

Product datasheet for **MG215244**

Scn2a (NM_001099298) Mouse Tagged ORF Clone

Product data:

Product Type:	Expression Plasmids
Product Name:	Scn2a (NM_001099298) Mouse Tagged ORF Clone
Tag:	TurboGFP
Symbol:	Scn2a
Synonyms:	6430408L10; A230052E19Rik; Nav; Nav1.2; Scn; Scn2a1
Mammalian Cell Selection:	Neomycin
Vector:	pCMV6-AC-GFP (PS100010)
E. coli Selection:	Ampicillin (100 ug/mL)
ORF Nucleotide Sequence:	>MG215244 representing NM_001099298 Red=Cloning site Blue=ORF Green=Tags(s)

TTTTGTAATACGACTCACTATAGGGCGGCCGGAATTCGTCGACTGGATCCGGTACCGAGGAGATCTGCC
GCC**CGATCGCC**

ATGGCACAATCAGTGCTGGTACCGCCAGGACCTGACAGCTTCGGCTTCTTTACCAGGGAATCCCTTGCTG
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TCAAAGTTAAGTTCCAAGAGTGAAAAGGAGCTGAAAAACAGAAGGAAGAAAAAGAAACAGAAAGAGCAGG
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ACGCGTACGCGGCCGCTCGAG - GFP Tag - GTTTAA

Protein Sequence:

>MG215244 representing NM_001099298

Red=Cloning site Green=Tags(s)

MAQSVLVPPGPDSEFRFFTTRESLAAIEQRIAEKAKRPKQERKDEDDENGPKPNSDLEAGKSLPFIYGDIP
 PEMVSEPLEDLDPYYINKKTFIVLNKKGKISRFSATSALYILTPFNPIRKLAIKILVHSLFNVLMCTIL
 TNCVFMSTNSPPDWTKNVEYFTGTIYTFESLIKILARGFCLEDFTLRDPWNWLDFTVITFAVYTFEVLN
 GNVSALRTFRVLRALKTISVIPGLKTIVGALIQSVKKLSDVMILTVFCLSVFALIGLQLFMGNLRNKCLQ
 WPPDNSTFEINITSSFNNSLDWNGTAFNRMTMFMNWDIYIEDKSHFYFLEGQNDALLCGNSSDAGQCPEG
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 KVVAYGFQMYFTNAWCWLDLIVDVSLVSLTANALGYSELGAIKSLRTRLRALRPLRALSRFEGMRVVVNA
 LLGAIPTIMNVLLVCLIFWLIFSIMGVNLFAGKFYHCINYYTTGEMFDVSVVNNYSECQALIESNQTARWK
 NVKVNFDNVGLGYLSLLQVATFKGWDIMYAAVDSRNVELQPKYEDNLYMYLYFVIFIFIGSFTLNLFI
 GVIIDNFNQKKKFGGQDIFMTEEQKYYNAMKKLGSKKPQKPIPRPANKFQGMVDFVTKQVFDISIMI
 LICLNMVMTMMVETDDQSQEMTNILYWINLVFIVLFTGECVLKLI SLRHYYFTIGWNIIFDVVVILSIVGM
 FLAELIEKYFVSPTLFRVIRLARIGRILRLIKGAKGIRTLFALMMSLPALFNIGLLLFLVMFIYAFGM
 SNFAYVKREVIGDDMFNFETFGNSMICLFQITTSAGWDGLLAPILNSGPPDCDPEKDHPGSSVKGDCGNP
 SVGIFFFVSYIIISFLVVNMYIAVILENF SVATEESAEPLEDDFEMFYEVEKFDPDATQFIEFCKLS
 DFAAALDPPLLIAKPNKVQLIAMDLPMVSGDRIHCLDILFAFTKRVLGESEMDALRIQMEERFMASNPS
 KVSYESITTTLKRKQEEVSAIIVIQRAYRRLKQKVKVSSIYKDKGKEDEGTPIKEDIITDKLNENST
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TRTRPLE - GFP Tag - V

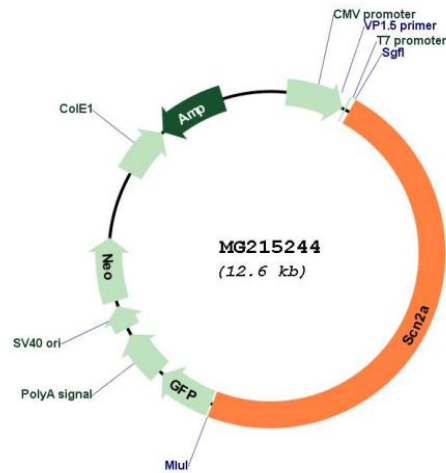
Restriction Sites:

Sgfl-MluI

Cloning Scheme:



Plasmid Map:



ACCN: NM_001099298

ORF Size: 6018 bp

OTI Disclaimer: 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. [More info](#)

OTI Annotation: This clone was engineered to express the complete ORF with an expression tag. Expression varies depending on the nature of the gene.

Components:	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).
Reconstitution Method:	<ol style="list-style-type: none">1. Centrifuge at 5,000xg for 5min.2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.3. Close the tube and incubate for 10 minutes at room temperature.4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid at the bottom.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.
RefSeq:	<u>NM_001099298.3</u>
RefSeq Size:	8690 bp
RefSeq ORF:	6021 bp
Locus ID:	110876
UniProt ID:	<u>B1AWN6</u>
Cytogenetics:	2 38.61 cM
Gene Summary:	Voltage-gated sodium channels are transmembrane glycoprotein complexes composed of a large alpha subunit with four repeat domains, each of which is composed of six membrane-spanning segments, and one or more regulatory beta subunits. Voltage-gated sodium channels are responsible for the generation and propagation of action potentials in neurons and muscle. This gene encodes one member of the sodium channel alpha subunit gene family. In humans, variants of this gene are associated with seizure disorders and autism spectrum disorder. Mice homozygous for a knockout mutation die with severe hypoxia and extensive neuronal cell death, while gain of function mutations result in progressive seizure disorder. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Nov 2016]