

## *Division of Signal Transduction Therapy*

### **Standard Operating Procedure**

#### **Preparation of DAZAP T269A [2 - 407]**

**Enzyme description:-** DAZAP T269A [2 - 407]

**Clone number:-** DU 3217

**Source:-** Recombinant

**Expression system:-** *E.coli*

**Tag:-** N-terminal GST

**Purification method:-** GSH Sepharose

**Calculated molecular mass:-**

Monoisotopic 70,001.64 daltons

Average Mass 70,046.34 daltons

[cysteines reduced, methionines have not been oxidised]

**Theoretical pI:-** 6.87

**Purity:-** >80 %

**Enzyme storage buffer:-**

50 mM Tris-HCl pH 7.5, 270 mM Sucrose, 150 mM NaCl, 0.1 mM EGTA,  
0.1 % 2-mercaptoethanol, 0.02 % Brij-35, 1 mM benzamidine, 0.2 mM PMSF

**Storage temperature:-** -70 °C

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### Clone Data Sheet

#### DAZAP T269A [2 - 407]

<b><u>Protein</u></b>	DAZAP T269A [2 - 407]
<b><u>Clone number</u></b>	DU 3217
<b><u>Species</u></b>	Human
<b><u>Accession number</u></b>	NM_018959.3
<b><u>Tags</u></b>	N-terminal GST
<b><u>Bacterially expressed protein</u></b>	<p>MSPILGYWKIKGLVQPTRLLLEYLEEKYEEHLYERDEGDKWRNKKFELG LEFPNLPPYYIDGDVKLTQSMAIIRYIADKHNMLGGCPKERAIEISMLEGA VLDIRYGVSR IAYS KDFETLKVDFLSKLP EMLKMFEDRLCHKTYLNGDH VTHPDFMLYDALDVVLYMDPMCLDAFPKLVCFKKRIEAIPOIDKYLKSS KYIAWPLQGWQATFGGGDHPPKSDLEVL FQGPLGSNNSGADEIGKLFVG <b>GLDWSTTQETLRSYFSQYGEVVDCVIMKDKT TNQSRGFGFVKFKDPNCV</b> <b>GTVLASRPHTLDGRNIDPKPCTPRGMQPERTRPKEGWQKGPRSDNSKSN</b> <b>KIFVGGIPHNCGETELREYFKKFVGVTEVVM IYDAEKQRPRGFGFITFE</b> <b>DEQSV DQAVNMHFHDIMGKKVEVKRAEPRDSKSQAPGQPGASQWGSRVV</b> <b>PNAANGWAGQPPPTWQOGYGPOGMWVPAGQAIGGYGPPPPAGRGAPPPP</b> <b>PFTSYIVSAPPGGFPPPQGFPOGYGAPPQFSFGYGPPPPPPDQFAPPGV</b> <b>PPPPATPGAAPLAFPPPPSQAAPDMSKPPTAQPDPYGYAGYGQDLSG</b> <b>FGQGFSDPSQPPPSYGGPSVPGSGGPPAGGSGFGRGQNHNVQGFHPYRR</b></p>
<b><u>Native sequence</u></b>	<p>Amino acids N2 – R407 (end) of human DAZAP. Residue N232 of the fusion protein is equivalent to N2 of the native enzyme. The GST tag is located at residues 1 – 220. The protein has a T269A mutation. Residue T269 is equivalent to residue A499 of the fusion protein.</p>
<b><u>Protease cleavage</u></b>	PreScission (LEVL FQGP) residues 221 - 228
<b><u>Cloning sites</u></b>	BamH1 sites of pGEX6P-1

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**Nucleotide**  
**Sequence of insert**

ggatccACAACCTCGGGCGCCGACGAGATCGGGAAGCTCTTCGTGGGCGGTCTT  
GACTGGAGCACGACCCAAGAGACTCTGCGCAGCTACTTTTCCCAATATGGAGAA  
GTCGTAGATTGTGTTATCATGAAAGATAAAACCACCAACCAGTCTCGAGGCTTT  
GGGTTTGTCAAATTTAAAGACCCAAACTGTGTGGGGACGGTGCTGGCCAGCAGA  
CCGCACACGCTAGATGGCCGAAACATCGACCCCAAGCCATGCACACCCCGGGGG  
ATGCAGCCGGAGAGAACACGGCCGAAGGAAGGATGGCAGAAAGGACCCAGGAGC  
GATAACAGTAAATCAAATAAGATATTTGTTCGGTGGAAATTCCTCACAAATTGTGGT  
GAGACAGAGCTCAGGGAATACTTCAAGAAGTTCGGAGTGGTCACGGAGGTAGTC  
ATGATCTATGACGCCGAGAAGCAGAGGCCCCCGAGGTTTTGGATTTATTACTTTC  
GAGGACGAACAATCAGTGGACCAGGCTGTCAACATGCATTTTTCACGACATCATG  
GGCAAAAAGTGAAGTTAAACGAGCTGAGCCTCGGGACAGCAAGAGCCAAGCG  
CCGGGACAGCCAGGTGCCAGCCAGTGGGGGAGCCGGGTTGTGCCAACGCTGCC  
AATGGCTGGGCAGGCCAGCCCCCGCCACGTGGCAGCAAGGATATGGCCCGCAA  
GGAATGTGGGTGCCGGCAGGACAGGCGATTGGTGGCTATGGACCCGCCCTGCA  
GGAAGAGGAGCCCCCGCCACCCACCGTTCACCTCCTACATCGTGTCCGCC  
CCTCCTGGAGGCTTTCCCCCTCCCCAGGGCTTCCCTCAGGGCTACGGTGCCCCG  
CCACAGTTCAGTTTTGGCTACGGGCCTCCACCTCCACCGCCAGATCAGTTTGCC  
CCTCCGGGGTTCTCCTCCACCAGCCACTCCCGGGGCAGCACCTCTGGCTTTC  
CCACCGCTCCGTCTCAGGCTGCCCCGGACATGAGCAAGCCCCCGACAGCTCAG  
CCAGACTTCCCCTATGGTCAGTATGCAGGTTACGGGCAGGACTTGAGTGGCTTC  
GGACAGGGCTTCTCAGACCCAGCCAGCAGCCTCCTTCTACGGGGTCCCTCC  
GTGCCAGGGTCGGGGGGCCCCCGCCGGCAGCGGCTTTGGACGAGGGCAG  
AACCACAACGTGCAAGGTTCCACCCCTACCGACGctaggatcc