

## Product datasheet for RC230776L4V

## OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

## MADH7 (SMAD7) (NM 001190822) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

**Product Name:** MADH7 (SMAD7) (NM\_001190822) Human Tagged ORF Clone Lentiviral Particle

Symbol: SMAD7

Synonyms: CRCS3; MADH7; MADH8

**Mammalian Cell** 

iammanan cen

Selection:

Puromycin

**Vector:** pLenti-C-mGFP-P2A-Puro (PS100093)

Tag: mGFP

**ACCN:** NM\_001190822

ORF Size: 1281 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC230776).

Sequence:

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.

**RefSeq:** NM 001190822.1, NP 001177751.1

 RefSeq Size:
 2293 bp

 RefSeq ORF:
 636 bp

 Locus ID:
 4092

 UniProt ID:
 015105

Cytogenetics: 18q21.1

**Protein Families:** Druggable Genome, Transcription Factors

**Protein Pathways:** TGF-beta signaling pathway





**MW:** 46.4 kDa

**Gene Summary:** The protein encoded by this gene is a nuclear protein that binds the E3 ubiquitin ligase

SMURF2. Upon binding, this complex translocates to the cytoplasm, where it interacts with TGF-beta receptor type-1 (TGFBR1), leading to the degradation of both the encoded protein and TGFBR1. Expression of this gene is induced by TGFBR1. Variations in this gene are a cause of susceptibility to colorectal cancer type 3 (CRCS3). Several transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, Jun 2010]