

## Product datasheet for **RC402002**

### **XPD (ERCC2) (NM\_000400) Human Mutant ORF Clone**

#### **Product data:**

<b>Product Type:</b>	Mutant ORF Clones
<b>Product Name:</b>	XPD (ERCC2) (NM_000400) Human Mutant ORF Clone
<b>Mutation Description:</b>	D312N
<b>Affected Codon#:</b>	312
<b>Affected NT#:</b>	934
<b>Nucleotide Mutation:</b>	ERCC2 Mutant (D312N), Myc-DDK-tagged ORF clone of Homo sapiens excision repair cross-complementing rodent repair deficiency, complementation group 2 (ERCC2), transcript variant 1 as transfection-ready DNA
<b>Effect:</b>	Inresed response o UV, ssoiion wih
<b>Symbol:</b>	ERCC2
<b>Synonyms:</b>	COFS2; EM9; TFIH; TTD; TTD1; XPD
<b>E. coli Selection:</b>	Kanamycin (25 ug/mL)
<b>Mammalian Cell Selection:</b>	Neomycin
<b>Vector:</b>	pCMV6-Entry (PS100001)
<b>Tag:</b>	Myc-DDK
<b>ACCN:</b>	NM_000400
<b>ORF Size:</b>	2280 bp
<b>Restriction Sites:</b>	Sgfl-Mlul



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ORF Nucleotide  
Sequence:

>RC402002 representing NM\_000400  
 Red=Cloning site Blue=ORF Green=Tags(s)

TTTTGTAATACGACTCACTATAGGGCGGCCGGGAATTCGTCGACTGGATCCGGTACCGAGGAGATCTGCC  
 GCC**CGGATCGCC**

ATGAAGCTCAACGTGGACGGGCTCCTGGTCTACTTCCCGTACGACTACATCTACCCCGAGCAGTTCTCCT  
 ACATGCGGGAGCTCAAACGCACGCTGGACGCCAAGGGTATGGAGTCTGGAGATGCCCTCAGGCACCGG  
 GAAGACAGTATCCCTGTTGGCCCTGATCATGGCATACAGAGAGCATATCCGCTGGAGGTGACCAAACCT  
 ATCTACTGCTCAAGAACTGTGCCAGAGATTGAGAAGGTGATTGAAGAGCTTCGAAAGTTGCTCAACTTCT  
 ATGAGAAGCAGGAGGGCGAGAAGCTGCCGTTTCTGGGACTGGCTCTGAGCTCCCGCAAAAATTGTGTAT  
 TCACCTGAGGTGACACCCCTGCGCTTTGGGAAGGACGTCGATGGGAAATGCCACAGCCTCACAGCCTCC  
 TATGTGCGGGCGCAGTACCAGCATGACACCAGCCTGCCCACTGCCGATTCTATGAGGAATTTGATGCC  
 ATGGGCGTGAGGTGCCCTCCCGCTGGCATCTACAACCTGGATGACCTGAAGGCCCTGGGGCGGCCA  
 GGGCTGGTGCCATACTTCTTGCTCGATACTCAATCCTGCATGCCAATGTGGTGGTTTATAGCTACCAC  
 TACCTCCTGGACCCCAAGATTGCAGACCTGGTGTCCAAGGAACTGGCCCGCAAGGCCGTCGTGGTCTTCG  
 ACGAGGCCACAACTTGACAACGTCTGCATCGACTCCATGAGCGTCAACCTCACCCGCCGACCCCTTGA  
 CCGGTGCCAGGGCAACCTGGAGACCTGCAGAAGACGGTCTCAGGATCAAAGAGACAGACGAGCAGCGC  
 CTGCGGGACGAGTACCGGCGTCTGGTGGAGGGGCTGCGGGAGGCCAGCGCCGCCCGGGAGACGGACGCC  
 ACCTGGCCAACCCGCTGCTGCCAACGAAGTCTGCAGGAGGCAGTGCCTGGCTCCATCCGCACGGCCGA  
 GCATTTCTGGGCTTCTGAGGGCGTCTGGAGTACGTGAAGTGGCGGCTGCGTGTGACAGCATGTGGT  
 CAGGAGAGCCCGCCCTTCTGAGCGGCTGGCCAGCGCGTGTGCATCCAGCGCAAGCCCTCAGAT  
 TCTGTGCTGAACGCCTCCGGTCCCTGCTGATACTCTGGAGATCACCGACCTTCTGACTTCTCCCGCT  
 CACCTCCTTGCTAACTTTGCCACCCTTGTGACACCTACGCCAAAGGCTTACCATCATCATCGAGCC  
 TTTGACGACAGAACCCCGACCATTGCCAACCCATCCTGCACTTCAGCTGCATGGACGCCTCGTGGCCA  
 TCAAACCCGATTTGAGCGTTTCCAGTCTGTCATCATCACATCTGGGACACTGTCCCGCTGGACATCTA  
 CCCAAGATCCTGGACTTCCACCCCGTACCATGGCAACCTTACCATGACGCTGGCAGGGTCTGCCTC  
 TGCCCTATGATCATCGGCCGTGGCAATGACCAGGTGGCCATCAGCTCAAATTTGAGACCCGGGAGGATA  
 TTGCTGTGATCCGGAATATGGGAACCTCCTGCTGGAGATGTCGCTGTGGTCCCTGATGGCATCGTGGC  
 CTTCTTACCAGTACCAGTACATGGAGAGACCGTGGCTCCTGGTATGAGCAGGGGATCCTTGAGAAC  
 ATCCAGAGGAACAAGCTGCTCTTTATTGAGACCAGGATGGTGGCGAAACAGTGTGCCCTGGAGAAGT  
 ACCAGGAGGCCTGCGAGAATGGCCGCGGGGCCATCCTGCTGTGAGTGGCCCGGGGCAAAGTGTCCGAGGG  
 AATCGACTTTGTGACCACTACGGGCGGGCCGTCATCATGTTTGGCGTCCCTACGTCTACACACAGAGC  
 CGCATTCTCAAGGCGGGCTGGAATACCTGCGGGACCAGTTCAGATTCGTGAGAATGACTTTCTTACCT  
 TCGATGCCATGCGCCACGCGGCCAGTGTGTGGTTCGGCCATCAGGGGCAAGACGGACTACGGCCTCAT  
 GGTCTTTGCCGACAAGCGGTTTGCCTGGGGACAAGCGGGGAAGTGCCTCCGCTGGATCCAGGAGCAC  
 CTCACAGATGCCAACCTCAACCTGACCGTGGACGAGGGTGTCCAGGTGGCCAAGTACTTCTGCGGCAGA  
 TGGCACAGCCCTTCCACCGGGAGGATCAGCTGGCCTGTCCCTGCTCAGCCTGGAGCAGCTAGAATCAGA  
 GGAGACGCTGAAGAGGATAGAGCAGATTGCTCAGCAGCTC

