

**BLOC1S3 Blocking Peptide (C-term)**  
**Synthetic peptide**  
**Catalog # BP19902B****Specification**

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**BLOC1S3 Blocking Peptide (C-term) - Product Information**

Primary Accession [Q6QNY0](#)  
Other Accession [NP\\_997715.1](#)

**BLOC1S3 Blocking Peptide (C-term) - Additional Information**

**Gene ID** 388552

**Other Names**

Biogenesis of lysosome-related organelles complex 1 subunit 3, BLOC-1 subunit 3, BLOC1S3, BLOS3

**Target/Specificity**

The synthetic peptide sequence is selected from aa 191-202 of HUMAN BLOC1S3

**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.

**BLOC1S3 Blocking Peptide (C-term) - Protein Information**

**Name** BLOC1S3

**Synonyms** BLOS3

**Function**

Component of the BLOC-1 complex, a complex that is required for normal biogenesis of lysosome-related organelles (LRO), such as platelet dense granules and melanosomes. In concert with the AP-3 complex, the BLOC-1 complex is required to target membrane protein cargos into vesicles assembled at cell bodies for delivery into neurites and nerve terminals. The BLOC-1 complex, in association with SNARE proteins, is also proposed to be involved in neurite extension. Plays a role in intracellular vesicle trafficking.

**Cellular Location**

Cytoplasm.

## **BLOC1S3 Blocking Peptide (C-term) - Protocols**

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

- [Blocking Peptides](#)

## **BLOC1S3 Blocking Peptide (C-term) - Images**

## **BLOC1S3 Blocking Peptide (C-term) - Background**

This gene encodes a protein that is a component of the BLOC1 multi-subunit protein complex. This complex is necessary for the biogenesis of specialized organelles of the endosomal-lysosomal system, including platelet dense granules and melanosomes. Mutations in this gene cause Hermansky-Pudlak syndrome 8, a disease characterized by lysosomal storage defects, bleeding due to platelet storage pool deficiency, and oculocutaneous albinism.

## **BLOC1S3 Blocking Peptide (C-term) - References**

Seshadri, S., et al. JAMA 303(18):1832-1840(2010)  
Lowe, J.K., et al. PLoS Genet. 5 (2), E1000365 (2009) :  
Olsen, J.V., et al. Cell 127(3):635-648(2006)  
Morgan, N.V., et al. Am. J. Hum. Genet. 78(1):160-166(2006)  
Starcevic, M., et al. J. Biol. Chem. 279(27):28393-28401(2004)