

## Product datasheet for MC208421

### Edn3 (NM\_007903) Mouse Untagged Clone

#### Product data:

Product Type:	Expression Plasmids
Product Name:	Edn3 (NM_007903) Mouse Untagged Clone
Tag:	Tag Free
Symbol:	Edn3
Synonyms:	ET-3; Is; PPET3; tmgc48
Mammalian Cell Selection:	Neomycin
Vector:	pCMV6-Entry (PS100001)
E. coli Selection:	Kanamycin (25 ug/mL)
Fully Sequenced ORF:	>NCBI ORF sequence for NM_007903, the custom clone sequence may differ by one or more nucleotides

ATGGAGCCGGGCTGTGGCTCCTTCTCGGGCTCACAGTGACCTCCGCTGCAGGACTTGTGCCTTGCCCC  
 AGTCTGGGACTCTGGCAGAGCCAGTGTGTCCCAGGGTCCCCCTGAAGCTGGATCAGAGAGGGGCTGTGA  
 AGAGACTGTGGCTGGCCCTGGTGAGAGGATTGTGTCCCCAACAGTTGCACTGCCTGCACAGCCTGAAAGC  
 GCTGGGCAGGAACGGGCACCGAGGCTGTGGGAAACAAGAGGACAAGGGGCTGCCTGCACACCACCGCC  
 CTCGCCGCTGCACGTGCTTCACCTACAAGGACAAGGAGTGTGTCTACTATTGCCACCTGGACATCATCTG  
 GATTAACACTCCTGAACAGACTGTGCCCTATGGACTGTCCAACACAGAGAAAAGCCTTCGGGAAAGAGG  
 TCCTTGGGGCCAGTTCCAGAAAGCTCCCAGCCTTCTCCGTGGACACGCTTGCCTTGTACTTGTATGGGG  
 CGGATGACAAGGCCTGTGCACACTTCTGTGCACGACAGAGATGTCACCAAGTTATCCGGGAGAGCAGA  
 AAGGCCAGCTGCAGAAGAGATGCGGGAGACTGGAGGCCACGTCAAAGGTTGATGTCAAGGACAGATAAA  
 GCCCACCAGCCTTAG

Restriction Sites:	SgfI-MluI
ACCN:	NM_007903
Insert Size:	645 bp


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<b>OTI Disclaimer:</b>	<p>Due to the inherent nature of this plasmid, standard methods to replicate additional amounts of DNA in E. coli are highly likely to result in mutations and/or rearrangements. Therefore, OriGene does not guarantee the capability to replicate this plasmid DNA. Additional amounts of DNA can be purchased from OriGene with batch-specific, full-sequence verification at a reduced cost. Please contact our customer care team at <a href="mailto:custsupport@origene.com">custsupport@origene.com</a> or by calling 301.340.3188 option 3 for pricing and delivery.</p> <p>The molecular sequence of this clone aligns with the gene accession number as a point of reference only. However, individual transcript sequences of the same gene can differ through naturally occurring variations (e.g. polymorphisms), each with its own valid existence. This clone is substantially in agreement with the reference, but a complete review of all prevailing variants is recommended prior to use. <a href="#">More info</a></p>
<b>Components:</b>	The ORF clone is ion-exchange column purified and shipped in a 2D barcoded Matrix tube containing 10ug of transfection-ready, dried plasmid DNA (reconstitute with 100 ul of water).
<b>Reconstitution Method:</b>	<ol style="list-style-type: none"> <li>1. Centrifuge at 5,000xg for 5min.</li> <li>2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.</li> <li>3. Close the tube and incubate for 10 minutes at room temperature.</li> <li>4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid at the bottom.</li> <li>5. Store the suspended plasmid at -20°C. The DNA is stable for at least one year from date of shipping when stored at -20°C.</li> </ol>
<b>Note:</b>	Plasmids are not sterile. For experiments where strict sterility is required, filtration with 0.22um filter is required.
<b>RefSeq:</b>	<a href="#">BC137727</a> , <a href="#">AAI37728</a>
<b>RefSeq Size:</b>	2412 bp
<b>RefSeq ORF:</b>	645 bp
<b>Locus ID:</b>	13616
<b>UniProt ID:</b>	<a href="#">P48299</a>
<b>Cytogenetics:</b>	2 98.1 cM
<b>Gene Summary:</b>	<p>This gene is a member of the endothelin family whose members encode proteins that act on G protein-coupled receptors. Endothelins are produced as large prepropeptide precursors that undergo a first cleavage by a subtilisin serine protease to form an inactive intermediate, which in turn is cleaved again by endothelin-converting enzyme 1 (ECE-1) to yield the active 21 amino acid peptide. This gene encodes a protein which is expressed in neural crest cells (NCC), binds to endothelin receptor b (Ednrb) and plays an essential role in the development of NCC-derived cell lineages including melanocytes and enteric neurons. Mutations in this gene are associated with terminal aganglionosis and white spotted coat in mice and Hirschsprung's disease and Waardenburg syndrome in humans. [provided by RefSeq, Apr 2013]</p>