

Product Data Sheet

Anti-H-FABP Antibody

Catalog #	Source	e Reactivity	Applications		
CQA1953	Rabbit	: H, M, R	WB, IH		
Description		Rabbit polyclonal antibody t	o H-FABP		
Immunogen		Recombinant full length prot	ein of human H-FABP		
Purification		The antibody was purified by	/ immunogen affinity chromatography.		
Specificity		Recognizes endogenous leve	ls of H-FABP protein.		
Clonality		Polyclonal			
Conjugation					
Form		Liquid 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 (1/50 - 1/200)			
Gene Symbol		FABP3			
Alternative Names		FABP11; MDGI; Fatty acid-binding protein, heart; Fatty acid-binding protein 3;			
		Heart-type fatty acid-binding	protein; H-FABP; Mammary-derived growth inhibitor;		
		MDGI; Muscle fatty acid-bin	ding protein; M-FABP		
Entrez Gene		2170 (Human); 14077 (Mouse); 79131 (Rat)			
SwissProt		P05413 (Human); P11404 (Mouse); P07483 (Rat)			
Storage/Stabi	lity	Shipped at 4 $^\circ$ C. Upon deliv	ery aliquot and store at -20 $^\circ~$ C for one year. Avoid		
		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

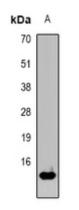
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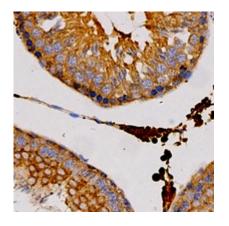
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Western blot analysis of H-FABP expression in mouse heart (A) whole cell lysates. (Predicted band size: 14 kD; Observed band size: 14 kD)



Immunohistochemical analysis of H-FABP staining in human testis 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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