

Division of Signal Transduction Therapy

Standard Operation Procedure

Preparation of Parkin

<u>Enzyme description:-</u>	Parkin 1-465 (full length)
<u>Clone number:-</u>	DU39835
<u>Source:-</u>	Recombinant
<u>Tag:-</u>	cleaved from N-terminal His ₆ -SUMO-1
<u>Purification method:-</u>	Ni ⁺⁺ -Sepharose, SEC
<u>Expression level:-</u>	1 mg/L
<u>Calculated molecular mass:-</u>	
Monoisotopic	51590 Da
Average Mass	51623 Da
[cysteines reduced, methionines have not been oxidised]	
<u>Theoretical pI:-</u>	7.21
<u>Purity:-</u>	95 %
<u>Enzyme storage buffer:-</u>	
50 mM HEPES pH 8.2, 20% glycerol, 150mM NaCl, 0.5mM TCEP, 0.03% Brij35	
<u>Storage temperature:-</u>	-80°C
<u>Assay:-</u>	

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Clone Data Sheet

Protein name Parkin S65A

<u>Protein</u>	Parkin 1 - 465 (Ser65 Ala) (full length)
<u>Synonyms</u>	PARK2, PRKN
<u>Clone Number</u>	DU23314
<u>Species</u>	Human
<u>Accession Number</u>	Protein: O60260
<u>Tags</u>	N-terminal His, followed by SUMO-1 to improve solubility
Aminoacid sequence of the expressed protein .	MGHHHHHSDQEAKPSTEDLGDKKEGEYIKLKVIGQDSSEIHFVKVMTTH LKKLKESYCORQGVPMNSLRFLFEGQRIADNHTPKELGMEEDVIEVYQE QTGGMIVFVRFNSSHGFPVEVDSDTSTIFQLKEVVAKRQGVADQLRVIFA GKELRNDWTVQNCDLDOQAIVHIVQRPWRKGQEMNATGGDDPRNAAGGCE REPOSLTRVDLSSSVLPGDSVGLAVILHTDSRKDSPPAGSPAGRSIYNSF YVYCKGPCQORVQPGKLRVQCSQATLTLTQGPSCWDDVLI PNRMSGEC QSPHCPGTSAEFFFKCGAHPTSDKETSVALHLIATNSRNITCITCTDVRS PVLVQFCNSRHVICLDCFHLYCVTRLNDRQFVHDPQLGYSLPCVAGCPNS LIKELHHFRILGEEQYNRYQQYGAEECVLQMGVLCPRPGCGAGLLPEPD QRKVTCEGGNGLGCGFAFCRECKEAYHEGECSAVFEASGTTTQAYRVDER AAEQARWEAASKETIKKTTKPCPRCHVPVEKNGGCMHMKCPQPQCRLEWC WNCGCEWNRVCMGDHWFVDV
SUMO-1 in grey, is removed during purification by SENP1	
The final product, in bold	
Native sequence	in bold
Protease cleavage	SENP1 protease site underlined
Cloning sites	Complex cloning, please inquire.

DNA sequence of cassette

ATGGGTCATCATCACCATCACCATTCTGACCAGGAGGCAAACCTTCAACT
GAGGACTTGGGGGATAAGAAGGAAGGTGAATATATTAAGTCAAAGTCATT
GGACAGGATAGCAGTGAGATTCACTTCAAAGTGAAAATGACAACACATCTC
AAGAACTCAAAGAATCATACTGTCAAAGACAGGGTGTCCAATGAACTCA
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