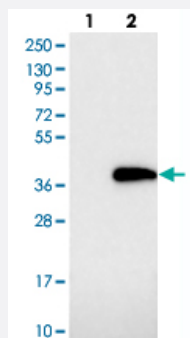


# MMADHC polyclonal antibody

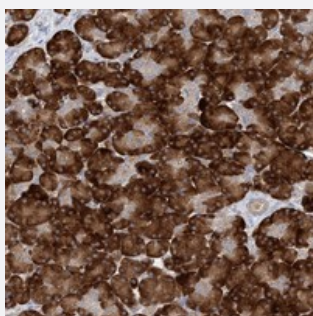
Catalog # PAB23126      Size 100 uL

## Applications



### Western Blot (Transfected lysate)

Western blot analysis of Lane 1: Negative control (vector only transfected HEK293T lysate), Lane 2: Over-expression Lysate (Co-expressed with a C-terminal myc-DDK tag (~3.1 kDa) in mammalian HEK293T cells) with MMADHC polyclonal antibody (Cat # PAB23126).



### Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections)

Immunohistochemical staining of human pancreas with MMADHC polyclonal antibody (Cat # PAB23126) shows strong cytoplasmic positivity in exocrine glandular cells.

## Specification

Product Description	Rabbit polyclonal antibody raised against recombinant MMADHC.
Immunogen	Recombinant protein corresponding to amino acids of human MMADHC.
Sequence	EVLLKFKINGAKEIC <sup>Y</sup> ALRAEGYWADFIDPSSGLAFFGPYTNNTLFETDERYRHLGFSVDDLGCCK VIRHSLW <sup>G</sup> THVVGSIFTNAT
Host	Rabbit
Reactivity	Human
Form	Liquid

<b>Purification</b>	Antigen affinity purification
<b>Isotype</b>	IgG
<b>Recommend Usage</b>	Immunohistochemistry (1:200-1:500) Western Blot (1:250-1:500) The optimal working dilution should be determined by the end user.
<b>Storage Buffer</b>	In PBS, pH 7.2 (40% glycerol, 0.02% sodium azide)
<b>Storage Instruction</b>	Store at 4°C. For long term storage store at -20°C. Aliquot to avoid repeated freezing and thawing.
<b>Note</b>	This product contains sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.

## Applications

- Western Blot (Transfected lysate)

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## Gene Info — MMADHC

<b>Entrez GeneID</b>	<a href="#">27249</a>
<b>Protein Accession#</b>	<a href="#">Q9H3L0</a>
<b>Gene Name</b>	MMADHC
<b>Gene Alias</b>	C2orf25, CL25022
<b>Gene Description</b>	methylmalonic aciduria (cobalamin deficiency) cblD type, with homocystinuria
<b>Gene Ontology</b>	<a href="#">Hyperlink</a>

**Gene Summary**

This gene encodes a mitochondrial protein that is involved in an early step of vitamin B12 metabolism. Vitamin B12 (cobalamin) is essential for normal development and survival in humans. Mutations in this gene cause methylmalonic aciduria and homocystinuria type cblD (MMADHC), a disorder of cobalamin metabolism that is characterized by decreased levels of the coenzymes adenosylcobalamin and methylcobalamin. Pseudogenes have been identified on chromosomes 11 and X

**Other Designations**

protein C2orf25, mitochondrial

**Disease**

- [Cardiovascular Diseases](#)
- [Diabetes Mellitus](#)
- [Disease Progression](#)
- [Disease Susceptibility](#)
- [Edema](#)
- [HIV Infections](#)