

*Division of Signal Transduction Therapy*

**Standard Operation Procedure**

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

**Enzyme description:-** His-USP21 [196-565]

**Clone number:-** DU22384

**Source:-** BL21 Recombinant

**Tag:-** N-terminal

**Purification method:-** N-terminal His<sub>6</sub> tag

**Expression level:-** 1.6 mg/L

**Calculated molecular mass:-**

Monoisotopic 42826 Da

Average Mass 42852 Da

[cysteines reduced, methionines have not been oxidised]

**Theoretical pI:-** 9.15

**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**

**His-USP21 [196-565]**

**Protein** His-USP21 [196-565]

**Synonyms**

**Clone Number** DU22384

**Species** Human

**Accession Number** Protein: Q9UK80 DNA: NM\_001014443.2

**Tags** N-terminal

**Amino acid sequence of expressed protein** **MHHHHHHVDSDDKMAHHTLLLLGSGHVGRLNGLNTCFLNAVLQCLSSSTRPLRDF  
CLRRDFRQEVPPGGRAQELTEAFADVIGALWHPDSCAVNPTRFRAVFQKYVP  
SFSGYSQQDAQEFLLKLLMERLHLEINRRGRRAPPILANGPVPSPPRRGGALLE  
EPELSDDDRANLMWKRYLEREDSKI VDLFVGLKSKLKCQACGYRSTTFEVFC  
DLSLP I P K K G F A G G K V S L R D C F N L F T K E E E L E S E N A P V C D R C R Q K T R S T K K L T  
V Q R F P R I L V L H L N R F S A S R G S I K K S S V G V D F P L Q R L S L G D F A S D K A G S P V Y Q L  
Y A L C N H S G S V H Y G H Y T A L C R C Q T G W H V Y N D S R V S P V S E N Q V A S S E G Y V L F Y Q L  
M Q E P P R C L**

**Native sequence** in bold

**Protease cleavage** No protease site in this construct

**Cloning sites** EcoR1/Sal1

**DNA sequence of insert** GTCGACTCTGATGACAAGATGGCTCATCACACACTCCTTCTGGGCTCTGGTCA  
TGTTGGCCTTCGAAACCTGGGAAACACGTGCTTCCTGAATGCTGTGCTGCAGT  
GTCTGAGCAGCACTCGACCTCTTCGGGACTTCTGTCTGAGAAGGGACTTCCGG  
CAAGAGGTGCCTGGAGGAGGCCGAGCCCAAGAGCTCACTGAAGCCTTTGCAGA  
TGTGATTGGTGCCCTCTGGCACCCCTGACTCCTGCGAAGCTGTGAATCCTACTC  
GATTCCGAGCTGTCTTCCAGAAATATGTTCCCTCCTTCTCTGGATACAGCCAG  
CAGGATGCCCAAGAGTTCCTGAAGCTCCTCATGGAGCGGCTACACCTTGAAAT  
CAACCGCCGAGGCCGCGGGCTCCACCGATACTTGCCAATGGTCCAGTTCCTT  
CTCCACCCGCGGAGGAGGGCTCTGCTAGAAGAACCTGAGTTAAGTGATGAT  
GACCGAGCCAACCTAATGTGGAACGTTACCTGGAGCGAGAGGACAGCAAGAT  
TGTGGACCTGTTTGTGGGCCAGTTGAAAAGTTGTCCTCAAGTGCCAGGCCTGTG  
GGTATCGCTCCACGACCTTCGAGGTTTTTTGTGACCTGTCCCTGCCATCCCC  
AAGAAAGGATTTGCTGGGGGCAAGGTGTCTCTGCGGGATTGTTTCAACCTTTT  
CACTAAGGAAGAAGAGCTAGAGTCGGAGAATGCCCCAGTGTGTGACCGATGTC  
GGCAGAAAACCTCGAAGTACCAAAAAGTTGACAGTACAAAGATTCCTCGAATC  
CTCGTGCTCCATCTGAATCGATTTTCTGCCTCCCGAGGCTCCATCAAAAAAAG  
TTCAGTAGGTGTAGACTTTCCACTGCAGCGACTGAGCCTAGGGGACTTTGCCA  
GTGACAAAGCCGGAAGTCTGTATACCAGCTGTATGCCCTTTGCAACCACTCA  
GGCAGCGTCCACTATGGCCACTACACAGCCCTGTGCCGGTGCCAGACTGGTTG  
GCATGTCTACAATGACTCTCGTGTCTCCCCTGTGAGTAAAAACCAGGTGGCAT  
CCAGCGAGGGCTACGTGCTGTTCTACCAACTGATGCAGGAGCCACCCCGGTG  
CTGTGAGCGGCCGC