

α 2ML1 (N-14): sc-138825

BACKGROUND

α -2-Macroglobulin (α -2M) is a homotetrameric serum protein consisting of four identical subunits that form dimers through disulfide bonds. Initially, α -2M was characterized as a pan-proteinase inhibitor that was able to "bait" proteinases into cleaving specific peptide sequences on α -2M. This interaction induces a conformational change in α -2M, thus enabling it to "trap" the proteinase and further inhibit its activity. Subsequently, α -2M has been shown to function as a carrier protein and regulator of cytokines during inflammation. Circulating transforming growth factor β (TGF β) in serum is primarily bound to α -2M, which renders TGF β inactive. Mutations and deletions in the gene encoding α -2M are associated with an increased incidence of Alzheimer's disease (AD). α -2-macroglobulin-like protein 1 (α 2ML1) is a related protein that is expressed in the epidermis and may play a role in keratinocyte differentiation.

REFERENCES

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CHROMOSOMAL LOCATION

Genetic locus: A2ML1 (human) mapping to 12p13.31.

SOURCE

α 2ML1 (N-14) is an affinity purified goat polyclonal antibody raised against a peptide mapping near the N-terminus of α 2ML1 of human origin.

STORAGE

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

PRODUCT

Each vial contains 200 μ g IgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-138825 P, (100 μ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

APPLICATIONS

α 2ML1 (N-14) is recommended for detection of α 2ML1 of human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2 μ g per 100-500 μ g of total protein (1 ml of cell lysate)], 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).

α 2ML1 (N-14) is also recommended for detection of α 2ML1 in additional species, including equine.

Suitable for use as control antibody for α 2ML1 siRNA (h): sc-95683, α 2ML1 shRNA Plasmid (h): sc-95683-SH and α 2ML1 shRNA (h) Lentiviral Particles: sc-95683-V.

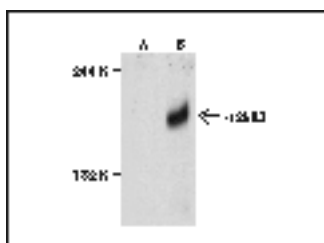
Molecular Weight of α 2ML1: 180 kDa.

Positive Controls: α 2ML1 (h): 293T Lysate: sc-372458.

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) Immunoprecipitation: use Protein A/G PLUS-Agarose: sc-2003 (0.5 ml agarose/2.0 ml). 3) Immunofluorescence: 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.

DATA



α 2ML1 (N-14) sc-138825. Western blot analysis of α 2ML1 expression in HeLa cells transfected with human α 2ML1 (sc-147792) and human α 2ML1 siRNA (sc-147792) (B) in control cells (A).

RESEARCH USE

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