

## **OBFC1 Antibody (C-term) Blocking Peptide**

Synthetic peptide Catalog # BP16810b

### **Specification**

### **OBFC1 Antibody (C-term) Blocking Peptide - Product Information**

**Primary Accession** 

**Q9H668** 

# **OBFC1** Antibody (C-term) Blocking Peptide - Additional Information

**Gene ID** 79991

#### **Other Names**

CST complex subunit STN1, Oligonucleotide/oligosaccharide-binding fold-containing protein 1, Suppressor of cdc thirteen homolog, OBFC1, STN1

#### **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.

### **OBFC1** Antibody (C-term) Blocking Peptide - Protein Information

Name STN1 (HGNC:26200)

Synonyms OBFC1

### **Function**

Component of the CST complex proposed to act as a specialized replication factor promoting DNA replication under conditions of replication stress or natural replication barriers such as the telomere duplex. The CST complex binds single-stranded DNA with high affinity in a sequence-independent manner, while isolated subunits bind DNA with low affinity by themselves. Initially the CST complex has been proposed to protect telomeres from DNA degradation (PubMed:<a href="http://www.uniprot.org/citations/19854130" target="\_blank">19854130</a>). However, the CST complex has been shown to be involved in several aspects of telomere replication. The CST complex inhibits telomerase and is involved in telomere length homeostasis; it is proposed to bind to newly telomerase-synthesized 3' overhangs and to terminate telomerase action implicating the association with the ACD:POT1 complex thus interfering with its telomerase stimulation activity. The CST complex is also proposed to be involved in fill-in synthesis of the telomeric C-strand probably implicating recruitment and activation of DNA polymerase alpha (PubMed:<a href="http://www.uniprot.org/citations/22964711" target="\_blank">22964711</a>
PubMed:<a href="http://www.uniprot.org/citations/22763445" target="\_blank">22763445</a>). The CST complex facilitates recovery from many forms of exogenous DNA damage; seems to be



involved in the re-initiation of DNA replication at repaired forks and/or dormant origins (PubMed:<a href="http://www.uniprot.org/citations/25483097" target="\_blank">25483097</a>). Required for efficient replication of the duplex region of the telomere. Promotes efficient replication of lagging-strand telomeres (PubMed:<a

href="http://www.uniprot.org/citations/22863775" target="\_blank">22863775</a>, PubMed:<a href="http://www.uniprot.org/citations/22964711" target="\_blank">22964711</a>). Promotes general replication start following replication-fork stalling implicating new origin firing (PubMed:<a href="http://www.uniprot.org/citations/22863775" target="\_blank">22863775</a>). May be in involved in C-strand fill-in during late S/G2 phase independent of its role in telomere duplex replication (PubMed:<a href="http://www.uniprot.org/citations/23142664" target="\_blank">23142664</a>).

**Cellular Location** 

Nucleus. Chromosome, telomere

## **OBFC1 Antibody (C-term) Blocking Peptide - Protocols**

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

Blocking Peptides

**OBFC1 Antibody (C-term) Blocking Peptide - Images** 

### OBFC1 Antibody (C-term) Blocking Peptide - Background

OBFC1 and C17ORF68 (MIM 613129) are subunits of an alphaaccessory factor (AAF) that stimulates the activity of DNApolymerase-alpha-primase (see MIM 176636), the enzyme that initiates DNA replication (Casteel et al., 2009 [PubMed 19119139]).OBFC1 also appears to function in a telomere-associated complexwith C17ORF68 and TEN1 (C17ORF106; MIM 613130) (Miyake et al., 2009[PubMed 19854130]).

# **OBFC1 Antibody (C-term) Blocking Peptide - References**

Levy, D., et al. Proc. Natl. Acad. Sci. U.S.A. 107(20):9293-9298(2010)Miyake, Y., et al. Mol. Cell 36(2):193-206(2009)Wan, M., et al. J. Biol. Chem. 284(39):26725-26731(2009)Casteel, D.E., et al. J. Biol. Chem. 284(9):5807-5818(2009)Lamesch. P., et al. Genomics 89(3):307-315(2007)