

Product datasheet for **RC400086**

p53 (TP53) (NM_000546) Human Mutant ORF Clone

Product data:

Product Type:	Mutant ORF Clones
Product Name:	p53 (TP53) (NM_000546) Human Mutant ORF Clone
Mutation Description:	R213X
Affected Codon#:	213
Affected NT#:	c.637
Nucleotide Mutation:	TP53 mutant (R213X), Myc-DDK-tagged ORF clone of Homo sapiens tumor protein p53 (TP53), transcript variant 1 as transfection-ready DNA
Effect:	nonsense
Symbol:	p53
Synonyms:	BCC7; BMFS5; LFS1; P53; TRP53
E. coli Selection:	Kanamycin (25 ug/mL)
Mammalian Cell Selection:	Neomycin
Vector:	pCMV6-Entry (PS100001)
Tag:	Myc-DDK
ACCN:	NM_000546
ORF Size:	1179 bp
Restriction Sites:	Sgfl-Mlul



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ORF Nucleotide Sequence:

>RC400086 representing NM_000546
 Red=Cloning site Blue=ORF Green=Tags(s)

TTTTGTAATACGACTCACTATAGGGCGGCCGGGAATTCGTCGACTGGATCCGGTACCGAGGAGATCTGCC
 GCC**GCGATCGCC**

ATGGAGGAGCCGAGTCAGATCCTAGCGTCGAGCCCCCTCTGAGTCAGGAAACATTTTCAGACCTATGGA
 AACTACTTCTGAAAACAACGTTCTGTCCCTTGCCGTCCTCAAGCAATGGATGATTTGATGCTGTCCCC
 GGACGATATTGAACAATGGTTCCTGAAGACCCAGGTCCAGATGAAGCTCCAGAAATGCCAGAGGCTGCT
 CCCCCGTGGCCCTGCACCAGCAGCTCTACCCGGCGGCCCTGCACCAGCCCCCTCCTGGCCCTGT
 CATCTTCTGTCCCTCCAGAAAACCTACCAGGGCAGCTACGTTTCCGTCCTGGGCTTCTTGCACTCTGG
 GACAGCCAAGTCTGTGACTTGCACGTAATCCCCTGCCCTCAACAAGATGTTTTGCCAACTGGCCAAGACC
 TGCCCTGTGCAGCTGTGGTTGATTCCACACCCCGCCCGCACCCGCTCCGCGCCATGCCCATCTACA
 AGCAGTCACAGCACATGACGGAGGTTGTGAGGCGCTGCCCCACCATGAGCGCTGCTCAGATAGCGATGG
 TCTGGCCCTCCTCAGCATCTTATCCGAGTGAAGGAAATTTGCGTGTGGAGATTTGGATGACAGAAAC
 ACTTTT

**ACGCGTACGCGGCCGCTCGAGCAGAAACTCATCTCAGAAGAGGATCTGGCAGCAAATGATATCCTGGATT
 ACAAGGATGACGACGATAAGGTTTAA**

Protein Sequence:

>RC400086 representing NM_000546
 Red=Cloning site Green=Tags(s)

MEEPQSDPSVEPPLSQETFSDLWKLLPENNVLSPLPSQAMDDLMLSPDDIEQWFTEDPGPDEAPRMPEAA
 PPVAPAPAAPTAAAPAPAPSWPLSSSVPSQKTYQGSYGFRLGFLHSGTAKSVTCTYSPALNKMFCQLAKT
 CPVQLWVDSTPPPGRVRAAIYKQSQHMTEVRRRCPPHHERCSDSDGLAPPQHLIRVEGNLRVEYLDRN
 TF

TRTRPLEQKLISEEDLAANDILDYKDDDDKV

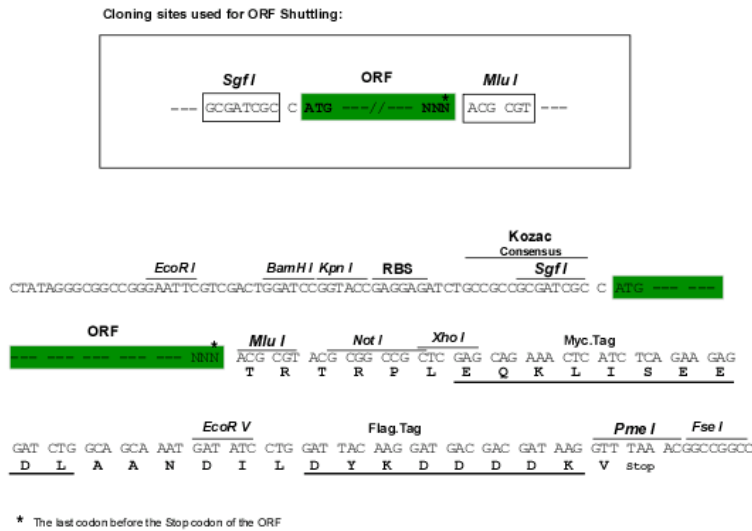
Chromatograms:

/chromatograms/ja1129_g07.zip

Restriction Sites:

Sgfl-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.
Components:	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).
RefSeq:	NP_000537
RefSeq Size:	2629 bp
RefSeq ORF:	1182 bp
Locus ID:	7157
Cytogenetics:	17p13.1
Domains:	P53
Protein Families:	Druggable Genome, Stem cell - Pluripotency, Transcription Factors
Protein Pathways:	Amyotrophic lateral sclerosis (ALS), Apoptosis, Basal cell carcinoma, Bladder cancer, Cell cycle, Chronic myeloid leukemia, Colorectal cancer, Endometrial cancer, Glioma, Huntington's disease, MAPK signaling pathway, Melanoma, Neurotrophin signaling pathway, Non-small cell lung cancer, p53 signaling pathway, Pancreatic cancer, Pathways in cancer, Prostate cancer, Small cell lung cancer, Thyroid cancer, Wnt signaling pathway
MW:	23.3 kDa
Gene Summary:	This gene encodes a tumor suppressor protein containing transcriptional activation, DNA binding, and oligomerization domains. The encoded protein responds to diverse cellular stresses to regulate expression of target genes, thereby inducing cell cycle arrest, apoptosis, senescence, DNA repair, or changes in metabolism. Mutations in this gene are associated with a variety of human cancers, including hereditary cancers such as Li-Fraumeni syndrome. Alternative splicing of this gene and the use of alternate promoters result in multiple transcript variants and isoforms. Additional isoforms have also been shown to result from the use of alternate translation initiation codons from identical transcript variants (PMIDs: 12032546, 20937277). [provided by RefSeq, Dec 2016]