

# granzyme A (GA6): sc-56116

## BACKGROUND

Granzyme A and granzyme B are serine proteases that mediate apoptotic signaling in cytotoxic T lymphocytes (CTL) and natural killer (NK) cells. Both granzyme A and granzyme B are synthesized as inactive proenzymes, and they are stored within cytolytic granules and released by effector cells during degranulation. In activated CTLs, granzyme A and granzyme B are processed and activated by cathepsin C, and they then function to induce apoptosis by two distinct pathways. Granzyme B proteolytically cleaves and activates members of the caspase family of cysteine proteases, including caspase-3, caspase-6, caspase-7 and caspase-9. When cleaved, these caspases assemble into active holoenzymes that then mediate apoptosis through a defined proteolytic cascade involving nuclear lamins and PARP (poly ADP ribose polymerase). Granzyme A mediates the activation of apoptosis by inducing single-strand DNA breaks, membrane perturbation and nuclear condensations in an alternative pathway that is independent from caspase activation or the caspase proteolytic cascade.

## REFERENCES

1. Gershenfeld, H.K., et al. 1988. Cloning and chromosomal assignment of a human cDNA encoding a T cell- and natural killer cell-specific trypsin-like serine protease. *Proc. Natl. Acad. Sci. USA* 85: 1184-1188.
2. Shresta, S., et al. 1995. Natural killer and lymphokine-activated killer cells require granzyme B for the rapid induction of apoptosis in susceptible target cells. *Proc. Natl. Acad. Sci. USA* 92: 5679-5683.
3. Trapani, J.A., et al. 1996. A putative role in the mechanism of cytotoxic lymphocyte-mediated apoptosis. Localization of granzyme B in the nucleus. *J. Biol. Chem.* 271: 4127-4133.
5. Atkinson, E.A., et al. 1998. Cytotoxic T lymphocyte-assisted suicide. caspase-3 activation is primarily the result of the direct action of granzyme B. *J. Biol. Chem.* 273: 21261-21266.

## CHROMOSOMAL LOCATION

Genetic locus: GZMB (human) mapping to 14q11.2.

## SOURCE

granzyme A (GA6) is a mouse monoclonal antibody raised against granzyme A of human origin.

## PRODUCT

Each vial contains 50 µg IgG<sub>1</sub> in 500 µl of PBS with < 0.1% sodium azide and 0.1% gelatin.

## STORAGE

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

## PROTOCOLS

See our web site at [www.scbt.com](http://www.scbt.com) for detailed protocols and support products.

## APPLICATIONS

granzyme A (GA6) is recommended for detection of granzyme A of human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2 µg per 100-500 µg of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and immunohistochemistry (including paraffin-embedded sections) (starting dilution 1:50, dilution range 1:50-1:500).

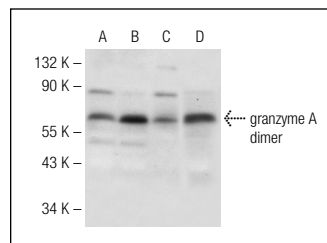
Suitable for use as control antibody for granzyme A siRNA (h): sc-37431, granzyme A shRNA Plasmid (h): sc-37431-SH and granzyme A shRNA (h) Lentiviral Particles: sc-37431-V.

Molecular Weight of granzyme A monomer: 28 kDa.

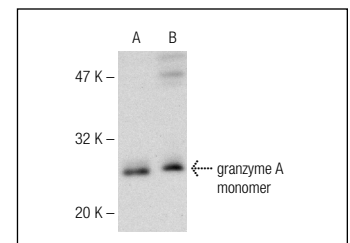
Molecular Weight of granzyme A homodimer: 43-65 kDa.

Positive Controls: HL-60 whole cell lysate: sc-2209, U-937 cell lysate: sc-2239 or K-562 whole cell lysate: sc-2203.

## DATA



granzyme A (GA6): sc-56116. Western blot analysis of granzyme A expression in U-937 (A), AML-193 (B), K-562 (C) and HL-60 (D) whole cell lysates.



granzyme A (GA6): sc-56116. Western blot analysis of granzyme A expression in CCRF-HSB-2 (A) and NK-92 (B) whole cell lysates.

## SELECT PRODUCT CITATIONS

1. Fogagnolo, L., et al. 2014. Cytotoxic granules in distinct subsets of cutaneous lupus erythematosus. *Clin. Exp. Dermatol.* 39: 835-839.

## RESEARCH USE

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