

# **Product Data Sheet**

### **Anti-GAMT Antibody**

Catalog #	Source	Reactivity	Applications
CQA1419	Rabbit	H, M, R	WB, IH, IF/IC
Description		Rabbit polyclonal antibody	to GAMT
Immunogen		Recombinant full length pro	tein of human GAMT
Purification		The antibody was purified b	y immunogen affinity chromatography.
Specificity		Recognizes endogenous lev	els of GAMT protein.
Clonality		Polyclonal	
Conjugation			
Form		Liquid in 0.42% Potassium p	hosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
		and 0.01% sodium azide.	
Dilution		WB (1/500 - 1/2000), IH (1/50	) - 1/200), IF/IC (1/50 - 1/200)
Gene Symbol		GAMT	
Alternative Na	ames	Guanidinoacetate N-methyl	transferase
Entrez Gene		2593 (Human); 14431 (Mou	se)
SwissProt		Q14353 (Human); O35969 (	Mouse); P10868 (Rat)
Storage/Stabi	lity	Shipped at 4°C. Upon delive	ry aliquot and store at -20°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

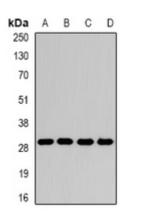
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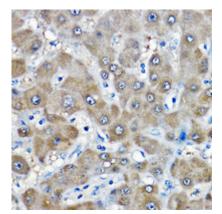




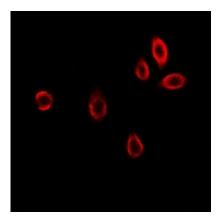
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Western blot analysis of GAMT expression in Jurkat (A), Hela (B), mouse liver (C), rat brain (D) whole cell lysates. (Predicted band size: 26; 29 kD; Observed band size: 29 kD)



Immunohistochemical analysis of GAMT 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.



Immunofluorescent analysis of GAMT staining in U2OS cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a hidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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