SANTA CRUZ BIOTECHNOLOGY, INC.

ECEL1 (P-15): sc-11339



BACKGROUND

ECEL1 (endothelin-converting enzyme-like 1, also designated XCE and DINE, damage-induced neuronal endopeptidase) is a member of the family of cellsurface zinc metallopeptidases. This family of metalloproteases includes endothelin-converting enzyme (ECE) and neutral endopeptidase (NEP). These peptidases are involved in the post-secretory processing and metabolism of neuropeptides and peptide hormones. Following neuronal damage, proteolytic activity of ECEL1 activates antioxidant enzymes suggesting a mechanism for how injured neurons protect themselves against death. Glycosylated ECEL1 is predominantly expressed in the central nervous system, including the spinal cord and medulla.

REFERENCES

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- Gomazkov, O.A. 1998. Endothelin-converting enzyme: its functional aspect. Biochemistry 63: 125-132.
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- Kiryu-Seo, S., et al. 2000. Damage-induced neuronal endopeptidase (DINE) is a unique metallo-peptidase expressed in response to neuronal damage and activates superoxide scavengers. Proc. Natl. Acad. Sci. USA 97: 4345-4350.

CHROMOSOMAL LOCATION

Genetic locus: ECEL1 (human) mapping to 2q37.1.

SOURCE

ECEL1 (P-15) is an affinity purified goat polyclonal antibody raised against a peptide mapping within an N-terminal cytoplasmic domain of ECEL1 of human origin.

STORAGE

Store at 4° C, **D0 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.

PRODUCT

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

Blocking peptide available for competition studies, sc-11339 P, (100 μ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

APPLICATIONS

ECEL1 (P-15) is recommended for detection of ECEL1 (also designated XCE or DINE) 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 solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

ECEL1 (P-15) is also recommended for detection of ECEL1 (also designated XCE or DINE) in additional species, including porcine.

Suitable for use as control antibody for ECEL1 siRNA (h): sc-106909, ECEL1 shRNA Plasmid (h): sc-106909-SH and ECEL1 shRNA (h) Lentiviral Particles: sc-106909-V.

Molecular Weight of ECEL1 cell surface: 105 kDa.

Molecular Weight of ECEL1 endoplasmic reticulum: 95 kDa.

Positive Controls: Jurkat whole cell lysate: sc-2204.

RECOMMENDED SECONDARY REAGENTS

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use donkey anti-goat IgG-HRP: sc-2020 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible donkey anti-goat IgG-HRP: sc-2033 (dilution range: 1:2000-1:5000), Cruz Marker™ Molecular Weight Standards: sc-2035, TBS Blotto A Blocking Reagent: sc-2333 and Western Blotting Luminol Reagent: sc-2048. 2) Immunoprecipitation: use Protein A/G PLUS-Agarose: sc-2003 (0.5 ml agarose/2.0 ml). 3) Immunofluorescence: use donkey anti-goat IgG-FITC: sc-2024 (dilution range: 1:100-1:400) or donkey anti-goat IgG-TR: sc-2783 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941.

DATA

132 K –			
90 K —	-	- ECEL1	
55 K –			
43 K –			
34 K –			

ECEL1 (P-15): sc-11339. Western blot analysis of ECEL1 expression in Jurkat whole cell lysate.

PROTOCOLS

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