

**BRD4 Antibody (C-term) Blocking Peptide**  
**Synthetic peptide**  
**Catalog # BP17153b****Specification**

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**BRD4 Antibody (C-term) Blocking Peptide - Product Information**Primary Accession [O60885](#)**BRD4 Antibody (C-term) Blocking Peptide - Additional Information****Gene ID** 23476**Other Names**

Bromodomain-containing protein 4, Protein HUNK1, BRD4, HUNK1

**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.

**BRD4 Antibody (C-term) Blocking Peptide - Protein Information****Name** BRD4**Synonyms** HUNK1**Function**

Chromatin reader protein that recognizes and binds acetylated histones and plays a key role in transmission of epigenetic memory across cell divisions and transcription regulation (PubMed:<a href="http://www.uniprot.org/citations/23086925" target="\_blank">23086925</a>, PubMed:<a href="http://www.uniprot.org/citations/23317504" target="\_blank">23317504</a>, PubMed:<a href="http://www.uniprot.org/citations/20871596" target="\_blank">20871596</a>, PubMed:<a href="http://www.uniprot.org/citations/29176719" target="\_blank">29176719</a>). Remains associated with acetylated chromatin throughout the entire cell cycle and provides epigenetic memory for postmitotic G1 gene transcription by preserving acetylated chromatin status and maintaining high-order chromatin structure (PubMed:<a href="http://www.uniprot.org/citations/23589332" target="\_blank">23589332</a>, PubMed:<a href="http://www.uniprot.org/citations/23317504" target="\_blank">23317504</a>, PubMed:<a href="http://www.uniprot.org/citations/22334664" target="\_blank">22334664</a>). During interphase, plays a key role in regulating the transcription of signal- inducible genes by associating with the P-TEFb complex and recruiting it to promoters (PubMed:<a href="http://www.uniprot.org/citations/23589332" target="\_blank">23589332</a>, PubMed:<a href="http://www.uniprot.org/citations/19596240" target="\_blank">19596240</a>, PubMed:<a href="http://www.uniprot.org/citations/19596240" target="\_blank">19596240</a>, PubMed:<a href="http://www.uniprot.org/citations/19596240" target="\_blank">19596240</a>).

[16109377](http://www.uniprot.org/citations/16109377), PubMed: [16109376](http://www.uniprot.org/citations/16109376), PubMed: [24360279](http://www.uniprot.org/citations/24360279)). Also recruits P-TEFb complex to distal enhancers, so called anti-pause enhancers in collaboration with JMJD6 (PubMed: [23589332](http://www.uniprot.org/citations/23589332), PubMed: [19596240](http://www.uniprot.org/citations/19596240), PubMed: [16109377](http://www.uniprot.org/citations/16109377), PubMed: [16109376](http://www.uniprot.org/citations/16109376), PubMed: [24360279](http://www.uniprot.org/citations/24360279)). BRD4 and JMJD6 are required to form the transcriptionally active P-TEFb complex by displacing negative regulators such as HEXIM1 and 7SKsnRNA complex from P-TEFb, thereby transforming it into an active form that can then phosphorylate the C-terminal domain (CTD) of RNA polymerase II (PubMed: [23589332](http://www.uniprot.org/citations/23589332), PubMed: [19596240](http://www.uniprot.org/citations/19596240), PubMed: [16109377](http://www.uniprot.org/citations/16109377), PubMed: [16109376](http://www.uniprot.org/citations/16109376), PubMed: [24360279](http://www.uniprot.org/citations/24360279)). Regulates differentiation of naive CD4(+) T-cells into T-helper Th17 by promoting recruitment of P-TEFb to promoters (By similarity). Promotes phosphorylation of 'Ser-2' of the C-terminal domain (CTD) of RNA polymerase II (PubMed: [23086925](http://www.uniprot.org/citations/23086925)). According to a report, directly acts as an atypical protein kinase and mediates phosphorylation of 'Ser-2' of the C-terminal domain (CTD) of RNA polymerase II; these data however need additional evidences in vivo (PubMed: [22509028](http://www.uniprot.org/citations/22509028)). In addition to acetylated histones, also recognizes and binds acetylated RELA, leading to further recruitment of the P-TEFb complex and subsequent activation of NF-kappa-B (PubMed: [19103749](http://www.uniprot.org/citations/19103749)). Also acts as a regulator of p53/TP53-mediated transcription: following phosphorylation by CK2, recruited to p53/TP53 specific target promoters (PubMed: [23317504](http://www.uniprot.org/citations/23317504)).

### Cellular Location

Nucleus. Chromosome. Note=Associates with acetylated chromatin (PubMed:21890894, PubMed:16109376). Released from chromatin upon deacetylation of histones that can be triggered by different signals such as activation of the JNK pathway or nocodazole treatment (PubMed:21890894, PubMed:16109376). Preferentially localizes to mitotic chromosomes, while it does not localize to meiotic chromosomes (PubMed:21890894, PubMed:16109376).

### Tissue Location

Ubiquitously expressed.

### BRD4 Antibody (C-term) Blocking Peptide - Protocols

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

- [Blocking Peptides](#)

### BRD4 Antibody (C-term) Blocking Peptide - Images

### BRD4 Antibody (C-term) Blocking Peptide - Background

The protein encoded by this gene is homologous to the murine protein MCAP, which associates with chromosomes during mitosis, and to the human RING3 protein, a serine/threonine kinase. Each of these proteins contains two bromodomains, a conserved sequence motif which may be involved in chromatin targeting. This gene has been implicated as the chromosome 19 target of translocation t(15;19)(q13;p13.1), which defines an upper respiratory tract carcinoma in young people. Two alternatively spliced transcript variants have been described. [provided by RefSeq].

**BRD4 Antibody (C-term) Blocking Peptide - References**

Reynoird, N., et al. EMBO J. 29(17):2943-2952(2010)Dow, E.C., et al. J. Cell. Physiol. 224(1):84-93(2010)Yan, J., et al. J. Virol. 84(1):76-87(2010)Weidner-Glunde, M., et al. Front. Biosci. 15, 537-549 (2010) :You, J., et al. Mol. Cell. Biol. 29(18):5094-5103(2009)