## SANTA CRUZ BIOTECHNOLOGY, INC.

# X11β (B-5): sc-377060



#### BACKGROUND

The  $\beta$ -Amyloid precursor protein ( $\beta$ -APP) is a major constituent of the amyloid deposits in patients with Alzheimer's disease. The  $\beta$ -Amyloid precursor is known to interact with several proteins, including X11 and the G heterotrimetric protein APP-BP1. The neuronal, transmembrane protein X11 is known to bind to the  $\beta$ -Amyloid precursor protein via a phosphotyrosine binding (PTB) domain, reducing the secretion of cellular  $\beta$ -APP and slowing  $\beta$ -APP processing pathways. X11 binds specifically to the YENPTY motif, which is involved in the internalization of  $\beta$ -APP. Multiple splice varitents of X11 have been identified, including X11 $\alpha$  (also designated Mint 1), X11 $\beta$  (Mint 2) and X11 $\gamma$  (Mint 3).

### REFERENCES

- Borg, J.P., et al. 1996. The phosphotyrosine interaction domains of X11 and Fe65 bind to distinct sites on the YENPTY motif of amyloid precursor protein. Mol. Cell. Biol. 16: 6229-6241.
- Okamoto, M., et al. 1997. Mints, Munc18-interacting proteins in synaptic vesicle exocytosis. J. Biol. Chem. 272: 31459-31464.
- Zhang, Z., et al. 1997. Sequence-specific recognition of the internalization motif of the Alzheimer's amyloid precursor protein by the X11 PTB domain. EMBO J. 16: 6141-6150.
- 4. Russo, T., et al. 1998. Fe65 and the protein network centered around the cytosolic domain of the Alzheimer's  $\beta$ -Amyloid precursor protein. FEBS Lett. 434: 1-7.
- Borg, J.P., et al. 1998. The X11α protein slows cellular amyloid precursor protein processing and reduces Aβ40 and Aβ42 secretion. J. Biol. Chem. 273: 14761-14766.
- Sastre, M., et al. 1998. X11 interaction with β-Amyloid precursor protein modulates its cellular stabilization and reduces amyloid β-protein secretion. J. Biol. Chem. 273: 22351-22357.

#### **CHROMOSOMAL LOCATION**

Genetic locus: APBA2 (human) mapping to 15q13.1; Apba2 (mouse) mapping to 7 C.

#### SOURCE

X11 $\beta$  (B-5) is a mouse monoclonal antibody raised against amino acids 1-220 mapping at the N-terminus of X11 $\beta$  of mouse origin.

#### PRODUCT

Each vial contains 200  $\mu g$  IgM kappa light chain in 1.0 ml 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.

### **RESEARCH USE**

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

#### **APPLICATIONS**

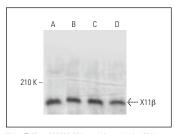
X11 $\beta$  (B-5) is recommended for detection of X11 $\beta$  of mouse, rat and human origin by Western Blotting (starting dilution 1:100, dilution range 1:100-1:1000), immunoprecipitation [1-2 µg per 100-500 µg of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500), immunohistochemistry (including paraffin-embedded sections) (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

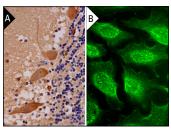
Suitable for use as control antibody for X11 $\beta$  siRNA (h): sc-36849, X11 $\beta$  siRNA (m): sc-36850, X11 $\beta$  shRNA Plasmid (h): sc-36849-SH, X11 $\beta$  shRNA Plasmid (m): sc-36850-SH, X11 $\beta$  shRNA (h) Lentiviral Particles: sc-36849-V and X11 $\beta$  shRNA (m) Lentiviral Particles: sc-36850-V.

Molecular Weight of X11<sub>B</sub>: 135 kDa.

Positive Controls: IMR-32 cell lysate: sc-2409, mouse brain extract: sc-2253 or rat cerebellum extract: sc-2398.

## DATA





 $X11\beta$  (B-5): sc-377060. Western blot analysis of  $X11\beta$  expression in IMR-32 whole cell lysate (A) and mouse brain (B), rat hippocampus (C) and rat cerebellum (D) tissue extracts.

X11 $\beta$  (B-5): sc-377060. Immunoperoxidase staining of formalin fixed, paraffin-embedded human cerebellum tissue showing cytoplasmic staining of Purkinje cells, cells in granular layer and cells in molecular layer (**A**). Immunofluorescence staining of methanol-fixed HeLa cells showing cytoplasmic and nuclear localization (**B**).

### SELECT PRODUCT CITATIONS

 Chung, Y., et al. 2020. Mint3 is dispensable for pancreatic and kidney functions in mice. Biochem. Biophys. Rep. 24: 100872.

#### PROTOCOLS

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