

Division of Signal Transduction Therapy

Standard Operation Procedure

Preparation of Parkin [R455A]

<u>Enzyme description:-</u>	Parkin 1-465 [R455A]
<u>Clone number:-</u>	DU46030
<u>Source:-</u>	BL21 recombinant
<u>Tag:-</u>	cleaved from N-terminal His-SUMO1 tag
<u>Purification method:-</u>	Ni ⁺⁺ -Sephareose, SENP-treatment, Ni ⁺⁺ -depletion, SEC
<u>Expression level:-</u>	0.2mg/L
<u>Calculated molecular mass:-</u>	
Monoisotopic	51521 Da
Average Mass	51554 Da
[cysteines reduced, methionines have not been oxidised]	
<u>Theoretical pI:-</u>	7.00
<u>Purity:-</u>	90%
<u>Enzyme storage buffer:-</u>	
50mM HEPES pH 8.2, 150mM NaCl, 20% glycerol, 1mM TCEP	
<u>Storage temperature:-</u>	-80°C
<u>Assay:-</u>	

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Clone Data Sheet

Parkin [R455A]

<u>Protein</u>	Parkin 1-465 [R455A]
<u>Synonyms</u>	PARK2, PRKN2
<u>Clone Number</u>	DU46030
<u>Species</u>	Human
<u>Accession Number</u>	Protein: O60260 Gene: NM_004562.2
<u>Tags</u>	N-terminal His ₆ -SUMO1-tag (cleaved)
Aminoacid sequence of the expressed protein	MGHHHHHSDQEAKPSTEDLGDKKEGEYIKLKVIGQDSSEIHFVKVMTTH LKKLKESYCRQGVPMNSLRFLFEGQRIADNHTPKELGMEEEDVIEVYQE QTGGMI VVFRFNSSHGFPVEVDS DTSIFQLKEVVAKRQGV PADQLRVIFA GKELRNDWTVQNC DLDQQSIVHIVQRPWRKQEMNATGGDDPRNAAGGCE REPQSLTRVDLSSSVLPGDSVGLAVILHTDSRKDSPPAGSPAGRSIYNSF YVYCKGPCQRVQPGKLRVQCSTCRQATLTLTQGPSCWDDVLI PNRMSGEC QSPHCPGTSAEFFFKCGAHPTSDKETSVALHLIATNSRNITCITCTD VRS PVLV FQCNSRHVICLDCFHLYCVTRLNDRQFVHDPQLGYS LPCVAGCPNS LIKELHHFRILGEEQYNRYQQYGAEECVLQMGVLCPRPGCGAGLLPEPD QRKVTCEGGNGLGCGFAFCRECKEAYHEGECSAVFEASGTTTQAYR VDER AAEQARWEAASKETIKKTTKPCPRCHVPVEKNGGCMHMKCPQPQCRLEWC WNCGCEWNAVCMGDHWF DV
Native sequence	in bold
Protease cleavage	SEN1 treatment after GG of SUMO1
Cloning sites	BamH1 / Not 1 (destroyed)
DNA sequence of the insert	CCATGGGTCATCATCACCATCACCATTCTGACCAGGAGGCAAAACCTTCA ACTGAGGACTTGGGGGATAAGAAGGAAGGTGAATATATTAACCTCAAAGT CATTGGACAGGATAGCAGTGAGATTCACCTTCAAAGTGAAAATGACAACAC ATCTCAAGAACTCAAAGAATCATACTGTCAAAGACAGGGTGTTC CAATG AACTCACTCAGGTTTCTCTTTGAGGGTCAGAGAATTGCTGATAATCATA C TCCAAAAGAACTGGGAATGGAGGAAGAAGATGTGATTGAAGTTTATCAGG AACAAACGGGGGAATGATAGTGT TTTGT CAGGTTCAACTCCAGCCATGGT TTCCCAGTGGAGGTCGATTCTGACACCAGCATCTTCCAGCTCAAGGAGGT GGTTGCTAAGCGACAGGGGGTTCCGGCTGACCAGTTGCGTGTGATTTTCG CAGGGAAGGAGCTGAGGAATGACTGGACTGTGCAGAATTGTGACCTGGAT CAGCAGAGCATTGTTACATTGTGCAGAGACCGTGGAGAAAAGGTCAAGA AATGAATGCAACTGGAGGCGACGACCC CAGAAACGCGGCGGGAGGCTGTG AGCGGGAGCCCAGAGCTTGACTCGGGTGGACCTCAGCAGCTCAGTCCTC CCAGGAGACTCTGTGGGGCTGGCTGT CATTCTGCACACTGACAGCAGGAA GGACTCACCACCAGCTGGAAGTCCAGCAGGTAGATCAATCTACAACAGCT TTTATGTGTATTGCAAAGGCCCTGTCAAAGAGTGCAGCCGGGAAAACCTC AGGTTACAGTGCAGCACCTGCAGGCAGGCAACGCTCACCTTGACCCAGG TCCATCTTGCTGGGATGATGTTTAAATTCCAAACCGGATGAGTGGTGAAT GCCAATCCCACACTGCCCTGGGACTAGTGCAGAATTTTTCTTTAAATGT GGAGCACACCCACCTCTGACAAGGAAACATCAGTAGCTTTGCACCTGAT CGCAACAAATAGTCGGAACATCACTTGCATTACGTGCACAGACGTCAGGA GCCCCGTCCTGGTTTTCCAGTGCAACTCCC GCCACGTGATTGCTTAGAC TGTTTCCACTTATACTGTGTGACAAGACTCAATGATCGGCAGTTTGTTC A

CGACCCTCAACTTGGCTACTCCCTGCCTTGTGTGGCTGGCTGTCCCAACT
CCTTGATTAAAGAGCTCCATCACTTCAGGATTCTGGGAGAAGAGCAGTAC
AACCGGTACCAGCAGTATGGTGCAGAGGAGTGTGTCTGCAGATGGGGGG
CGTGTTATGCCCCGCCCTGGCTGTGGAGCGGGGCTGCTGCCGGAGCCTG
ACCAGAGGAAAGTCACCTGCGAAGGGGGCAATGGCCTGGGCTGTGGGTTT
GCCTTCTGCCGGGAATGTAAAGAAGCGTACCATGAAGGGGAGTGCAGTGC
CGTATTTGAAGCCTCAGGAACAACACTCAGGCCTACAGAGTCGATGAAA
GAGCCGCCGAGCAGGCTCGTTGGGAAGCAGCCTCCAAAGAAACCATCAAG
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AGGCTGCATGCACATGAAGTGTCCGCAGCCCCAGTGCAGGCTCGAGTGGT
GCTGGAAGTGTGGCTGCGAGTGGAACGCCGTCTGCATGGGGGACCACTGG
TTCGACGTGTAGGCGGCCGC