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# Datasheet for ABIN6259637 anti-CCAR2 antibody (Internal Region)

2 Images



### Overview

Quantity:	100 µL
Target:	CCAR2
Binding Specificity:	Internal Region
Reactivity:	Human
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This CCAR2 antibody is un-conjugated
Application:	Western Blotting (WB), ELISA, Immunofluorescence (IF), Immunocytochemistry (ICC)

## Product Details

Immunogen:	A synthesized peptide derived from human DBC1, corresponding to a region within the internal amino acids.
Isotype:	lgG
Specificity:	DBC1 Antibody detects endogenous levels of total DBC1.
Predicted Reactivity:	Horse
Purification:	The antiserum was purified by peptide affinity chromatography using SulfoLink <sup>TM</sup> Coupling Resin (Thermo Fisher Scientific).
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Target Details
Target: CCAR2

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Target Details				
Alternative Name:	CCAR2 (CCAR2 Products)			
Background:	Description: Core component of the DBIRD complex, a multiprotein complex that acts at the			
	interface between core mRNP particles and RNA polymerase II (RNAPII) and integrates			
	transcript elongation with the regulation of alternative splicing: the DBIRD complex affects local			
	transcript elongation rates and alternative splicing of a large set of exons embedded in (A + T)-			
	rich DNA regions. Inhibits SIRT1 deacetylase activity leading to increasing levels of p53/TP53			
	acetylation and p53-mediated apoptosis. Inhibits SUV39H1 methyltransferase activity. As part			
	of a histone H3-specific methyltransferase complex may mediate ligand-dependent			
	transcriptional activation by nuclear hormone receptors. Plays a critical role in maintaining			
	genomic stability and cellular integrity following UV-induced genotoxic stress. Regulates the			
	circadian expression of the core clock components NR1D1 and ARNTL/BMAL1. Enhances the			
	transcriptional repressor activity of NR1D1 through stabilization of NR1D1 protein levels by			
	preventing its ubiquitination and subsequent degradation (PubMed:18235501,			
	PubMed:18235502, PubMed:19131338, PubMed:19218236, PubMed:22446626,			
	PubMed:23352644, PubMed:23398316). Represses the ligand-dependent transcriptional			
	activation function of ESR2 (PubMed:20074560). Acts as a regulator of PCK1 expression and			
	gluconeogenesis by a mechanism that involves, at least in part, both NR1D1 and SIRT1			
	(PubMed:24415752). Negatively regulates the deacetylase activity of HDAC3 and can alter its			
	subcellular localization (PubMed:21030595). Positively regulates the beta-catenin pathway			
	(canonical Wnt signaling pathway) and is required for MCC-mediated repression of the beta-			
	catenin pathway (PubMed:24824780). Represses ligand-dependent transcriptional activation			
	function of NR1H2 and NR1H3 and inhibits the interaction of SIRT1 with NR1H3			
	(PubMed:25661920). Plays an important role in tumor suppression through p53/TP53			
	regulation, stabilizes p53/TP53 by affecting its interaction with ubiquitin ligase MDM2			
	(PubMed:25732823). Represses the transcriptional activator activity of BRCA1			
	(PubMed:20160719). Inhibits SIRT1 in a CHEK2 and PSEM3-dependent manner and inhibits the			
	activity of CHEK2 in vitro (PubMed:25361978).			
	Gene: CCAR2			
Molecular Weight:	102 kDa			
Gene ID:	57805			
UniProt:	Q8N163			

# Application Details

Application Notes:

WB 1:500-1:1000, IF/ICC 1:100-1:500, ELISA(peptide) 1:20000-1:40000

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Application Details		

#### Restrictions:

For Research Use only

## Handling

Format:	Liquid
Concentration:	1 mg/mL
Buffer:	Rabbit IgG in phosphate buffered saline , pH 7.4, 150 mM NaCl, 0.02 % sodium azide and 50 %
	giycerol.
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which
	should be handled by trained staff only.
Storage:	-20 °C
Storage Comment:	Store at -20 °C. Stable for 12 months from date of receipt.
Expiry Date:	12 months

### Images



### Immunofluorescence (fixed cells)

**Image 1.** ABIN6274424 staining Hela by IF/ICC. The sample were fixed with PFA and permeabilized in 0.1% Triton X-100,then blocked in 10% serum for 45 minutes at 25°C. The primary antibody was diluted at 1/200 and incubated with the sample for 1 hour at 37°C. An Alexa Fluor 594 conjugated goat anti-rabbit IgG (H+L) Ab, diluted at 1/600, was used as the secondary antibody.

### Images



#### Western Blotting

**Image 2.** Western blot analysis of extracts from Jurkat cells, using KIAA1967 antibody.

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