

## Product datasheet for **SC205203**

### OGDH (NM\_001003941) Human 3' UTR Clone

#### Product data:

Product Type:	3' UTR Clones
Product Name:	OGDH (NM_001003941) Human 3' UTR Clone
Vector:	pMirTarget (PS100062)
Symbol:	OGDH
Synonyms:	AKGDH; E1k; KGD1; OGDC; OGDH2
ACCN:	NM_001003941
Insert Size:	385 bp
Insert Sequence:	>SC205203 3'UTR clone of NM_001003941 The sequence shown below is from the reference sequence of NM_001003941. The complete sequence of this clone may contain minor differences, such as SNPs. Blue=Stop Codon Red=Cloning site  GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAAGCCAAGAAGGGCGGAAAGATCGCCGTG TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC TCCAGCATGGAGTTCGGCTCACCAACA <b>TA</b> CCCAGAGCCCTGGGTGCATCTAGACTTTAAAAAATATT TAAAGTCGGCCGGCGCAGTGTCTCACGCCTGTAATCCCAGCACTTTGGGAGGCCGAGGTGGGCAGATC ACCTGAGTTCGGGAGTTGGAGACCAGCCTGACCAACATGGAGAACTCCATCTCTACTAAAAATACAAA ATTAGCTGGGCGTGGTGGCGCGCCTGTAAATCCCAGCTACTCAGGAGGCTGAGGCAGGAGAATCGCTT GAACCCGGGAGGTGGAGTTGCACTGAGCCGAGATTACGCCATTGCACTCCAGCCTGGGCCAACAAAGAG CGAAACTCTGTCTCAAAGAAAAAATAAATAAATAAAAAA <b>ACGCGT</b> AAGCGGCCGCGCATCTAGATTCAAGAAAATGACCGACCAAGCGACGCCAACCTGCCATCA CGAGATTCGATTCCACCGCCCTTCTATGAAAGG
Restriction Sites:	Sgfl-MluI
OTI Disclaimer:	Our molecular clone sequence data has been matched to the sequence identifier above as a point of reference. Note that the complete sequence of this clone is largely the same as the reference sequence but may contain minor differences , e.g., single nucleotide polymorphisms (SNPs).
Components:	The cDNA clone is shipped in a 2-D bar-coded Matrix tube as 10 ug dried plasmid DNA. The package also includes 100 pmols of both the corresponding 5' and 3' vector primers in separate vials.
RefSeq:	<u><a href="#">NM_001003941.3</a></u>



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**Summary:** This gene encodes one subunit of the 2-oxoglutarate dehydrogenase complex. This complex catalyzes the overall conversion of 2-oxoglutarate (alpha-ketoglutarate) to succinyl-CoA and CO<sub>2</sub> during the Krebs cycle. The protein is located in the mitochondrial matrix and uses thiamine pyrophosphate as a cofactor. A congenital deficiency in 2-oxoglutarate dehydrogenase activity is believed to lead to hypotonia, metabolic acidosis, and hyperlactatemia. Alternative splicing results in multiple transcript variants encoding distinct isoforms.[provided by RefSeq, Sep 2009]

**Locus ID:** 4967

**MW:** 14.3