

*Division of Signal Transduction Therapy*

**Standard Operation Procedure**

**Preparation of GST-USP21 [196-565]**

<b><u>Enzyme description:-</u></b>	GST-USP21
<b><u>Clone number:-</u></b>	DU22385
<b><u>Source:-</u></b>	BL21 Recombinant
<b><u>Tag:-</u></b>	N-terminal GST
<b><u>Purification method:-</u></b>	GSH sepharose
<b><u>Expression level:-</u></b>	1.5 mg/L

**Calculated molecular mass:-**

Monoisotopic	67,325 Da
Average Mass	67,366 Da
[cysteines reduced, methionines have not been oxidised]	

**Theoretical pI:-** 8.6

**Purity:-** 80%

**Enzyme storage buffer:-**

50 mM HEPES pH 7.5, 10% glycerol, 150mM NaCl, 1mM DTT

**Storage temperature:-** -80°C

**Assay:-**

Ub-Rho110-Gly cleavage assay monitored by Ex/Em 485/535 nm

**Assay buffer:-**

40 mM Tris pH 7.5, 100 mM NaCl, 5 mM DTT, 0.01% Triton X-100, 0.005% Ovalbumin, 0.5 µM Ub-Rho110-Gly

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**Clone Data Sheet**

**GST-USP21 [196-565]**

<b><u>Protein</u></b>	GST-USP21 [196-565]
<b><u>Synonyms</u></b>	
<b><u>Clone Number</u></b>	DU22385
<b><u>Species</u></b>	Human
<b><u>Accession Number</u></b>	Protein: Q9UK80 DNA: NM_001014443.2
<b><u>Tags</u></b>	N-terminal GST
<b><u>Amino acid sequence of expressed protein</u></b>	<b>MSPILGYWKIKGLVQPTRLLEYLEEKYEEHLYERDEGDKWRNKKFELGLEFP NLPYYIDGDVKLTQSMAIIRYIADKHNMLGGCPKERAEISMLEGAVLDIRYGV SRIAYSKDFETLKVDFLSKLPPEMLKMFEDRLCHKTYLNGDHVTHPDFMLYDAL DVVLYMDPMCLDAFPKLVCFKKRIEAIPOIDKYLKSSKYIAWPLQGWQATFGG GDHPPKSDSDDKMAHHTLLLGSQHVGLRNLGNTCFLNAVLQCLSSTRPLRDFC LRRDFRQEVPGGGRAQELTEAFADVIGALWHPDSCEAVNPTRFRAVFQKYVPS FSGYSQQDAQEFLKLLMERLHLEINRRGRRAPPILANGPVPSPPRRGGALLEE PELSDDDRANLMWKRYLEREDSKIIVDLFVGQLKSCLKCQACGYRSTTFEVFCD LSLPIPKKGFAGGKVSRLRDCFNLFTEKEEELSENAPVCDRCRQKTRSTKKLTV QRFPRILVLHLNRFSAASRSIKKSSVGVDFPLQRLSLGDFASDKAGSPVYQLY ALCNHSGSVHYGHYALCQCQTGWVHYNDSRVSPVSENQVASSEGYVLFYQLM QEPPRCL</b>
<b><u>Native sequence</u></b>	in bold
<b><u>Protease cleavage</u></b>	No cleavage sites in this construct
<b><u>Cloning sites</u></b>	Sal1 / Not1
<b><u>DNA sequence of insert</u></b>	TCTGATGACAAGATGGCTCATCACACTCCTTCTGGGCTCTGGTCATGTTGG CCTTCGAAACCTGGGAAACACGTGCTTCCTGAAATGCTGTGCTGCAGTGTCTGA GCAGCACTCGACCTCTTCGGGACTTCTGTCTGAGAAGGGACTTCCGGCAAGAG GTGCCTGGAGGAGGCCGAGCCCAAGAGCTCACTGAAGCCTTTGCAGATGTGAT TGGTGCCCTCTGGCACCCTGACTCCTGCGAAGCTGTGAATCCTACTCGATTCC GAGCTGTCTTCCAGAAATATGTTCCCTCCTTCTCTGGATACAGCCAGCAGGAT GCCCAAGAGTTCTGAAGCTCCTCATGGAGCGGCTACACCTTCAAATCAACCG CCGAGGCCCGGGCTCCACCGATACTTGCCAAATGGTCCAGTTCCTCTCCAC CCCGCCGAGGAGGGGCTCTGCTAGAAGAACCTGAGTTAAGTGATGATGACCGA GCCAACCTAATGTGGAAACGTTACCTGGAGCGAGAGGACAGCAAGATTGTGGA CCTGTTTGTGGGCCAGTTGAAAAGTTGTCTCAAGTGCCAGGCCTGTGGGTATC GCTCCACGACCTTCGAGGTTTTTTGTGACCTGTCCCTGCCCATCCCCAAGAAA GGATTTGCTGGGGCAAGGTGTCTCTGCGGGATTGTTTCAACCTTTTCACTAA GGAAGAAGAGCTAGAGTCCGAGAAATGCCCCAGTGTGTGACCGATGTCCGGCAGA AAACTCGAAGTACCAAAAAGTTGACAGTACAAAAGATTCCCTCGAATCCTCGTG CTCCATCTGAATCGATTTTTCTGCCCTCCCGAGGCTCCATCAAAAAAAGTTTCAGT AGGTGTAGACTTTTCCACTGCAGCGACTGAGCCTAGGGGACTTTGCCAGTGACA AAGCCGGAAGTCTGTATAACAGCTGTATGCCCTTTGCAACCACTCAGGCAGC GTCCACTATGGCCACTACACAGCCCTGTGCCGGTGCCAGACTGGTTGGCATGT CTACAATGACTCTCGTGTCTCCCTGTGAGTAAAACCAGGTGGCATCCAGCG AGGGCTACGTGCTGTTCTACCAACTGATGCAGGAGCCACCCCGGTGCCTGTGA GCGGCCGC