# SANTA CRUZ BIOTECHNOLOGY, INC.

# CIDE-A (4B9): sc-293289



BACKGROUND

The DNA fragmentation factor (DFF) is involved in the caspase-3 apoptotic pathway. DFF is composed of two subunits, DFF-45 (also designated ICAD, for inhibitor of CAD), and CPAN (caspase-activated nuclease), also designated CAD (caspase-activated deoxyribonuclease). CPAN is a DNase that is responsible for DNA degradation during apoptosis. CPAN is inhibited by DFF-45. Caspase-3 acts to dissociate CPAN from DFF-45, allowing CPAN to enter the nucleus and degrade DNA. CIDE-A and CIDE-B have been identified as proteins that share homology with the N-terminal region of DFF-45. Like CPAN, CIDE-A and CIDE-B promote cell death and DNA fragmentation and are inhibited by DFF-45.

# REFERENCES

- Liu, X., et al. 1997. DFF, a heterodimeric protein that functions downstream of caspase-3 to trigger DNA fragmentation during apoptosis. Cell 89: 175-184.
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- Halenbeck, R., et al. 1998. CPAN, a human nuclease regulated by the caspase-sensitive inhibitor DFF45. Curr. Biol. 8: 537-540.
- Sakahira, H., et al. 1998. Cleavage of CAD inhibitor in CAD activation and DNA degradation during apoptosis. Nature 391: 96-99.
- Inohara, N., et al. 1998. CIDE, a novel family of cell death activators with homology to the 45 kDa subunit of the DNA fragmentation factor. EMBO J. 17: 2526-2533.
- 7. Inohara, N., et al. 1999. Identification of regulatory and catalytic domains in the apoptosis nuclease DFF40/CAD. J. Biol. Chem. 274: 270-274.
- Lin, S.C., et al. 2004. CIDE-A, a novel link between brown adipose tissue and obesity. Trends Mol. Med. 10: 434-439.
- Novikova, S.I., et al. 2005. Cocaine-induced changes in the expression of apoptosis-related genes in the fetal mouse cerebral wall. Neurotoxicol. Teratol. 27: 3-14.

### CHROMOSOMAL LOCATION

Genetic locus: CIDEA (human) mapping to 18p11.21.

#### SOURCE

CIDE-A (4B9) is a mouse monoclonal antibody raised against amino acids 1-253 of CIDE-A of human origin.

# PRODUCT

Each vial contains 100  $\mu g$   $lgG_{2a}$  kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

### APPLICATIONS

CIDE-A (4B9) is recommended for detection of CIDE-A of human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2  $\mu$ g per 100-500  $\mu$ 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 CIDE-A siRNA (h): sc-37439, CIDE-A shRNA Plasmid (h): sc-37439-SH and CIDE-A shRNA (h) Lentiviral Particles: sc-37439-V.

Molecular Weight of CIDE-A: 26 kDa.

Positive Controls: CIDE-A transfected 293T whole cell lysate.

#### **RECOMMENDED SUPPORT REAGENTS**

To ensure optimal results, the following support reagents are recommended: 1) Western Blotting: use m-IgGκ BP-HRP: sc-516102 or m-IgGκ BP-HRP (Cruz Marker): sc-516102-CM (dilution range: 1:1000-1:10000), Cruz Marker™ Molecular Weight Standards: sc-2035, UltraCruz<sup>®</sup> Blocking Reagent: sc-516214 and Western Blotting Luminol Reagent: sc-2048. 2) Immunoprecipitation: use Protein A/G PLUS-Agarose: sc-2003 (0.5 ml agarose/2.0 ml).

#### DATA



CIDE-A (4B9): sc-293289. Western blot analysis of CIDE-A expression in non-transfected (**A**) and CIDE-A transfected (**B**) 293T whole cell lysates.

#### SELECT PRODUCT CITATIONS

 Duan, Y.N., et al. 2020. Diphyllin improves high-fat diet-induced obesity in mice through brown and beige adipocytes. Front. Endocrinol. 11: 592818.

human recombinant CIDE-A fusion protein

#### **STORAGE**

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

#### **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.