

# Rabbit Anti-MARS antibody

# SL18682R

<b>Product Name:</b>	MARS
Chinese Name:	甲硫氨酸转运RNA合成酶抗体
Alias:	cytoplasmic; Mars; Methionine tRNA ligase 1, cytoplasmic; Methionine tRNA ligase; Methionine tRNA synthetase; MethioninetRNA ligase; Methionyl tRNA synthetase; Methionyl-tRNA synthetase; MetRS; MTRNS; SYMC_HUMAN.
Organism Species:	Rabbit
Clonality:	Polyclonal
React Species:	Human, Mouse, Rat, Pig, Cow, Sheep,
Applications:	WB=1:500-2000ELISA=1:500-1000IHC-P=1:400-800IHC-F=1:400-800ICC=1:100-500IF=1:100-500 (Paraffin sections need antigen repair) not yet tested in other applications. optimal dilutions/concentrations should be determined by the end user.
Molecular weight:	101kDa
Cellular localization:	cytoplasmic
Form:	Lyophilized or Liquid
Concentration:	1mg/ml
immunogen:	KLH conjugated synthetic peptide derived from human MARS:1-100/900
Lsotype:	IgG
Purification:	affinity purified by Protein A
Storage Buffer:	0.01M TBS(pH7.4) with 1% BSA, 0.03% Proclin300 and 50% Glycerol.
Storage:	Store at -20 °C for one year. Avoid repeated freeze/thaw cycles. The lyophilized antibody is stable at room temperature for at least one month and for greater than a year when kept at -20 °C. When reconstituted in sterile pH 7.4 0.01M PBS or diluent of antibody the antibody is stable for at least two weeks at 2-4 °C.
PubMed:	<u>PubMed</u>
Product Detail:	This gene encodes a member of the class I family of aminoacyl-tRNA synthetases. These enzymes play a critical role in protein biosynthesis by charging tRNAs with their cognate amino acids. The encoded protein is a component of the multi-tRNA synthetase complex and catalyzes the ligation of methionine to tRNA molecules. [provided by RefSeq, Jan 2011]

### **Subunit:**

Component of the multisynthetase complex which is comprised of a bifunctional glutamyl-prolyl-tRNA synthetase, the monospecific isoleucyl, leucyl, glutaminyl, methionyl, lysyl, arginyl, and aspartyl-tRNA synthetases as well as three auxiliary proteins, p18, p48 and p43.

# **Subcellular Location:**

Cytoplasm.

#### **DISEASE:**

Infantile liver failure syndrome 2 (ILFS2) [MIM:615486]: A life-threatening disorder of hepatic function that manifests with liver failure in the first months of life. Clinical features include failure to thrive, hypotonia, intermittent lactic acidosis, aminoaciduria, hypothyroidism, interstitial lung disease, anemia, liver canalicular cholestasis, steatosis, and iron deposition. Note=The disease is caused by mutations affecting the gene represented in this entry.

# Similarity:

Belongs to the class-I aminoacyl-tRNA synthetase family.

Contains 1 GST C-terminal domain.

Contains 1 WHEP-TRS domain.

# **SWISS:**

P56192

# Gene ID:

4141

#### Database links:

Entrez Gene: 4141 Human

Entrez Gene: 216443 Mouse

Entrez Gene: 299851 Rat

Omim: 156560 Human

SwissProt: P56192 Human

SwissProt: Q68FL6 Mouse

Unigene: 632707 Human

Unigene: 28173 Mouse

	Important Note: This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications.

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