abcam

Product datasheet

Recombinant Human YARS2/TyRS protein ab126680

1 Image

Description

Product name Recombinant Human YARS2/TyRS protein

Purity > 85 % SDS-PAGE.

ab126680 is purified using conventional chromatography techniques.

Expression system Escherichia coli

Accession Q9Y2Z4

Protein length Full length protein

Animal free No

Nature Recombinant

Species Human

Sequence MGSSHHHHHH SSGLVPRGSH MTLNLSVLLP

LGLRKAHSGA QGLLAAQKAR GLFKDFFPET

GTKIELPELF DRGTASFPQT IYCGFDPTAD SLHVGHLLAL

LGLFHLQRAG HNVIALVGGA TARLGDPSGR
TKEREALETE RVRANARALR LGLEALAANH
QQLFTDGRSW GSFTVLDNSA WYQKQHLVDF
LAAVGGHFRM GTLLSRQSVQ LRLKSPEGMS
LAEFFYQVLQ AYDFYYLFQR YGCRVQLGGS

DQLGNIMSGY EFINKLTGED VFGITVPLIT STTGAKLGKS

AGNAVWLNRD KTSPFELYQF FVRQPDDSVE

RYLKLFTFLP LPEIDHIMQL HVKEPERRGP QKRLAAEVTK

LVHGREGLDS AKRCTQALYH SSIDALEVMS DQELKELFKE APFSEFFLDP GTSVLDTCRK

ANAIPDGPRG YRMITEGGVS INHQQVTNPE SVLIVGQHIL

KNGLSLLKIG KRNFYIIKWL QL

Predicted molecular weight 54 kDa including tags

Amino acids 17 to 477

Tags His tag N-Terminus

Specifications

Our <u>Abpromise guarantee</u> covers the use of ab126680 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Applications SDS-PAGE

Form Liquid

Additional notes Protein previously labeled as YARS2.

Preparation and Storage

Stability and Storage Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C or -

80°C. Avoid freeze / thaw cycle.

pH: 8.00

Constituents: 0.02% DTT, 0.32% Tris HCl, 20% Glycerol (glycerin, glycerine), 0.58% Sodium

chloride

General Info

Function Catalyzes the attachment of tyrosine to tRNA(Tyr) in a two-step reaction: tyrosine is first activated

by ATP to form Tyr-AMP and then transferred to the acceptor end of tRNA(Tyr).

Involvement in diseaseDefects in YARS2 are the cause of myopathy with lactic acidosis and sideroblastic anemia type 2

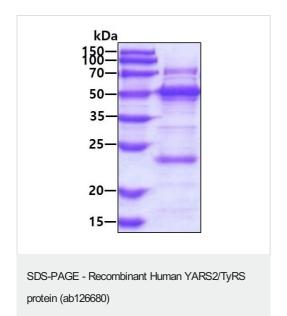
(MLASA2) [MIM:613561]. MLASA2 is a rare oxidative phosphorylation disorder specific to skeletal muscle and bone marrow. Affected individuals manifest sideroblastic anemia, progressive lethargy, muscle weakness, and exercise intolerance associated with persistent

lactic acidemia.

Sequence similaritiesBelongs to the class-I aminoacyl-tRNA synthetase family.

Cellular localization Mitochondrion matrix.

Images



3ug by SDS-PAGE under reducing conditions and visualized by coomassie blue stain.

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