

**CUL4A Antibody (N-term) Blocking peptide**  
**Synthetic peptide**  
**Catalog # BP13873a**

**Specification**

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**CUL4A Antibody (N-term) Blocking peptide - Product Information**

Primary Accession [Q13619](#)

**CUL4A Antibody (N-term) Blocking peptide - Additional Information**

**Gene ID** 8451

**Other Names**

Cullin-4A, CUL-4A, CUL4A

**Target/Specificity**

The synthetic peptide sequence used to generate the antibody AP13873a was selected from the N-term region of CUL4A. A 10 to 100 fold molar excess to antibody is recommended. Precise conditions should be optimized for a particular assay.

**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.

**CUL4A Antibody (N-term) Blocking peptide - Protein Information**

**Name** CUL4A {ECO:0000303|PubMed:9721878, ECO:0000312|HGNC:HGNC:2554}

**Function**

Core component of multiple cullin-RING-based E3 ubiquitin- protein ligase complexes which mediate the ubiquitination of target proteins (PubMed:<a href="http://www.uniprot.org/citations/14578910" target="\_blank">14578910</a>, PubMed:<a href="http://www.uniprot.org/citations/15811626" target="\_blank">15811626</a>, PubMed:<a href="http://www.uniprot.org/citations/15548678" target="\_blank">15548678</a>, PubMed:<a href="http://www.uniprot.org/citations/15448697" target="\_blank">15448697</a>, PubMed:<a href="http://www.uniprot.org/citations/14739464" target="\_blank">14739464</a>, PubMed:<a href="http://www.uniprot.org/citations/16678110" target="\_blank">16678110</a>, PubMed:<a href="http://www.uniprot.org/citations/17041588" target="\_blank">17041588</a>, PubMed:<a href="http://www.uniprot.org/citations/24209620" target="\_blank">24209620</a>, PubMed:<a href="http://www.uniprot.org/citations/30166453" target="\_blank">30166453</a>, PubMed:<a href="http://www.uniprot.org/citations/33854232" target="\_blank">33854232</a>, PubMed:<a href="http://www.uniprot.org/citations/33854239" target="\_blank">33854239</a>). As a scaffold

