

α -2M (9A3): sc-81541

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 inhibit its further activity. Subsequently, α -2M has also 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. α -2M also binds to IL-6 and, thereby, increases the concentration of IL-6 near lymphocytes, hepatocytes and stem cells involved in mediating the inflammatory cascade. Mutations and deletions in the gene encoding α -2M are associated with an increased incidence of Alzheimer's disease (AD), which is consistent with the role of α -2M in mediating the clearance and degradation of A β , the major component of β -Amyloid deposits accumulated during AD.

REFERENCES

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2. Tsuchiya, Y., et al. 1987. Sequence analysis of the putative regulatory region of rat α -2 macroglobulin gene. *Gene* 57: 73-80.
3. Borth, W., et al. 1990. Binding of IL-1 β to α macroglobulins and release by thioredoxin. *J. Immunol.* 145: 3747-3754.
4. Poller, W., et al. 1992. Cloning of the human α -2 macroglobulin gene and detection of mutations in two functional domains: the bait region and the thiolester site. *Hum. Genet.* 88: 313-319.
5. Webb, D.J., et al. 1998. Localization of the binding site for TGF β in human α -2 macroglobulin to a 20 kDa peptide that also contains the bait region. *J. Biol. Chem.* 273: 13339-13346.
6. Blacker, D., et al. 1998. α -2 macroglobulin is genetically associated with Alzheimer disease. *Nat. Genet.* 19: 357-360.

CHROMOSOMAL LOCATION

Genetic locus: A2M (human) mapping to 12p13.31.

SOURCE

α -2M (9A3) is a mouse monoclonal antibody raised against α -2M from plasma of human origin.

PRODUCT

Each vial contains 100 μ g IgG_{2b} in 1.0 ml PBS with < 0.1% sodium azide and 5% glycerol.

STORAGE

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

APPLICATIONS

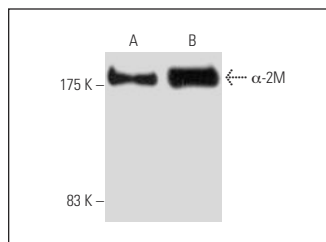
α -2M (9A3) is recommended for detection of α -2M of human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2 μ g per 100-500 μ g of total protein (1 ml of cell lysate)] and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

Suitable for use as control antibody for α -2M siRNA (h): sc-40297, α -2M shRNA Plasmid (h): sc-40297-SH and α -2M shRNA (h) Lentiviral Particles: sc-40297-V.

Molecular Weight of α -2M subunits: 185 kDa.

Molecular Weight of tetrameric α -2M: 718 kDa.

DATA



Immunoprecipitation of α -2M from human plasma (**B**) using α -2M (9A3): sc-81541 (mouse monoclonal antibody) followed by Western blot analysis using α -2M (2D9): sc-69750 (mouse monoclonal antibody) as compared with human plasma (**A**).

RESEARCH USE

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

PROTOCOLS

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