

CDT2 Antibody

Rabbit mAb Catalog # AP91651

Product Information

| Application | WB, IHC, IF, FC, ICC, IHF |
|-------------------|--------------------------------|
| Primary Accession | <u>Q9NZJ0</u> |
| Reactivity | Rat, Human, Mouse |
| Clonality | Monoclonal |
| Other Names | CDW1; DCAF2; Dtl; L2DTL; RAMP; |
| Isotype | Rabbit IgG Rabbit |
| Host | |
| Calculated MW | 79468 |

Additional Information

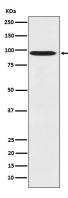
| Dilution Purification Immunogen Description | WB 1:1000~1:5000 IHC 1:50~1:100 ICC/IF 1:50~1:100 FC 1:80 Affinity-chromatography A synthesized peptide derived from human CDT2 Substrate-specific adapter of a DCX (DDB1-CUL4-X-box) E3 ubiquitin-protein ligase complex required for cell cycle control, DNA damage response and translesion DNA synthesis. The DCX(DTL) complex, also named CRL4(CDT2) complex, mediates the polyubiquitination and subsequent degradation of CDT1 and CDKN1A/p21(CIP1). |
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| Storage Condition and Buffer | |

Protein Information

| Name | DTL |
|----------|---|
| Synonyms | CDT2, CDW1, DCAF2, L2DTL, RAMP |
| Function | Substrate-specific adapter of a DCX (DDB1-CUL4-X-box) E3 ubiquitin-protein ligase complex required for cell cycle control, DNA damage response and translesion DNA synthesis. The DCX(DTL) complex, also named CRL4(CDT2) complex, mediates the polyubiquitination and subsequent degradation of CDT1, CDKN1A/p21(CIP1), FBH1, KMT5A and SDE2 (PubMed:16861906, PubMed:16949367, PubMed:16964240, PubMed:17085480, PubMed:18703516, PubMed:18794347, PubMed:18794348, PubMed:19332548, PubMed:20129063, PubMed:23478441, PubMed:23478445, PubMed:23677613, PubMed:27906959). CDT1 degradation in response to DNA damage is necessary to ensure proper cell cycle regulation of DNA replication (PubMed:16861906, PubMed:16949367, PubMed:17085480). CDKN1A/p21(CIP1) degradation during S phase or |

| | following UV irradiation is essential to control replication licensing (PubMed: <u>18794348</u> , PubMed: <u>19332548</u>). KMT5A degradation is also important for a proper regulation of mechanisms such as TGF-beta signaling, cell cycle progression, DNA repair and cell migration (PubMed: <u>23478445</u>). Most substrates require their interaction with PCNA for their polyubiquitination: substrates interact with PCNA via their PIP-box, and those containing the 'K+4' motif in the PIP box, recruit the DCX(DTL) complex, leading to their degradation. In undamaged proliferating cells, the DCX(DTL) complex also promotes the 'Lys-164' monoubiquitination of PCNA, thereby being involved in PCNA- dependent translesion DNA synthesis (PubMed: <u>20129063</u> , PubMed: <u>23478441</u> , PubMed: <u>23478445</u> , PubMed: <u>23677613</u>). The DDB1-CUL4A-DTL E3 ligase complex regulates the circadian clock function by mediating the ubiquitination and degradation of CRY1 (PubMed: <u>26431207</u>). |
|-------------------|--|
| Cellular Location | Nucleus. Nucleus membrane; Peripheral membrane protein; Nucleoplasmic side. Cytoplasm, cytoskeleton, microtubule organizing center, centrosome. Chromosome Note=Nuclear matrix-associated protein. Translocates from the interphase nucleus to the metaphase cytoplasm during mitosis |
| Tissue Location | Expressed in placenta and testis, very low expression seen in skeletal muscle. Detected in all hematopoietic tissues examined, with highest expression in thymus and bone marrow. A low level detected in the spleen and lymph node, and barely detectable level in the peripheral leukocytes. RA treatment down-regulated the expression in NT2 cell. |

Images



Western blot analysis of CDT2 expression in 293T cell lysate.

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