protein may contribute to catalysis through positioning of the substrate and the ubiquitin-conjugating enzyme (PubMed:<a href="http://www.uniprot.org/citations/14578910" target="\_blank">14578910</a>, PubMed:<a href="http://www.uniprot.org/citations/15811626" target="\_blank">15811626</a>, PubMed:<a href="http://www.uniprot.org/citations/15548678" target="\_blank">15548678</a>, PubMed:<a href="http://www.uniprot.org/citations/15448697" target="\_blank">15448697</a>, PubMed:<a href="http://www.uniprot.org/citations/14739464" target="\_blank">14739464</a>, PubMed:<a href="http://www.uniprot.org/citations/16678110" target="\_blank">16678110</a>, PubMed:<a href="http://www.uniprot.org/citations/17041588" target="\_blank">17041588</a>, PubMed:<a href="http://www.uniprot.org/citations/24209620" target="\_blank">24209620</a>). The E3 ubiquitin-protein ligase activity of the complex is dependent on the neddylation of the cullin subunit and is inhibited by the association of the deneddylated cullin subunit with TIP120A/CAND1 (PubMed:<a href="http://www.uniprot.org/citations/14578910" target="\_blank">14578910</a>, PubMed:<a href="http://www.uniprot.org/citations/15811626" target="\_blank">15811626</a>, PubMed:<a href="http://www.uniprot.org/citations/15548678" target="\_blank">15548678</a>, PubMed:<a href="http://www.uniprot.org/citations/15448697" target="\_blank">15448697</a>, PubMed:<a href="http://www.uniprot.org/citations/14739464" target="\_blank">14739464</a>, PubMed:<a href="http://www.uniprot.org/citations/16678110" target="\_blank">16678110</a>, PubMed:<a href="http://www.uniprot.org/citations/17041588" target="\_blank">17041588</a>, PubMed:<a href="http://www.uniprot.org/citations/24209620" target="\_blank">24209620</a>). The functional specificity of the E3 ubiquitin-protein ligase complex depends on the variable substrate recognition component (PubMed:<a href="http://www.uniprot.org/citations/14578910" target="\_blank">14578910</a>, PubMed:<a href="http://www.uniprot.org/citations/15811626" target="\_blank">15811626</a>, PubMed:<a href="http://www.uniprot.org/citations/15548678" target="\_blank">15548678</a>, PubMed:<a href="http://www.uniprot.org/citations/15448697" target="\_blank">15448697</a>, PubMed:<a href="http://www.uniprot.org/citations/14739464" target="\_blank">14739464</a>, PubMed:<a href="http://www.uniprot.org/citations/16678110" target="\_blank">16678110</a>, PubMed:<a href="http://www.uniprot.org/citations/17041588" target="\_blank">17041588</a>, PubMed:<a href="http://www.uniprot.org/citations/24209620" target="\_blank">24209620</a>). DCX(DET1-COP1) directs ubiquitination of JUN (PubMed:<a href="http://www.uniprot.org/citations/14739464" target="\_blank">14739464</a>). DCX(DDB2) directs ubiquitination of XPC (PubMed:<a href="http://www.uniprot.org/citations/15811626" target="\_blank">15811626</a>). DCX(DDB2) ubiquitinates histones H3-H4 and is required for efficient histone deposition during replication-coupled (H3.1) and replication-independent (H3.3) nucleosome assembly, probably by facilitating the transfer of H3 from ASF1A/ASF1B to other chaperones involved in histone deposition (PubMed:<a href="http://www.uniprot.org/citations/16678110" target="\_blank">16678110</a>, PubMed:<a href="http://www.uniprot.org/citations/17041588" target="\_blank">17041588</a>, PubMed:<a href="http://www.uniprot.org/citations/24209620" target="\_blank">24209620</a>). DCX(DTL) plays a role in PCNA-dependent polyubiquitination of CDT1 and MDM2-dependent ubiquitination of p53/TP53 in response to radiation-induced DNA damage and during DNA replication (PubMed:<a href="http://www.uniprot.org/citations/14578910" target="\_blank">14578910</a>, PubMed:<a href="http://www.uniprot.org/citations/15548678" target="\_blank">15548678</a>, PubMed:<a href="http://www.uniprot.org/citations/15448697" target="\_blank">15448697</a>). DCX(DTL) directs autoubiquitination of DTL (PubMed:<a href="http://www.uniprot.org/citations/23478445" target="\_blank">23478445</a>). In association with DDB1 and SKP2 probably is involved in ubiquitination of CDKN1B/p27kip (PubMed:<a href="http://www.uniprot.org/citations/16537899" target="\_blank">16537899</a>). Is involved in ubiquitination of HOXA9 (PubMed:<a href="http://www.uniprot.org/citations/14609952" target="\_blank">14609952</a>). The DDB1-CUL4A-DTL E3 ligase complex regulates the circadian clock function by mediating the ubiquitination and degradation of CRY1 (PubMed:<a href="http://www.uniprot.org/citations/26431207" target="\_blank">26431207</a>). A number of DCX complexes (containing either TRPC4AP or DCAF12 as substrate-recognition component) are part of the DesCEND (destruction via C-end degrons) pathway, which recognizes a C-degron located at the extreme C terminus of target proteins, leading to their ubiquitination and degradation (PubMed:<a href="http://www.uniprot.org/citations/29779948" target="\_blank">29779948</a>). The DCX(AMBRA1) complex is a master regulator of the

transition from G1 to S cell phase by mediating ubiquitination of phosphorylated cyclin-D (CCND1, CCND2 and CCND3) (PubMed:<a href="http://www.uniprot.org/citations/33854232" target="\_blank">33854232</a>, PubMed:<a href="http://www.uniprot.org/citations/33854239" target="\_blank">33854239</a>). The DCX(AMBRA1) complex also acts as a regulator of Cul5-RING (CRL5) E3 ubiquitin-protein ligase complexes by mediating ubiquitination and degradation of Elongin-C (ELOC) component of CRL5 complexes (PubMed:<a href="http://www.uniprot.org/citations/30166453" target="\_blank">30166453</a>). With CUL4B, contributes to ribosome biogenesis (PubMed:<a href="http://www.uniprot.org/citations/26711351" target="\_blank">26711351</a>).

### **CUL4A Antibody (N-term) Blocking peptide - Protocols**

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

- [Blocking Peptides](#)

### **CUL4A Antibody (N-term) Blocking peptide - Images**

### **CUL4A Antibody (N-term) Blocking peptide - Background**

CUL4A is the ubiquitin ligase component of a multimeric complex involved in the degradation of DNA damage-response proteins(Liu et al., 2009 [PubMed 19481525]).

### **CUL4A Antibody (N-term) Blocking peptide - References**

Aggarwal, P., et al. Cancer Cell 18(4):329-340(2010)Abbas, T., et al. Mol. Cell 40(1):9-21(2010)Lv, X.B., et al. J. Biol. Chem. 285(24):18234-18240(2010)Kerzendorfer, C., et al. Hum. Mol. Genet. 19(7):1324-1334(2010)Melchor, L., et al. Breast Cancer Res. 11 (6), R86 (2009) :