

# **Product Data Sheet**

### Anti-Alpha-fetoglobulin Antibody

Catalog #	Source	Reactivity	Applications			
CPA3119	Rabbit	H, M, R	WB, IH			
Description	Rab	Rabbit polyclonal antibody to Alpha-fetoglobulin				
Immunogen	KLH	-conjugated synthetic	peptide encompassing a sequence within the center			
	regi	region of human Alpha-fetoglobulin. The exact sequence is proprietary.				
Purification	The	antibody was purified	by immunogen affinity chromatography.			
Specificity	Reco	ognizes endogenous le	vels of Alpha-fetoglobulin protein.			
Clonality	Poly	rclonal				
Conjugation						
Form	Liqu	id in 0.42% Potassium	phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	and	0.01% sodium azide.				
Dilution	WB	(1/500 - 1/1000), IH (1/5	60 - 1/100)			
Gene Symbol	AFP					
Alternative Na	ames HPA	FP; Alpha-fetoprotein;	Alpha-1-fetoprotein; Alpha-fetoglobulin			
Entrez Gene	174	(Human); 11576 (Mou	se)			
SwissProt	P02	771 (Human); P02772 (	Mouse); P02773 (Rat)			
Storage/Stabi	lity Ship	ped at 4°C. Upon deliv	ery aliquot and store at -20°C for one year. Avoid			
	free	ze/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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Western blot analysis of Alpha-fetoglobulin expression in HEK293T (A), PC12 (B), U2OS (C), mouse liver (D) whole cell lysates. (Predicted band size: 68 kD; Observed band size: 69 kD)



Immunohistochemical analysis of Alpha-fetoglobulin 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.

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