

## Datasheet for ABIN966088 anti-ERCC2 antibody (C-Term)

Publication



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## Overview

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Quantity:	0.1 mg
Target:	ERCC2
Binding Specificity:	C-Term
Reactivity:	Human, Mouse
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This ERCC2 antibody is un-conjugated
Application:	Immunohistochemistry (IHC)
Product Details	
Immunogen:	Polyclonal antibody produced in rabbits immunizing with a synthetic peptide corresponding to C-terminal residues of human ERCC2(TFIIH basal transcription factor complex helicase subunit)
Purification:	Purified by antigen-specific affinity chromatography.
Target Details	
Target:	ERCC2
Alternative Name:	ERCC2 (ERCC2 Products)
Background:	ERCC2(TFIIH basal transcription factor complex helicase subunit) is an ATP-dependent 5'-3'

DNA helicase, component of the core-TFIIH basal transcription factor. ERCC2 is involved in

nucleotide excision repair (NER) of DNA by opening DNA around the damage, and in RNA

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	transcription by RNA polymerase II by anchoring the CDK-activating kinase (CAK) complex, composed of CDK7, cyclin H and MAT1, to the core-TFIIH complex. ERCC2 might also have a role in aging process and could play a causative role in the generation of skin cancers. One of the six subunits forming the core-TFIIH basal transcription factor. The interaction with p44 results in the stimulation of the 5'>3' helicase activity. Defects in ERCC2 are the cause of xeroderma pigmentosum complementation group D (XP-D), xeroderma pigmentosum group D
	trichothiodystrophy (TTD) and COFS syndrome. ERCC2 belongs to the helicase family and
	RAD3/XPD subfamily.
Pathways:	DNA Damage Repair
Application Details	
Application Notes:	ELISA, Western blotting: 1µg/ml for 2hrs.
Restrictions:	For Research Use only
Handling	
Format:	Liquid
Buffer:	This antibody is stored in PBS, 50% glycerol
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
Publications	
Product cited in:	Coin, Marinoni, Rodolfo, Fribourg, Pedrini, Egly: "Mutations in the XPD helicase gene result in XP
	and TTD phenotypes, preventing interaction between XPD and the p44 subunit of TFIIH." in:
	Nature genetics, Vol. 20, Issue 2, pp. 184-8, (1998) (PubMed).