

DCAF1 抗原(重组蛋白)

中文名称: DCAF1 抗原(重组蛋白)

英文名称: DCAF1 Antigen (Recombinant Protein)

别 名: DDB1 and CUL4 associated factor; RIP; VPRBP

相关类别: 抗原

储 存: 冷冻(-20℃)

概述

Fusion protein corresponding to a region derived from 1308-1507 amino acids of human DCAF1 $\,$

技术规格

Full name:	DDB1 and CUL4 associated factor
Synonyms:	RIP; VPRBP
Swissprot:	Q9Y4B6
Gene Accession:	BC022792
Purity:	>85%, as determined by Coomassie blue stained SDS-PAGE
Expression system:	Escherichia coli
Tags:	His tag C-Terminus, GST tag N-Terminus
Background:	Acts both as a substrate recognition component of E3 ubiquitin-protein ligase complexes and as an atypical serine/threonine-protein kinase, playing key roles in various processes such as cell cycle, telomerase regulation and histone modification. Probable substrate-specific adapter of a DCX (DDB1-CUL4-X-box) E3 ubiquitin-protein ligase complex, named CUL4A-RBX1-DDB1-DCAF1/VP RBP complex, which mediates ubiquitination and proteasome-de



pendent degradation of proteins such as NF2. Involved in the tu rnover of methylated proteins: recognizes and binds methylated proteins via its chromo domain, leading to ubiquitination of targ et proteins by the RBX1-DDB1-DCAF1/VPRBP complex (PubMed: 23063525). The CUL4A-RBX1-DDB1-DCAF1/VPRBP complex is als o involved in B-cell development: DCAF1 is recruited by RAG1 t o ubiquitinate proteins, leading to limit error-prone repair durin g V(D)J recombination. Also part of the EDVP complex, an E3 lig ase complex that mediates ubiquitination of proteins such as TE RT, leading to TERT degradation and telomerase inhibition (Pub Med:23362280). Also acts as an atypical serine/threonine-protein kinase that specifically mediates phosphorylation of 'Thr-120' of histone H2A (H2AT120ph) in a nucleosomal context, thereby rep ressing transcription. H2AT120ph is present in the regulatory reg ion of many tumor suppresor genes, down-regulates their transc ription and is present at high level in a number of tumors (Pub Med:24140421). Involved in JNK-mediated apoptosis during cell competition process via its interaction with LLGL1 and LLGL2 (Pu bMed:20644714).