

## ERCC4 Polyclonal Antibody

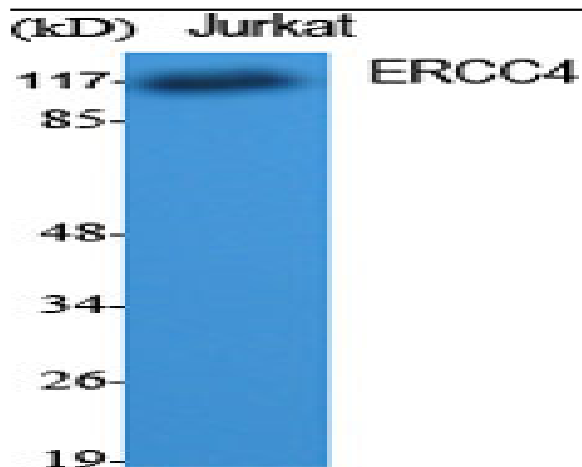
<b>Catalog No :</b>	YT1615
<b>Reactivity :</b>	Human;Mouse
<b>Applications :</b>	WB;ELISA
<b>Target :</b>	ERCC4
<b>Fields :</b>	>>Nucleotide excision repair;>>Fanconi anemia pathway
<b>Gene Name :</b>	ERCC4
<b>Protein Name :</b>	DNA repair endonuclease XPF
<b>Human Gene Id :</b>	2072
<b>Human Swiss Prot No :</b>	Q92889
<b>Mouse Gene Id :</b>	50505
<b>Mouse Swiss Prot No :</b>	Q9QZD4
<b>Immunogen :</b>	The antiserum was produced against synthesized peptide derived from human XPF. AA range:801-850
<b>Specificity :</b>	ERCC4 Polyclonal Antibody detects endogenous levels of ERCC4 protein.
<b>Formulation :</b>	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
<b>Source :</b>	Polyclonal, Rabbit,IgG
<b>Dilution :</b>	WB 1:500 - 1:2000. ELISA: 1:10000. Not yet tested in other applications.
<b>Purification :</b>	The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.
<b>Concentration :</b>	1 mg/ml

---

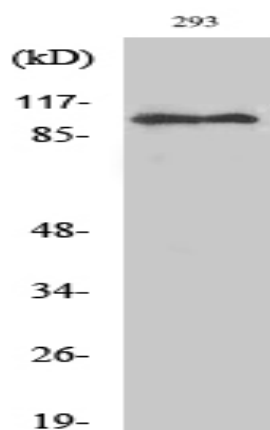
<b>Storage Stability :</b>	-15°C to -25°C/1 year(Do not lower than -25°C)
<b>Observed Band :</b>	103kD
<b>Cell Pathway :</b>	Nucleotide excision repair;
<b>Background :</b>	The protein encoded by this gene forms a complex with ERCC1 and is involved in the 5' incision made during nucleotide excision repair. 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).[provided by RefSeq, Mar 2009],
<b>Function :</b>	cofactor:Magnesium.,disease:Defects in ERCC4 are a cause of XFE progeroid syndrome [MIM:610965]. This syndrome is illustrated by one patient who presented with dwarfism, cachexia and microcephaly.,disease:Defects in ERCC4 are the cause of xeroderma pigmentosum complementation group F (XP-F) [MIM:278760]; also known as xeroderma pigmentosum VI (XP6). XP-F is an autosomal recessive disease characterized by hypersensitivity of the skin to sunlight followed by high incidence of skin cancer and frequent neurologic abnormalities.,function:Structure-specific DNA repair endonuclease responsible for the 5-prime incision during DNA repair. Involved in homologous recombination that assists in removing interstrand cross-link.,similarity:Belongs to the XPF family.,subunit:Heterodimer composed of ERCC1 and XPF/ERCC4. Interacts with EME1.,
<b>Subcellular Location :</b>	Nucleus . Chromosome . Localizes to sites of DNA damage. .
<b>Expression :</b>	Epithelium,Fibroblast,
<b>Tag :</b>	hot
<b>Sort :</b>	5724
<b>No4 :</b>	1
<b>Host :</b>	Rabbit
<b>Modifications :</b>	Unmodified

---

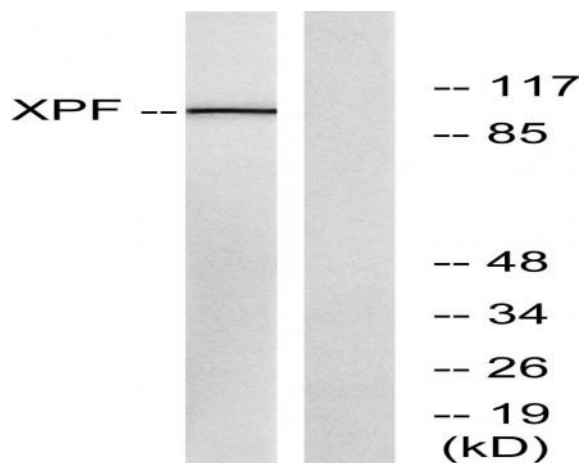
## Products Images



Western Blot analysis of various cells using ERCC4 Polyclonal Antibody cells nucleus extracted by Minute TM Cytoplasmic and Nuclear Fractionation kit (SC-003, Inventbiotech, MN, USA).



Western Blot analysis of 293 cells using ERCC4 Polyclonal Antibody cells nucleus extracted by Minute TM Cytoplasmic and Nuclear Fractionation kit (SC-003, Inventbiotech, MN, USA).



Western blot analysis of lysates from 293 cells, using XPF Antibody. The lane on the right is blocked with the synthesized peptide.