

### POLYCLONAL ANTIBODY

For research use only. Not for clinical diagnosis.

Catalog No. 74-108EX

## Anti-APP-C31 (C-terminal fragment of the caspase 3-cleaved APP) antibody, (ACT1)

#### **BACKGROUND**

The Alzheimer amyloid precursor protein (APP) is a transmembrane protein whose abnormal processing is associated with the pathogenesis of Alzheimer's disease. APP695 lacking the protease inhibitor domain is the predominant form in neuronal tissues. APP695 is cleaved by caspases into the 664-residue amino (N)-terminal fragment that lacks the carboxyl C-terminal 31-residues (APP□C31) and the 31-residues C-terminal fragment (APP-C31). Both fragments might be potent inducers of neuronal apoptosis. An antibody (named ACT1) against the N-terminus of caspase 3-generated APP C-terminal 31 aa of human APP695 (APP-C31) was raised in rabbit.

Primary antibodies **Product type** 

Rabbit Host Serum Source

Antiserum added with 0.05% sodium azide **Form** 

100µL Volume

Concentration

Synthetic peptide corresponding to the N-terminus of the caspase 3-generated **Immunogen** 

APP C-terminal 31 amino acids (aa 665-670 of human APP695)

#### Application notes

**Data Link** 

References

ref.3.

This antibody was used in

- 1. Western blotting (dilution: 1/3,000-1/1,000) 2. Immunocytechemistry (dilution: 1/1,000-1/500)
- 3. ELISA

These applications were confirmed in the laboratory of Prof. K, Yoshikawa of Osaka University. (ref.3).

Other applications have not been tested.

Optimal dilutions/concentrations should be determined by the end user.

UniProtKB/Swiss-Prot P05067 (A4\_HUMAN)

Reactive to human, mouse and rat. Specific to the N-terminal end of the caspase Reactivity

3-generated APP-C31

Shipped at 4°C and stored at -20°C

Storage

1. Kang HG et al (1987) "The precursor of Alzheimer's disease amyloid A4 protein resembles a cell-surface receptor." Nature 325: 33-736 PMID: 2881207

- 2. Selkoe DJ (1994) "Normal and abnormal biology of the beta-amyloid precursor protein." Annu Rev Neurosci 17: 489-517 PMID: 8210185
- 3. Nishimura I et al (2002) "Cell death induced by a caspase-cleaved transmembrane fragment of the Alzheimer amyloid precursor protein." Cell Death Differ 9: 199-208 PMID: 11840170
- 4. Nishimura I et al (2003) "Upregulation and antiapoptotic role of endogenous Alzheimer amyloid precursor protein in dorsal root ganglion neurons." Exp Cell Res **286**: 241-251 PMID: <u>12749853</u>

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Anti-APP-C31 (C-terminal fragment of the caspase 3-cleaved APP) antibody, rabbit serum (ACT1)

Related product

#<u>74-102EX</u> anti-Activated caspase3 antibody
#<u>74-104EX</u> anti-APP (C-terminus) antibody
#<u>74-106EX</u> anti-APP (N-terminus) antibody
#<u>74-110EX</u> anti-APPΔ31 (specific to C-terminal APPΔ31) antibody

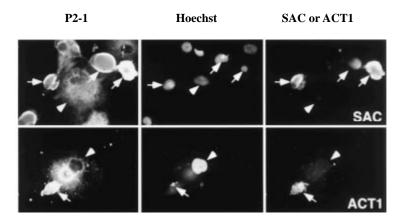


Fig.1 Immunocytochemistry for APPΔC31 and APP-C31: Generation of the caspase-cleaved fragments in NT2 neurons (neurally differentiated human NT2 embryonic carcinoma cells) overexpressing wild type APP (ref.3).

NT2 neurons were fixed 72 h after infection with adenovirus vector expressing wild-type APP and stained for the N-terminus of APP (P2-1, mouse monoclonal antibody), chromosomal DNA (Hoechst), the C-terminus of APPΔC31 (SAC) or the N-terminus of APP-C31 (ACT1). Most of wild-type APP-accumulating neurons with shrunken and fragmented nuclei contain SAC- and ACT1-immunoreactivities (arrows), but non-neuronal cells are hardly labeled with SAC and ACT1 (arrowheads).

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