

MRCPPU REAGENTS and SERVICES

Standard Operating Procedure

Preparation of CMTR1 [143 – 835]

Enzyme description:- CMTR1 [143 - 835]

Clone number:- DU 46710

Source:- Recombinant

Expression system:- *E.coli*

Tag:- N-terminal GST

Purification method:- GSH Agarose

Calculated molecular mass:-

Monoisotopic 107, 417.41 daltons

Average Mass 107, 486.75 daltons

[cysteines reduced, methionines have not been oxidized]

Theoretical pI:- 6.20

Purity:- >80 %

Enzyme storage buffer:-

50 mM Tris-HCl pH 7.5, 0.1 mM EGTA, 150 mM NaCl, 0.1 % 2-mercaptoethanol,
270 mM Sucrose

Storage temperature:- -70 °C

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Clone Data Sheet

CMTR1 [143 - 835]

<u>Protein</u>	CMTR1 [143 – 835]
<u>Clone number</u>	DU 46710
<u>Species</u>	Human
<u>Accession number</u>	Q8N1G2
<u>Tags</u>	N-terminal GST
<u>Bacterially expressed protein</u>	<p>MSPILGYWKIKGLVQPTRLLEYLEEKYEEHLYERDEGDKWRNKKFELG LEFPNLPYYIDGDVKLTQSMAIIRYIADKHNMLGGCPKERAELSMLEGA VLDIRYGVSR IAYS KDFETLKVDFLSKLP EMLKMFEDRLCHKTYLNGDH VTHPDFMLYDALDVVLYMDPMCLDAFPKLVCFKKRIEAI PQIDKYLKSS KYIAWPLQGWQATFGGGDHPPKSDLEVL<u>FQGPL</u>GSPNSRVEWRDEPEPS ACEQVSWFPECTTEIPDTQEMSDWMVVGKRKMI IEDETEFCGEELLHSV LQCKSVFDVLDGEEMRRARTRANPYEMIRGVFFLNRAAMKMANMDFVFD RMFTNPRDSYGKPLVKDREAELLYFADV CAGPGGFSEYVLWRKKWHAKG FGMTLKGPNDFKLEDFYSASSELFEPYYGEGGIDGDGDITRPENISAFR NFVLDNDRKGVHFLMADGGFSVEGQENLQEILSKQLLLCQFLMALSIV RTGGHFICKTFDLFTPFVGLVYLLYCCFERVCLFKPITSRPANSERYV VCKGLKVGIDDVRDYLFAVNIKLNQLRNTSDVNLVVPLEVIKGDHEFT DYMIRSNESHCSLQIKALAKIHAFVQDTTLSEPRQAEIRKECLRLWGIP DQARVAPSSSDPKSKFFELIQGTEIDIFSYPKPTLLTSKTLEKIRPVFDY RCMVSGSEQFLIGLGSQIYTW DGRQSDRWIKLDLKTLPDRTLLSVE IVHELKGEKQAQRKISAIHILDVVLNGLTDVREQHFNQRIQLAEKFKVKA VSKPSRPDMNPIRVKEVYRLEEMEKIFVRLEMKIIKGSSTPKLSYTGR DDRHFVPMGLYIVRTVNEPWTMGFSKSEFKKFFYNKKT KDSTFDLPADS IAPFHICYGRLEFEWGDGIRVHDSQKPQDQDKLSKEDVLSFIQM HRA</p>
<u>Native sequence</u>	Amino acids W143 – A835 (end residue) of human CMTR1. Residue W238 of the fusion protein is equivalent to W143 of the native enzyme. The GST tag is located at residues 1 – 220.
<u>Protease cleavage</u>	PreScission (<u>LEVL FQGP</u>) residues 221 - 229
<u>Cloning sites</u>	<i>Sal</i> I and <i>Not</i> I into <i>Xho</i> I and <i>Not</i> I sites of pGEX 6P-1

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Nucleotide Sequence

ggctcgagTGGCGAGATGAGCCAGAGCCCAGTGCTTGTGAGCAGGTGTC
ATGGTTTTCCAGAATGTACCACTGAAATTCCTGACACTCAGGAAATGAGC
GATTGGATGGTGGTGGGAAAGAGAAAGATGATTATTGAAGATGAAACAG
AGTTTTGTGGGAAGAGCTGCTTCACAGTGTGTTGCAGTGTAAAGAGCGT
GTTTGATGTCTTGGATGGGGAAGAGATGCGGCGAGCTCGGACTCGGGCC
AATCCCTATGAGATGATCCGAGGAGTCTTCTTTCTAAACAGGGCAGCAA
TGAAGATGGCTAACATGGATTTTGTATTTGATCGCATGTTCACAAATCC
GCGGGACTCTTATGGGAAGCCACTGGTGAAGGACCGGGAAGCTGAGCTT
CTGTACTTTGCTGATGTCTGCGCAGGCCAGGTGGCTTCTCAGAGTATG
TGCTGTGGAGGAAGAAGTGGCATGCAAAGGGCTTTGGAATGACTTTGAA
GGGCCCTAATGACTTCAAGCTGGAGGACTTCTACTCTGCTTCCAGTGAA
CTCTTCGAACCCACTATGGTGAGGGTGGGATTGATGGAGATGGAGATA
TCACCCGCCAGAGAACATCTCTGCTTTTCGGAATTTTGTCTGGATAA
CACAGATCGCAAGGGTGTCCATTTTCTGATGGCTGATGGGGGTTTCTCG
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TGTGTCAGTTCCTCATGGCGCTGTCCATTGTCCGGACAGGAGGCCACTT
CATCTGTAAAACCTTTGACCTGTTACACCGTTTAGTGTGGGGCTTGTC
TACCTGCTGTACTGCTGCTTTGAACGAGTTTGTCTCTTCAAGCCTATTA
CCAGCCGTCCTGCCAACTCAGAGAGGTATGTGGTGTGCAAGGGCCTGAA
GGTGGGCATAGATGATGTTCCGGATTACCTCTTCGCAGTGAATATTTAAA
CTCAATCAGCTGCGGAACACGGATTCCGACGTCAACTTGGTGGTCCCC
TGGAGGTGATCAAGGGAGACCATGAATTTACTGACTACATGATACGGTC
CAATGAGAGCCACTGTAGTCTGCAGATCAAAGCTCTGGCGAAAATCCAT
GCCTTTGTTCAAGACACGACACTGAGTGAGCCTCGACAGGCAGAGATAC
GGAAGGAGTGCCCTCCGACTCTGGGGGATCCAGACCAGGCTCGTGTGGC
TCCTTCTCCTCCGACCCATAATCGAAGTTCTTTGAGCTAATCCAGGGC
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GCCCAGCATGAATCCCATCAGGGTGAAGGAGGTGTACAGACTGGAAGAG
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AGGACTCTACTTTTGACCTCCCTGCAGACTCCATTGCCCCATTTACAT
TTGCTACTATGGCCGGCTCTTCTGGGAGTGGGGGGATGGCATTTCGTGTG
CATGACTCCCAGAAGCCCCAGGACCAGGACAAGCTGTCCAAGGAGGACG
TCCTCTCCTTCATCCAGATGCACAGGGCCtaagcgccgc