

AMPD3 Antibody (Center) Blocking Peptide
Synthetic peptide
Catalog # BP7869c**Specification**

AMPD3 Antibody (Center) Blocking Peptide - Product InformationPrimary Accession [Q01432](#)**AMPD3 Antibody (Center) Blocking Peptide - Additional Information****Gene ID** 272**Other Names**

AMP deaminase 3, AMP deaminase isoform E, Erythrocyte AMP deaminase, AMPD3

Target/Specificity

The synthetic peptide sequence used to generate the antibody [AP7869c](/products/AP7869c) was selected from the Center region of human AMPD3. A 10 to 100 fold molar excess to antibody is recommended. Precise conditions should be optimized for a particular assay.

Format

Peptides are lyophilized in a solid powder format. Peptides can be reconstituted in solution using the appropriate buffer as needed.

Storage

Maintain refrigerated at 2-8°C for up to 6 months. For long term storage store at -20°C.

Precautions

This product is for research use only. Not for use in diagnostic or therapeutic procedures.

AMPD3 Antibody (Center) Blocking Peptide - Protein Information**Name** AMPD3 ([HGNC:470](#))**Function**

AMP deaminase plays a critical role in energy metabolism.

AMPD3 Antibody (Center) Blocking Peptide - Protocols

Provided below are standard protocols that you may find useful for product applications.

- [Blocking Peptides](#)

AMPD3 Antibody (Center) Blocking Peptide - Images**AMPD3 Antibody (Center) Blocking Peptide - Background**

AMPD3 is a member of the AMP deaminase gene family. This protein is a highly regulated enzyme that catalyzes the hydrolytic deamination of adenosine monophosphate to inosine monophosphate, a branch point in the adenylate catabolic pathway. The protein is the erythrocyte (E) isoforms, whereas other family members isoforms predominate in muscle (M) and liver (L) cells. Mutations in this gene lead to the clinically asymptomatic, autosomal recessive condition erythrocyte AMP deaminase deficiency.

AMPD3 Antibody (Center) Blocking Peptide - References

Mahnke-Zizelman D.K., Eddy R. Biochim. Biophys. Acta 1306:75-92(1996) Yamada Y., Goto H., Wakamatsu N., Ogasawara N. Hum. Mutat. 17:78-78(2001)