

**CUSABIO TECHNOLOGY LLC** 

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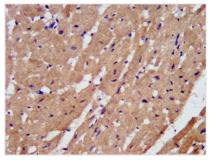
## PAM Antibody

Product Code	CSB-PA017417LA01HU	
Storage	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.	
Uniprot No.	P19021	
Immunogen	Recombinant Human Peptidyl-glycine alpha-amidating monooxygenase protein (338-497AA)	
Raised In	Rabbit	
Species Reactivity	Human	
Tested Applications	ELISA, WB, IHC, IF; Recommended dilution: WB:1:500-1:5000, IHC:1:200-1:500, IF:1:50-1:200	
Form	Liquid	
Conjugate	Non-conjugated	
Storage Buffer	Preservative: 0.03% Proclin 300 Constituents: 50% Glycerol, 0.01M PBS, pH 7.4	
Purification Method	>95%, Protein G purified	
Isotype	lgG	
Clonality	Polyclonal	
Alias	Peptidyl-glycine alpha-amidating monooxygenase (PAM) [Includes: Peptidylglycine alpha-hydroxylating monooxygenase (PHM) (EC 1.14.17.3); Peptidyl-alpha-hydroxyglycine alpha-amidating lyase (EC 4.3.2.5) (Peptidylamidoglycolate lyase) (PAL)], PAM	
Immunogen Species	Homo sapiens (Human)	
Research Area	Neuroscience	
Target Names	PAM	
Image	$250 \text{KD} \rightarrow 130 \text{KD} \rightarrow 95 \text{KD} \rightarrow 72 \text{KD} \rightarrow 55 \text{KD} \rightarrow 36 \text{KD} \rightarrow 72 KD$	Western Blot Positive WB detected in: Hela whole cell lysate All lanes: PAM antibody at 5.6µg/ml Secondary Goat polyclonal to rabbit IgG at 1/50000 dilution Predicted band size: 109, 97, 101, 99, 107 kDa Observed band size: 109 kDa

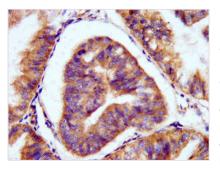
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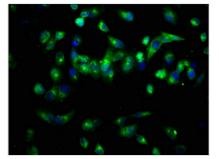
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IHC image of CSB-PA017417LA01HU diluted at 1:200 and staining in paraffin-embedded human heart tissue performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.



IHC image of CSB-PA017417LA01HU diluted at 1:200 and staining in paraffin-embedded human endometrial cancer performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.



Immunofluorescence staining of Hela cells with CSB-PA017417LA01HU at 1:66, counter-stained with DAPI. The cells were fixed in 4% formaldehyde, permeabilized using 0.2% Triton X-100 and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4°C. The secondary antibody was Alexa Fluor 488-congugated AffiniPure Goat Anti-Rabbit IgG(H+L).