

**DC12 Antibody (C-term) Blocking peptide**  
**Synthetic peptide**  
**Catalog # BP10997b****Specification**

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**DC12 Antibody (C-term) Blocking peptide - Product Information**Primary Accession  
Other Accession[O96FZ2](#)  
[NP\\_001006109.1](#), [NP\\_064572.2](#)**DC12 Antibody (C-term) Blocking peptide - Additional Information****Gene ID** 56941**Other Names**

Embryonic stem cell-specific 5-hydroxymethylcytosine-binding protein, ES cell-specific 5hmC-binding protein, Putative peptidase SRAPD1, 34--, SRAP domain-containing protein 1, HMCES, C3orf37, DC12, SRAPD1

**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.

**DC12 Antibody (C-term) Blocking peptide - Protein Information****Name** HMCES {ECO:0000303|PubMed:30554877, ECO:0000312|HGNC:HGNC:24446}**Function**

Sensor of abasic sites in single-stranded DNA (ssDNA) required to preserve genome integrity by promoting error-free repair of abasic sites (PubMed:<a href="http://www.uniprot.org/citations/30554877" target="\_blank">30554877</a>, PubMed:<a href="http://www.uniprot.org/citations/31235913" target="\_blank">31235913</a>, PubMed:<a href="http://www.uniprot.org/citations/31235915" target="\_blank">31235915</a>, PubMed:<a href="http://www.uniprot.org/citations/32307824" target="\_blank">32307824</a>, PubMed:<a href="http://www.uniprot.org/citations/32492421" target="\_blank">32492421</a>). Acts as an enzyme that recognizes and binds abasic sites in ssDNA at replication forks and chemically modifies the lesion by forming a covalent cross-link with DNA: forms a stable thiazolidine linkage between a ring-opened abasic site and the alpha-amino and sulfhydryl substituents of its N-terminal catalytic cysteine residue (PubMed:<a href="http://www.uniprot.org/citations/30554877" target="\_blank">30554877</a>, PubMed:<a href="http://www.uniprot.org/citations/31235913" target="\_blank">31235913</a>). Promotes error-free repair by protecting abasic sites from translesion synthesis (TLS) polymerases and endonucleases that are error-prone and would generate mutations and double-strand breaks

(PubMed:<a href="http://www.uniprot.org/citations/30554877" target="\_blank">30554877</a>). The HMCES DNA- protein cross-link is then either reversed or degraded (PubMed:<a href="http://www.uniprot.org/citations/30554877" target="\_blank">30554877</a>, PubMed:<a href="http://www.uniprot.org/citations/36608669" target="\_blank">36608669</a>, PubMed:<a href="http://www.uniprot.org/citations/37519246" target="\_blank">37519246</a>, PubMed:<a href="http://www.uniprot.org/citations/37950866" target="\_blank">37950866</a>). HMCES is able to catalyze the reversal of its thiazolidine cross-link and cycle between a cross-link and a non-cross-linked state depending on DNA context: mediates self-reversal of the thiazolidine cross-link in double stranded DNA, allowing APEX1 to initiate downstream repair of abasic sites (PubMed:<a href="http://www.uniprot.org/citations/37519246" target="\_blank">37519246</a>, PubMed:<a href="http://www.uniprot.org/citations/37950866" target="\_blank">37950866</a>). The HMCES DNA-protein cross-link can also be degraded by the SPRTN metalloprotease following unfolding by the BRIP1/FANCD1 helicase (PubMed:<a href="http://www.uniprot.org/citations/36608669" target="\_blank">36608669</a>). Has preference for ssDNA, but can also accommodate double-stranded DNA with 3' or 5' overhang (dsDNA), and dsDNA-ssDNA 3' junction (PubMed:<a href="http://www.uniprot.org/citations/31235915" target="\_blank">31235915</a>, PubMed:<a href="http://www.uniprot.org/citations/31806351" target="\_blank">31806351</a>). Plays a protective role during somatic hypermutation of immunoglobulin genes in B-cells: acts via its ability to form covalent cross-links with abasic sites, thereby limiting the accumulation of deletions in somatic hypermutation target regions (PubMed:<a href="http://www.uniprot.org/citations/35450882" target="\_blank">35450882</a>). Also involved in class switch recombination (CSR) in B-cells independently of the formation of a DNA-protein cross-link: acts by binding and protecting ssDNA overhangs to promote DNA double-strand break repair through the microhomology-mediated alternative-end-joining (Alt-EJ) pathway (By similarity). Acts as a protease: mediates autocatalytic processing of its N-terminal methionine in order to expose the catalytic cysteine (By similarity).

#### **Cellular Location**

Chromosome. Note=Recruited to chromatin following DNA damage (PubMed:30554877) Localizes to replication forks (PubMed:30554877)

#### **DC12 Antibody (C-term) Blocking peptide - Protocols**

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

- [Blocking Peptides](#)

#### **DC12 Antibody (C-term) Blocking peptide - Images**

#### **DC12 Antibody (C-term) Blocking peptide - References**

Gerhard, D.S., et al. Genome Res. 14 (10B), 2121-2127 (2004) :