AG**CGGACCG**ACGCGTACGCGGCCGCTCGAGCAGAACTCATCTCAGAAGAGGATCTGGCAGCAAATGATATCC  
 TGGATTACAAGGATGACGACGA TAAGGTTTAA

**Protein Sequence:** >RC402002 representing NM\_000400  
 Red=Cloning site Green=Tags(s)

MKLNVDGLLVYFPYDIYPEQFSYMRELKRTLDAKGHVLEMPSTGKTVSLALIMAYQRAYPLEVTKL  
 IYCSRTVPEIEKVIEELRKLNFYEKQEGEKL PFLGLALSSRNLCIHPVTPLRFGKDVGKCHSLTAS  
 YVRAQYQHDTSLPHCRFYEEFDAHGREVPLPAGIYNLDDLKALGRRQGWCPYFLARYSILHANVVVSYH  
 YLLDPKIALVSKELARKAVVVFDEAHNIDNVCIDSMSVNLTRRTLDRQCQGNLETLQKTVLRRIKETDEQR  
 LRDEYRRLVEGLREASAARETDAHLANPVL PNEVLQEAVPGSIRTAEHFLGFLRRLLEYVKWRLRVQHVV  
 QESPPAFLSGLAQRVCIQRKPLRFCAERLRSLLHLEITDLADF SPLTLLANFATLVSTYAKGFTIIIEP  
 FDDRTPTIANPILHFSCMDASLAIKPVFERFQSVIITSGLSPLDIYPKILDFHPVTMATFTMTLARVCL  
 CPMIIGRGNDQVAISSKFETREDIAVIRNYGNLLEMSAVVPDGIVAFFTSYQYMESTVASWYEQGILEN  
 IQRNKLLFIETQDGAETSVALEKYQEACENGRGAILLSVARGKVSEGIDFVHHYGRAVIMFGVPYVYTQS  
 RILKARLEYLRDQFQIRENFLTFDAMRHAAQCVGRAIRGKTDYGLMVFADKRFARGDKRGLPRWIQEH  
 LTDANLNLTVDEGVQVAKYFLRQMAQPFHREDQLGLSLLSLEQLESEETLKRIEQIAQQQL

SGPTRRRLEQKLI SEEDLAANDILDYKDDDDKV

**Restriction Sites:**

SgfI-MluI

**Cloning Scheme:**



**OTI Disclaimer:**

The molecular sequence of this clone aligns with the gene accession number as a point of reference only. However, individual transcript sequences of the same gene can differ through naturally occurring variations (e.g. polymorphisms), each with its own valid existence. This clone is substantially in agreement with the reference, but a complete review of all prevailing variants is recommended prior to use. [More info](#)

**OTI Annotation:**

This clone was engineered to express the complete ORF with an expression tag. Expression varies depending on the nature of the gene.

<b>Components:</b>	The ORF clone is ion-exchange column purified and shipped in a 2D barcoded Matrix tube containing 10ug of transfection-ready, dried plasmid DNA (reconstitute with 100 ul of water).
<b>RefSeq:</b>	<a href="#">NP_000391</a>
<b>RefSeq Size:</b>	2280 bp
<b>RefSeq ORF:</b>	2283 bp
<b>Locus ID:</b>	2068
<b>Cytogenetics:</b>	19q13.32
<b>Domains:</b>	DEXDc2, HELICc2
<b>Protein Families:</b>	Druggable Genome, Transcription Factors
<b>Protein Pathways:</b>	Nucleotide excision repair
<b>MW:</b>	83.6 kDa
<b>Gene Summary:</b>	<p>The nucleotide excision repair pathway is a mechanism to repair damage to DNA. The protein encoded by this gene is involved in transcription-coupled nucleotide excision repair and is an integral member of the basal transcription factor BTF2/TFIIH complex. The gene product has ATP-dependent DNA helicase activity and belongs to the RAD3/XPD subfamily of helicases. Defects in this gene can result in three different disorders, the cancer-prone syndrome xeroderma pigmentosum complementation group D, trichothiodystrophy, and Cockayne syndrome. Alternatively spliced transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, Aug 2008]</p>