

Product datasheet for RG214427

SAMD9L (NM_152703) Human Tagged ORF Clone

Product data:

Product Type: Expression Plasmids

Product Name: SAMD9L (NM_152703) Human Tagged ORF Clone

Tag: TurboGFP Symbol: SAMD9L

Synonyms: ATXPC; C7orf6; DRIF2; M7MLS1; UEF1

Mammalian Cell Neomycin

Selection:

Vector:

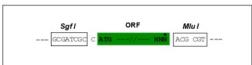
pCMV6-AC-GFP (PS100010)

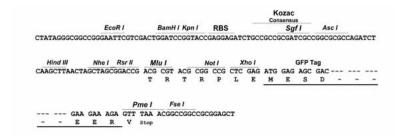
E. coli Selection: Ampicillin (100 ug/mL)

Restriction Sites: Sgfl-Mlul

Cloning Scheme:

Cloning sites used for ORF Shuttling:







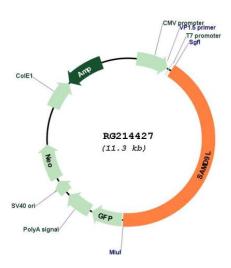
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Plasmid Map:



ACCN: NM_152703

ORF Size: 4752 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: 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 152703.5</u>



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RefSeq Size: 7133 bp
RefSeq ORF: 4755 bp
Locus ID: 219285
UniProt ID: Q8IVG5
Cytogenetics: 7q21.2

Gene Summary: This gene encodes a cytoplasmic protein that acts as a tumor suppressor but also plays a key

role in cell proliferation and the innate immune response to viral infection. The encoded protein contains an N-terminal sterile alpha motif domain. Naturally occurring mutations in this gene are associated with myeloid disorders such as juvenile myelomonocytic leukemia, acute myeloid leukemia, and myelodysplastic syndrome. Naturally occurring mutations are also associated with hepatitis-B related hepatocellular carcinoma, normophosphatemic familial tumoral calcinosis, and ataxia-pancytopenia syndrome. [provided by RefSeq, Apr

2017]