HNP-3; HP-3; HP3		
Entrez Gene		1668 (Human)			
SwissProt	Р	59666 (Human)			
Storage/Stabi	lity S	hipped at 4°C. Upon deliv	very aliquot and store at -20°C for one year. Avoid		
	fr	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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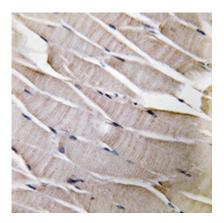
kDa A

70

For research purposes only, not for human use

# **Product Data Sheet**

Western blot analysis of Defensin alpha 3 expression in H1688 (A) whole cell lysates. (Predicted band size: 10 kD; Observed band size: 11 kD)



Immunohistochemical analysis of Defensin alpha 3 staining in human skeletal muscle 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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