

**ERCC4 Antibody (Center) Blocking peptide**  
**Synthetic peptide**  
**Catalog # BP5681c****Specification**

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**ERCC4 Antibody (Center) Blocking peptide - Product Information**

Primary Accession [O92889](#)  
Other Accession [NP\\_005227.1](#)

**ERCC4 Antibody (Center) Blocking peptide - Additional Information**

**Gene ID** 2072

**Other Names**

DNA repair endonuclease XPF, 31--, DNA excision repair protein ERCC-4, DNA repair protein complementing XP-F cells, Xeroderma pigmentosum group F-complementing protein, ERCC4, ERCC11, XPF

**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.

**ERCC4 Antibody (Center) Blocking peptide - Protein Information**

**Name** ERCC4 {ECO:0000303|PubMed:8887684, ECO:0000312|HGNC:HGNC:3436}

**Function**

Catalytic component of a structure-specific DNA repair endonuclease responsible for the 5-prime incision during DNA repair, and which is essential for nucleotide excision repair (NER) and interstrand cross-link (ICL) repair.

**Cellular Location**

Nucleus. Chromosome. Note=Localizes to sites of DNA damage

**ERCC4 Antibody (Center) Blocking peptide - Protocols**

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

- [Blocking Peptides](#)

**ERCC4 Antibody (Center) Blocking peptide - Images**

**ERCC4 Antibody (Center) Blocking peptide - Background**

The protein encoded by this gene forms a complex with ERCC1 and is involved in the 5' incision made during nucleotide excision repair. This complex is a structure specific DNA repair endonuclease that interacts with EME1. Defects in this gene are a cause of xeroderma pigmentosum complementation group F (XP-F), or xeroderma pigmentosum VI (XP6).

**ERCC4 Antibody (Center) Blocking peptide - References**

Sijbers, A.M., et al. Cell 86(5):811-822(1996) Park, C.H., et al. J. Biol. Chem. 270(39):22657-22660(1995) Park, C.H., et al. Proc. Natl. Acad. Sci. U.S.A. 91(11):5017-5021(1994) Liu, P., et al. Mutagenesis 8(3):199-205(1993)