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SNX9 Protein (AA 1-595) (His tag)



Image



Go to Product page

Overview

| Quantity: | 1 mg |
|-------------------------------|--|
| Target: | SNX9 |
| Protein Characteristics: | AA 1-595 |
| Origin: | Human |
| Source: | Insect Cells |
| Protein Type: | Recombinant |
| Purification tag / Conjugate: | This SNX9 protein is labelled with His tag. |
| Application: | Western Blotting (WB), SDS-PAGE (SDS), ELISA, Crystallization (Crys) |

Product Details

Sequence:

MATKARVMYD FAAEPGNNEL TVNEGEIITI TNPDVGGGWL EGRNIKGERG LVPTDYVEIL
PSDGKDQFSC GNSVADQAFL DSLSASTAQA SSSAASNNHQ VGSGNDPWSA WSASKSGNWE
SSEGWGAQPE GAGAQRNTNT PNNWDTAFGH PQAYQGPATG DDDDWDEDWD GPKSSSYFKD
SESADAGGAQ RGNSRASSSS MKIPLNKFPG FAKPGTEQYL LAKQLAKPKE KIPIIVGDYG
PMWVYPTSTF DCVVADPRKG SKMYGLKSYI EYQLTPTNTN RSVNHRYKHF DWLYERLLVK
FGSAIPIPSL PDKQVTGRFE EEFIKMRMER LQAWMTRMCR HPVISESEVF QQFLNFRDEK
EWKTGKRKAE RDELAGVMIF STMEPEAPDL DLVEIEQKCE AVGKFTKAMD DGVKELLTVG
QEHWKRCTGP LPKEYQKIGK ALQSLATVFS SSGYQGETDL NDAITEAGKT YEEIASLVAE
QPKKDLHFLM ECNHEYKGFL GCFPDIIGTH KGAIEKVKES DKLVATSKIT LQDKQNMVKR
VSIMSYALQA EMNHFHSNRI YDYNSVIRLY LEQQVQFYET IAEKLRQALS RFPVM

Sequence without tag. Tag location is at the discretion of the manufacturer. If you have a special request, please contact us.

Characteristics:

- Made in Germany from design to production by highly experienced protein experts.
- Human SNX9 Protein (raised in Insect Cells) purified by multi-step, protein-specific process to ensure crystallization grade.
- State-of-the-art algorithm used for plasmid design (Gene synthesis).

This protein is a made to order protein and will be made for the first time for your order. Our experts in the lab will ensure that you receive a correctly folded protein.

The big advantage of ordering our made-to-order proteins in comparison to ordering custom made proteins from other companies is that there is no financial obligation in case the protein cannot be expressed or purified.

In the unlikely event that the protein cannot be expressed or purified we do not charge anything (other companies might charge you for any performed steps in the expression process for custom-made proteins, e.g. fees might apply for the expression plasmid, the first expression experiments or purification optimization).

When you order this made-to-order protein you will only pay upon receival of the correctly folded protein. With no financial risk on your end you can rest assured that our experienced protein experts will do everything to make sure that you receive the protein you ordered.

The concentration of our recombinant proteins is measured using the absorbance at 280nm.

The protein's absorbance will be measured in several dilutions and is measured against its specific reference buffer.

The concentration of the protein is calculated using its specific absorption coefficient. We use the Expasy's protparam tool to determine the absorption coefficient of each protein.

Purification:

Two step purification of proteins expressed in baculovirus infected SF9 insect cells:

- 1. In a first purification step, the protein is purified from the cleared cell lysate using three different His-tag capture materials: high yield, EDTA resistant, or DTT resistant. Eluate fractions are analyzed by SDS-PAGE.
- 2. Protein containing fractions of the best purification are subjected to second purification step through size exclusion chromatography. Eluate fractions are analyzed by SDS-PAGE and Western blot.

Purity:

>95 % as determined by SDS PAGE, Size Exclusion Chromatography and Western Blot.

Sterility:

0.22 µm filtered

Endotoxin Level:

Protein is endotoxin free.

Grade:

Crystallography grade

Target Details

| Target: | SNX9 |
|---------------------|---|
| Alternative Name: | SNX9 (SNX9 Products) |
| Background: | Involved in endocytosis and intracellular vesicle trafficking, both during interphase and at the |
| | end of mitosis. Required for efficient progress through mitosis and cytokinesis. Required for |
| | normal formation of the cleavage furrow at the end of mitosis. Plays a role in endocytosis via |
| | clathrin-coated pits, but also clathrin-independent, actin-dependent fluid-phase endocytosis. |
| | Plays a role in macropinocytosis. Promotes internalization of TNFR. Promotes degradation of |
| | EGFR after EGF signaling. Stimulates the GTPase activity of DNM1. Promotes DNM1 |
| | oligomerization. Promotes activation of the Arp2/3 complex by WASL, and thereby plays a role |
| | in the reorganization of the F-actin cytoskeleton. Binds to membranes enriched in |
| | phosphatidylinositol 4,5-bisphosphate and promotes membrane tubulation. Has lower affinity |
| | for membranes enriched in phosphatidylinositol 3-phosphate. |
| | {ECO:0000269 PubMed:11799118, ECO:0000269 PubMed:12952949, |
| | ECO:0000269 PubMed:15703209, ECO:0000269 PubMed:17609109, |
| | ECO:0000269 PubMed:17948057, ECO:0000269 PubMed:18388313, |
| | ECO:0000269 PubMed:20427313, ECO:0000269 PubMed:21048941, |
| | ECO:0000269 PubMed:22718350}. |
| Molecular Weight: | 67.5 kDa Including tag. |
| UniProt: | Q9Y5X1 |
| Application Details | |
| Application Notes: | In addition to the applications listed above we expect the protein to work for functional studies |
| | as well. As the protein has not been tested for functional studies yet we cannot offer a gurante |
| | though. |
| Comment: | In cases in which it is highly likely that the recombinant protein with the default tag will be |
| | insoluble our protein lab may suggest a higher molecular weight tag (e.g. GST-tag) instead to |
| | increase solubility. We will discuss all possible options with you in detail to assure that you |
| | receive your protein of interest. |
| Restrictions: | For Research Use only |
| Handling | |
| | |

Handling

| Buffer: | 100 mM NaCL, 20 mM Hepes, 10% glycerol. pH value is at the discretion of the manufacturer. |
|------------------|--|
| Handling Advice: | Avoid repeated freeze-thaw cycles. |
| Storage: | -80 °C |
| Storage Comment: | Store at -80°C. |
| Expiry Date: | Unlimited (if stored properly) |
| | |

Images

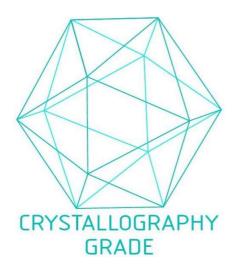


Image 1. "Crystallography Grade" protein due to multi-step, protein-specific purification process