SANTA CRUZ BIOTECHNOLOGY, INC.

IleRS (D-16): sc-34676



BACKGROUND

The fidelity of protein synthesis requires efficient discrimination of amino acid substrates by aminoacyl-tRNA synthetases. Accurate discrimination of the structurally similar amino acids valine and isoleucine by isoleucyl-tRNA synthetase (IIeRS) results, in part, from a hydrolytic editing reaction, which prevents misactivated valine from being stably joined to tRNAIIe. IIeRS joins IIe to tRNA(IIe) at its synthetic active site and hydrolyzes incorrectly acylated amino acids at its editing active site. A member of the aminoacyl-tRNA synthetase family, human IIeRS has been identified as a target of antibodies in the autoimmune disease polymyositis.

REFERENCES

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- Nordin, B.E., et al. 1999. RNA determinants for translational editing. Mischarging a minihelix substrate by a tRNA synthetase. J. Biol. Chem. 274: 6835-6838.
- Silvian, L.F., et al. 1999. Insights into editing from an IIe-tRNA synthetase structure with tRNAIIe and mupirocin. Science 285: 1074-1077.
- Nakama, T., et al. 2001. Structural basis for the recognition of isoleucyladenylate and an antibiotic, mupirocin, by isoleucyl-tRNA synthetase. J. Biol. Chem. 276: 47387-47393.
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- 6. SWISS-PROT/TrEMBL (P41252). World Wide Web URL: http://www.expasy. ch/sprot/sprot-top.html

CHROMOSOMAL LOCATION

Genetic locus: IARS (human) mapping to 9q22.31; lars (mouse) mapping to 13 A5.

SOURCE

IleRS (D-16) is an affinity purified goat polyclonal antibody raised against a peptide mapping within an internal region of IleRS of human origin.

PRODUCT

Each vial contains 200 μg lgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-34676 P, (100 μ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

STORAGE

Store at 4° C, **D0 NOT FREEZE**. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

RESEARCH USE

For research use only, not for use in diagnostic procedures.

APPLICATIONS

IleRS (D-16) is recommended for detection of IleRS of mouse, rat and human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

IleRS (D-16) is also recommended for detection of IleRS in additional species, including equine, canine, bovine, porcine and avian.

Suitable for use as control antibody for IIeRS siRNA (h): sc-45473, IIeRS siRNA (m): sc-45474, IIeRS shRNA Plasmid (h): sc-45473-SH, IIeRS shRNA Plasmid (m): sc-45474-SH, IIeRS shRNA (h) Lentiviral Particles: sc-45473-V and IIeRS shRNA (m) Lentiviral Particles: sc-45474-V.

Molecular Weight of IleRS: 145 kDa.

Positive Controls: Ramos cell lysate: sc-2216.

RECOMMENDED SECONDARY REAGENTS

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use donkey anti-goat IgG-HRP: sc-2020 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible donkey anti-goat IgG-HRP: sc-2033 (dilution range: 1:2000-1:5000), Cruz Marker™ Molecular Weight Standards: sc-2035, TBS Blotto A Blocking Reagent: sc-2333 and Western Blotting Luminol Reagent: sc-2048. 2) Immunofluo-rescence: use donkey anti-goat IgG-FITC: sc-2024 (dilution range: 1:100-1:400) or donkey anti-goat IgG-TR: sc-2783 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941.

PROTOCOLS

See our web site at www.scbt.com or our catalog for detailed protocols and support products.