

Product datasheet for SC329874

METTL23 (NM 001206986) Human Untagged Clone

Product data:

Product Type: Expression Plasmids

Product Name: METTL23 (NM_001206986) Human Untagged Clone

Tag: Tag Free Symbol: METTL23

Synonyms: C17orf95; MRT44

Vector: pCMV6-Entry (PS100001)

Fully Sequenced ORF: >SC329874 representing NM_001206986.

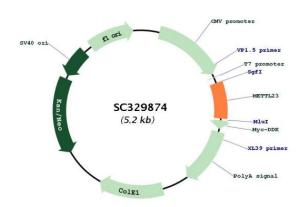
Blue=Insert sequence Red=Cloning site Green=Tag(s)

ATGAATAACCTGCCACATCTGCAGGTGGTAGGACTAACATGGGGTCATATATCTTTGGGATCTTCTGGCT CTACCACCACAAGATATTATCCTTGCATCTGATGTGTTCTTTGAACCAGAAGATTTTGAAGACATTTTG GCTACAAATATATTTTTTGATGCACAAGAATCCCAAGGTCCAATTGTGGTCTACTTATCAAGTTAGGAGT GCTGACTGGTCACTTGAAGCTTTACTCTACAAATGGGATATGAAATGTGTCCACATTCCTCTTGAGTCT TTTGATGCAGACAAAGAAGATATAGCAGAATCTACCCTTCCAGGAAGACATACAGTTGAAATGCTGGTC

ATTTCCTTTGCAAAGGACAGTCTCTGA

Restriction Sites: Sgfl-Mlul

Plasmid Map:



ACCN: NM_001206986



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Insert Size: 372 bp

OTI Disclaimer: Our molecular clone sequence data has been matched to the reference identifier above as a

point of reference. Note that the complete sequence of our molecular clones may differ from the sequence published for this corresponding reference, e.g., by representing an alternative

RNA splicing form or single nucleotide polymorphism (SNP).

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 001206986.1</u>

 RefSeq Size:
 1082 bp

 RefSeq ORF:
 372 bp

 Locus ID:
 124512

 UniProt ID:
 Q86XA0

 Cytogenetics:
 17q25.1

MW:

Gene Summary: The protein encoded by this gene functions as a transcription factor regulator in the

transcriptional pathway for human cognition. It is a partner of the alpha subunit of the GA-binding protein transcription factor. Mutations in this gene cause mild autosomal recessive intellectual disability. Alternative splicing results in multiple transcript variants. [provided by

RefSeq, Nov 2014]

14.2 kDa

Transcript Variant: This variant (5) uses a different splice site in the 5' UTR, lacks an alternate exon that includes a portion of the 5' coding region, and uses a downstream in-frame start

codon, compared to variant 1. The encoded isoform (2) is shorter at the N-terminus,

compared to isoform 1. Variants 4, 5, 6 and 9 all encode isoform 2.