SANTA CRUZ BIOTECHNOLOGY, INC.

AIF (H-300): sc-5586



BACKGROUND

A key event in the apoptotic process is the opening of the mitochondrial permeability transition pore, an event that is regulated by Bcl-2 family proteins, resulting in the release of several proteins from the mitochondrial intermembrane space. Several of these proteins participate in apoptosis, including cytochrome c, procaspases-2, -3 and -9, and AIF (apoptosis-inducing factor). AIF was shown to cause DNA fragmentation and chromatin condensation, and to induce the release of cytochrome c and caspase-9 from mitochondria. Bcl-2 overexpression was shown to prevent the release of AIF from mitochondria, but not to block its apoptogenic activity.

REFERENCES

- Zamzami, N., et al. 1996. Mitochondrial control of nuclear apoptosis. J. Exp. Med. 183: 1533-1544.
- Susin, S.A., et al. 1996. Bcl-2 inhibits the mitochondrial release of an apoptogenic protease. J. Exp. Med. 184: 1331-1341.

CHROMOSOMAL LOCATION

Genetic locus: AIFM1 (human) mapping to Xq26.1; Aifm1 (mouse) mapping to X A4.

SOURCE

AIF (H-300) is a rabbit polyclonal antibody raised against amino acids 1-300 mapping at the N-terminus of AIF of human origin.

PRODUCT

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

Available as agarose conjugate for immunoprecipitation, sc-7384 AC, $500 \mu g/0.25 \text{ ml}$ agarose in 1 ml.

APPLICATIONS

AlF (H-300) is recommended for detection of AlF of mouse, rat and 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).

AIF (H-300) is also recommended for detection of AIF in additional species, including equine, canine, bovine and porcine.

Suitable for use as control antibody for AIF siRNA (h): sc-29193, AIF siRNA (m): sc-29194, AIF shRNA Plasmid (h): sc-29193-SH, AIF shRNA Plasmid (m): sc-29194-SH, AIF shRNA (h) Lentiviral Particles: sc-29193-V and AIF shRNA (m) Lentiviral Particles: sc-29194-V.

Molecular Weight of AIF: 57 kDa.

Positive Controls: CCRF-CEM cell lysate: sc-2225, Hep G2 cell lysate: sc-2227 or AML-193 whole cell lysate: sc-364182.

STORAGE

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

DATA





of normal mouse intestine frozen section showing

cytoplasmic and nuclear staining

AIF (H-300): sc-5586. Western blot analysis of AIF expression in AML-193 (A), CCRF-CEM (B), Hep G2 (C) and MOLT-4 (D) whole cell lysates.

SELECT PRODUCT CITATIONS

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- 3. Vorburger, S.A., et al. 2003. The mitochondrial apoptosis-inducing factor plays a role in E2F-1-induced apoptosis in human colon cancer cells. Ann. Surg. Oncol. 10: 314-322.
- Malina, H.Z., et al. 2003. Abnormal signalling of 14-3-3 proteins in cells with accumulated xanthurenic acid. Biochem. Biophys. Res. Commun. 310: 646-650.
- 5. Son, Y.O., et al. 2010. Cadmium induces intracellular Ca²⁺⁻ and H₂O₂-dependent apoptosis through JNK- and p53-mediated pathways in skin epidermal cell line. Toxicol. Sci. 113: 127-137.
- Kumar, A., et al. 2011. A novel parthenin analog exhibits anti-cancer activity: activation of apoptotic signaling events through robust NO formation in human leukemia HL-60 cells. Chem. Biol. Interact. 193: 204-215.
- Hojka-Osinska, A., et al. 2012. Combined treatment with fenretinide and indomethacin induces AIF-mediated, non-classical cell death in human acute T-cell leukemia Jurkat cells. Biochem. Biophys. Res. Commun. 419: 590-595.
- Khan, S., et al. 2012. A novel cyano derivative of 11-keto-β-boswellic acid causes apoptotic death by disrupting PI3K/AKT/Hsp-90 cascade, mitochondrial integrity, and other cell survival signaling events in HL-60 cells. Mol. Carcinog. 51: 679-695.
- Tardito, S., et al. 2012. Copper-dependent cytotoxicity of 8-hydroxyquinoline derivatives correlates with their hydrophobicity and does not require caspase activation. J. Med. Chem. 55: 10448-10459.