

## *Division of Signal Transduction Therapy*

### **Standard Operating Procedure**

#### **Preparation of hnRNP A1 [2 - 189]**

<b><u>Protein description:-</u></b>	hnRNP A1 [2 - 189]
<b><u>Clone number:-</u></b>	DU 4003
<b><u>Source:-</u></b>	Recombinant
<b><u>Expression system:-</u></b>	<i>E.coli</i>
<b><u>Tag:-</u></b>	N-terminal GST and HA
<b><u>Purification method:-</u></b>	GSH Sepharose
<b><u>Expression level:-</u></b>	10 mg/L
<b><u>Calculated molecular mass:-</u></b>	49, 552 daltons
<b><u>Purity:-</u></b>	95 %

#### **Enzyme storage buffer:-**

50 mM Tris-HCl pH 7.5, 50 % glycerol, 150 mM NaCl, 0.1 mM EGTA, 0.1 % 2-mercaptoethanol, 0.02 % Brij-35, 0.2 mM PMSF, 1 mM Benzamidine.

**Storage temperature:-** -20 °C

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**CLONE DATA SHEET**

**hnRNP A1 [2 – 189]**

<b><u>Protein</u></b>	hnRNP A1 [2 - 189]
<b><u>Clone number</u></b>	DU 4003
<b><u>Species</u></b>	Human
<b><u>Accession no</u></b>	NM_031157
<b><u>Tags</u></b>	N-terminal GST and HA
<b><u>Bacterially expressed protein</u></b>	<p>MSPILGYWKIKGLVQPTRLLEYLEEKYEHLRYERDEGDKW RNKKFELGLEFPNLPYYIDGDVKLTQSMAIIRYIADKHNML GGCPKERAESMLEGAVLDIRYGVSRIAYSKDFETLKVDFL SKLPEMLKMFEDRLCHKTYLNGDHVTHPDFMLYDALDVVLY MDPMCLDAFPKLVCFKKRIEAI PQIDKYLKSSKYIAWPLQG WQATFGGGDHPPKSDLEVL FQGPLGSATMYPYDVPDYASKS <b>ESPKEPEQLRKLFIGGLSFETTDESLRSHFEQWGTLTDCVV</b> <b>MRDPNTKRSRGGFVTYATVEEVDAAMNARPHKVDGRVVEP</b> <b>KRAVSREDSQRPGAHLTVKKIFVGGIKEDTEEHHLRDYFEQ</b> <b>YGKIEVIEIMTDRGSGKKRGFAFVTFDDHDSVDKIVI QKYH</b> <b>TVNGHNCEVRKALSKQEMASA</b></p>
<b><u>Native sequence</u></b>	<p>Amino acids S2 – A189 of human hnRNP A1. [Full length protein ends at residue F372] Residue G244 of the fusion protein is equivalent to S2 of the native protein. The GST tag is located at residues 1 – 220 and the HA tag is located at residues 235 – 243.</p>
<b><u>Protease cleavage</u></b>	PreScission ( <u>LEVLFQGPL</u> ) residues 221 - 229
<b><u>Cloning sites</u></b>	<i>Bam</i> H1 of pGEX-6P-1

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**Complete  
nucleotide sequence**

ATGTCCCCTATACTAGGTTATTGGAAAATTAAGGGCCTTGT  
GCAACCCACTCGACTTCTTTTTGGAATATCCTTGAAGAAAAAT  
ATGAAGAGCATTGTATGAGCGCGATGAAGGTGATAAATGG  
CGAAACAAAAAGTTTGAATTGGGTTTGGAGTTTCCCAATCT  
TCCTTATTATATTGATGGTGATGTTAAATTAACACAGTCTA  
TGGCCATCATACTTATATAGCTGACAAGCACAAACATGTTG  
GGTGGTTGTCCAAAAGAGCGTGCAGAGATTTCAATGCTTGA  
AGGAGCGGTTTTTGGATATTAGATACGGTGTTTTCGAGAATTG  
CATATAGTAAAGACTTTGAAACTCTCAAAGTTGATTTTCTT  
AGCAAGCTACCTGAAATGCTGAAAATGTTCGAAGATCGTTT  
ATGTCATAAAACATATTTAAATGGTGATCATGTAACCCATC  
CTGACTTCATGTTGTATGACGCTCTTGATGTTGTTTTTATAC  
ATGGACCCAATGTGCCTGGATGCGTTCCCAAATTAGTTTG  
TTTTAAAAACGTATTGAAGCTATCCCACAAATTGATAAGT  
ACTTGAAATCCAGCAAGTATATAGCATGGCCTTTGCAGGGC  
TGGCAAGCCACGTTTGGTGGTGGCGACCATCTCCAAAATC  
GGATCTGGAAGTCTGTTCCAGGGGCCCTGGGATCCGCCA  
CCATGTACCCATACGATGTGCCAGATTACGCCTCTAAGTCA  
GAGTCTCCTAAAGAGCCCGAACAGCTGAGGAAGCTCTTCAT  
TGGAGGGTTGAGCTTTGAAACAACCTGATGAGAGCCTGAGGA  
GCCATTTTGAGCAATGGGGAACGCTCACGGACTGTGTGGTA  
ATGAGAGATCCAAACACCAAGCGCTCCAGGGGCTTTGGGTT  
TGTCACATATGCCACTGTGGAGGAGGTGGATGCAGCTATGA  
ATGCAAGGCCACACAAGGTGGATGGAAGAGTTGTGGAACCA  
AAGAGAGCTGTCTCCAGAGAAGATTCTCAAAGACCAGGTGC  
CCACTTAACTGTGAAAAAGATATTTGTTGGTGGCATTAAAG  
AAGACACTGAAGAACATCACCTAAGAGATTATTTTGAACAG  
TATGGAAAAATTGAAGTGATTGAAATCATGACTGACCGAGG  
CAGTGGCAAGAAAAGGGGCTTTGCCTTTGTAACCTTTGACG  
ACCATGACTCCGTGGATAAGATTGTCATTTCAGAAATACCAT  
ACTGTGAATGGCCACAACCTGTGAAGTTAGAAAAGCCCTGTC  
AAAGCAAGAGATGGCTAGTGCTtaa