

## **Product datasheet for SC308648**

## BLOC1S3 (NM 212550) Human Untagged Clone

## **Product data:**

**Product Type:** Expression Plasmids

**Product Name:** BLOC1S3 (NM\_212550) Human Untagged Clone

Tag: Tag Free Symbol: BLOC1S3

Synonyms: BLOS3; HPS8; RP

Mammalian Cell None

Selection:

Vector: pCMV6-XL5

E. coli Selection: Ampicillin (100 ug/mL)

Fully Sequenced ORF: >OriGene ORF sequence for NM\_212550 edited

CGGGCCTAG

Restriction Sites: Notl-Notl
ACCN: NM\_212550
Insert Size: 2400 bp

**OTI Disclaimer:** Our molecular clone sequence data has been matched to the reference identifier above as a

point of reference. Note that the complete sequence of our molecular clones may differ from the sequence published for this corresponding reference, e.g., by representing an alternative

RNA splicing form or single nucleotide polymorphism (SNP).

**OTI Annotation:** This clone has been fully sequenced and found to be a perfect match to the protein

associated with this reference, NM 212550.2



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## BLOC1S3 (NM\_212550) Human Untagged Clone - SC308648

**Components:** The ORF clone is ion-exchange column purified and shipped in a 2D barcoded Matrix tube

containing 10ug of transfection-ready, dried plasmid DNA (reconstitute with 100 ul of water).

**Reconstitution Method:** 1. Centrifuge at 5,000xg for 5min.

2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.

3. Close the tube and incubate for 10 minutes at room temperature.

4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid

at the bottom.

5. Store the suspended plasmid at -20°C. The DNA is stable for at least one year from date of

shipping when stored at -20°C.

RefSeq: <u>NM 212550.1</u>, <u>NP 997715.1</u>

 RefSeq Size:
 609 bp

 RefSeq ORF:
 609 bp

 Locus ID:
 388552

 UniProt ID:
 Q6QNY0

 Cytogenetics:
 19q13.32

**Gene Summary:** 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. [provided by RefSeq, Jul 2008]