

YBB250Ra01 50µg

5'-Nucleotidase, Ecto (NT5E)

Organism Species: Rattus norvegicus (Rat)

Instruction manual

FOR IN VITRO USE AND RESEARCH USE ONLY NOT FOR USE IN CLINICAL DIAGNOSTIC PROCEDURES

9th Edition (Revised in Jul, 2013)

[PROPERTIES]

Residues: Trp29~Phe550 (Accession # P21588),

with N-terminal His-Tag.

Host: E. coli

Subcellular Location: Cell membrane;

Lipid-anchor, GPI-anchor.

Purity: >95%

Endotoxin Level: <1.0EU per 1µg

(determined by the LAL method).

Formulation: Supplied as lyophilized form in PBS,

pH7.4, containing 5% sucrose, 0.01% sarcosyl.

Predicted isoelectric point: 6.2

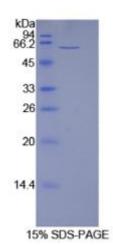
Predicted Molecular Mass: 59.6kDa

Applications: SDS-PAGE; WB; ELISA; IP.

(May be suitable for use in other assays to be determined by the end user.)

[USAGE]

Reconstitute in sterile PBS, pH7.2-pH7.4.





[STORAGE AND STABILITY]

Storage: Avoid repeated freeze/thaw cycles.

Store at 2-8°C for one month.

Aliquot and store at -80°C for 12 months.

Stability Test: The thermal stability is described by the loss rate of the target protein. The loss rate was determined by accelerated thermal degradation test, that is, incubate the protein at 37°C for 48h, and no obvious degradation and precipitation were observed. (Referring from China Biological Products Standard, which was calculated by the Arrhenius equation.) The loss of this protein is less than 5% within the expiration date under appropriate storage condition.

[SEQUENCES]

The target protein is fused with N-terminal His-Tag, its sequence is listed below. MGHHHHHHSGSEF-WE LTIMHTNDVH SRLEQTSDDS TKCLNASLCV GGVARLFTKV QQIRKEEPNV LLLDAGDQYQ GTIWFTVYKG LEVAHFMNLL GYDAMALGNH EFDNGVEGLI DPLLRNVKFP ILSANIKARG PLAPQISGLY LPYKVLSVGG EVVGIVGYTS KETPFLSNPG TNLVFEDEVT ALQPEVDKLK TLNVNKIIAL GHSGFEMDKL IAQKVRGVDV VVGGHTNTFL YTGNPPSKEV PAGKYPFIVT SDDGRKVPVV QAYAFGKYLG YLKVEFDDKG NVVTSYGNPI LLNSTIREDA AIKADINOWR IKLDNYSTQE LGRTIVYLNG SAQECRFREC NMGNLICDAM INNNLRHPDE MFWNHVSMCI VNGGGIRSPI DERNNGTITW ENLAAVLPFG GTFDLVQLKG STLKKAFEHS VHRYGQSTGE FLQVGGIHVV YDISRKPWDR VVQLKVLCTK CRVPIYEPLE MDKVYKVVLP SYLVNGGDGF QMIKDELLKH DSGDQDISVV SEYISKMKVI YPAVEGRIKF

[REFERENCES]

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- 2. Stanojevic I., et al. (2011) Int. J. Dev. Neurosci. 29:397-403.
- 3. Bjelobaba I., et al. (2011) J. Neurosci. Res. 89:862-873.
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