

Product Data Sheet

Anti-BAAT Antibody

Catalog #	Source	Reactivity	Applications		
CQA1844	Rabbit	Н	WB, IH		
Description	Rat	bbit polyclonal antibod [,]	y to BAAT		
Immunogen	Rec	combinant full length p	rotein of human BAAT		
Purification	The	e antibody was purified	by immunogen affinity chromatography.		
Specificity	Rec	cognizes endogenous le	evels of BAAT protein.		
Clonality	Pol	lyclonal			
Conjugation					
Form	Liq	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	and	d 0.01% sodium azide.			
Dilution	WE	3 (1/500 - 1/2000), IH (1/	50 - 1/200)		
Gene Symbol	BA	AT			
Alternative Na	ames Bile	e acid-CoA:amino acid I	N-acyltransferase; BACAT; BAT; Glycine		
	N-c	choloyltransferase; Lon	g-chain fatty-acyl-CoA hydrolase		
Entrez Gene	570) (Human); 12012 (Moเ	use); 29725 (Rat)		
SwissProt	Q1-	4032 (Human); Q91X34	l (Mouse); Q63276 (Rat)		
Storage/Stabi	lity Shi	pped at 4°C. Upon deliv	very aliquot and store at -20°C for one year. Avoid		
	fre	eze/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

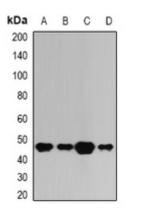
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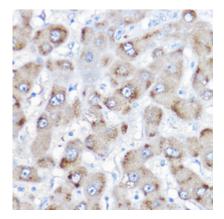


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Western blot analysis of BAAT expression in A549 (A), HepG2 (B), mouse liver (C), rat liver (D) whole cell lysates. (Predicted band size: 46 kD; Observed band size: 40-46 kD)



Immunohistochemical analysis of BAAT staining in human liver 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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