granzyme A (CB9): sc-56115



The Power to Question

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

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- 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.
- 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.
- 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.
- 5. Trapani, J.A., et al. 1998. Efficient nuclear targeting of granzyme B and the nuclear consequences of apoptosis induced by granzyme B and perforin are caspase-dependent, but cell death is caspase-independent. J. Biol. Chem. 273: 27934-27938.
- Pham, C.T., et al. 1999. Dipeptidyl peptidase I is required for the processing and activation of granzymes A and B *in vivo*. Proc. Natl. Acad. Sci. USA 96: 8627-8632.
- 7. Shresta, S., et al. 1999. granzyme A initiates an alternative pathway for granule-mediated apoptosis. Immunity 10: 595-605.

CHROMOSOMAL LOCATION

Genetic locus: GZMA (human) mapping to 5q11.2.

SOURCE

granzyme A (CB9) is a mouse monoclonal antibody raised against purified full length native granzyme A of human origin.

RESEARCH USE

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

PRODUCT

Each vial contains 200 $\mu g \ lgG_1$ kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

granzyme A (CB9) is available conjugated to either phycoerythrin (sc-56115 PE) or fluorescein (sc-56115 FITC), 200 μ g/ml, for WB (RGB), IF, IHC(P) and FCM.

APPLICATIONS

granzyme A (CB9) is recommended for detection of granzyme A of human origin by 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), flow cytometry (1 μ g per 1 x 10⁶ cells) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

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.

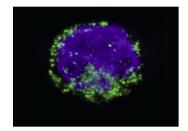
Molecular Weight of granzyme A monomer: 28 kDa.

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

RECOMMENDED SUPPORT REAGENTS

To ensure optimal results, the following support reagents are recommended: 1) Immunoprecipitation: use Protein A/G PLUS-Agarose: sc-2003 (0.5 ml agarose/2.0 ml). 2) Immunofluorescence: use m-lgG κ BP-FITC: sc-516140 or m-lgG κ BP-PE:sc-516141 (dilution range: 1:50-1:200) with UltraCruz® Mounting Medium:sc-24941 or UltraCruz® Hard-set Mounting Medium: sc-359850.

DATA



granzyme A (CB9): sc-56115. Immunofluorescence staining of human Cytotoxic T-Lymphocyte cells showing cytoplasmic granule (green color) localization. Kindly provided by Jerome Thiery and Judy Lieberman.

SELECT PRODUCT CITATIONS

 Shi, Z., et al. 2022. Microglia drive transient insult-induced brain injury by chemotactic recruitment of CD8+T lymphocytes. Neuron. E-published.

STORAGE

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