antibodies

## Datasheet for ABIN3137430 SYT7 Protein (AA 1-403) (rho-1D4 tag)



1 Image



## Overview

| Quantity:                     | 1 mg   |
|-------------------------------|--|
| Target:                       | SYT7   |
| Protein Characteristics:      | AA 1-403   |
| Origin:                       | Mouse  |
| Source:                       | Insect Cells   |
| Protein Type:                 | Recombinant  |
| Purification tag / Conjugate: | This SYT7 protein is labelled with rho-1D4 tag.                      |
| Application:                  | ELISA, Western Blotting (WB), Crystallization (Crys), SDS-PAGE (SDS) |

## Product Details

| Sequence:        | MYRDPEAASP GAPTRDVLLV SAIITVSLSV TIVLCGLCHW CQRKLGKRYK NSLETVGTPD   |
|------------------|---|
|                  | SGRGRGEKKA IKLPAGGKAV NTAPVPGQTP HDESDRRTET RSSVSDLVNS LTSEMLMLSP   |
|                  | GSEEDEAHEG CSRENLGRIQ FSVGYNFQES TLTVKVMKAQ ELPAKDFSGT SDPFVKIYLL   |
|                  | PDKKHKLETK VKRKNLNPHW NETFLFEGFP YEKVVQRVLY LQVLDYDRFS RNDPIGEVSI   |
|                  | PLNKVDLTQM QTFWKDLKPC SDGSGSRGEL LLSLCYNPSA NSIIVNIIKA RNLKAMDIGG   |
|                  | TSDPYVKVWL MYKDKRVEKK KTVTKKRNLN PIFNESFAFD IPTEKLRETT IIITVMDKDK   |
|                  | LSRNDVIGKI YLSWKSGPGE VKHWKDMIAR PRQPVAQWHQ LKA   |
|                  | Sequence without tag. Tag location is at the discretion of the manufacturer. If you have a  |
|                  | special request, please contact us.   |
| Characteristics: | <ul> <li>Made in Germany - from design to production - by highly experienced protein experts.</li> <li>Mouse Syt7 Protein (raised in Insect Cells) purified by multi-step, protein-specific process to ensure crystallization grade.</li> </ul> |

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|                  | 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:    | Three step purification of membrane proteins expressed in baculovirus infected SF9 insect   |
|                  | cells:  |
|                  | 1. Membrane proteins are fractioned by ultracentrifugation and subsequently solubilized with different detergents (detergent screen). Samples are analyzed by Western blot. |
|                  | 2. The best performing detergent is used for solubilization and the proteins are purified via their   |
|                  | rho1D4 tag via two rho1D4 antibody columns: one DTT resistant, the other one not. Eluate  |
|                  | fractions are analyzed by Western blot.<br>3. Protein containing fractions of the best purification are subjected to second purification step                               |
|                  | through size exclusion chromatograph. 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   |

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| Target Details |
|----------------|
|----------------|

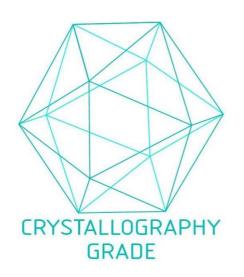
| Target:           | SYT7  |
|-------------------|---|
| Alternative Name: | Syt7 (SYT7 Products)  |
| Background:       | Ca(2+) sensor involved in Ca(2+)-dependent exocytosis of secretory and synaptic vesicles            |
|                   | through Ca(2+) and phospholipid binding to the C2 domain. Ca(2+) induces binding of the C2-         |
|                   | domains to phospholipid membranes and to assembled SNARE-complexes, both actions                    |
|                   | contribute to triggering exocytosis. SYT7 binds Ca(2+) with high affinity and slow kinetics         |
|                   | compared to other synaptotagmins (PubMed:26738595). Involved in Ca(2+)-triggered                    |
|                   | lysosomal exocytosis, a major component of the plasma membrane repair (By similarity).              |
|                   | Ca(2+)-regulated delivery of lysosomal membranes to the cell surface is also involved in the        |
|                   | phagocytic uptake of particles by macrophages (PubMed:16982801, PubMed:21041449).                   |
|                   | Ca(2+)-triggered lysosomal exocytosis also plays a role in bone remodeling by regulating            |
|                   | secretory pathways in osteoclasts and osteoblasts (PubMed:18539119). Involved in cholester          |
|                   | transport from lysosome to peroxisome by promoting membrane contacts between lysosome               |
|                   | and peroxisomes: probably acts by promoting vesicle fusion by binding phosphatidylinositol-         |
|                   | 4,5-bisphosphate on peroxisomal membranes (PubMed:25860611). Acts as a key mediator of              |
|                   | synaptic facilitation, a process also named short-term synaptic potentiation: synaptic              |
|                   | facilitation takes place at synapses with a low initial release probability and is caused by influx |
|                   | of Ca(2+) into the axon terminal after spike generation, increasing the release probability of      |
|                   | neurotransmitters (PubMed:24569478, PubMed:26738595). Probably mediates synaptic                    |
|                   | facilitation by directly increasing the probability of release (PubMed:26738595). May also          |
|                   | contribute to synaptic facilitation by regulating synaptic vesicle replenishment, a process         |
|                   | required to ensure that synaptic vesicles are ready for the arrival of the next action potential:   |
|                   | SYT7 is required for synaptic vesicle replenishment by acting as a sensor for Ca(2+) and by         |
|                   | forming a complex with calmodulin (PubMed:24569478). Also acts as a regulator of Ca(2+)-            |
|                   | dependent insulin and glucagon secretion in beta-cells (PubMed:18308938,                            |
|                   | PubMed:19171650). Triggers exocytosis by promoting fusion pore opening and fusion pore              |
|                   | expansion in chromaffin cells (PubMed:20956309). Also regulates the secretion of some non-          |
|                   | synaptic secretory granules of specialized cells (By similarity).                                   |
|                   | {EC0:0000250 UniProtKB:Q62747, EC0:0000269 PubMed:16982801,   |
|                   | ECO:0000269 PubMed:18308938, ECO:0000269 PubMed:18539119,   |
|                   | ECO:0000269 PubMed:19171650, ECO:0000269 PubMed:20956309,   |
|                   | ECO:0000269 PubMed:21041449, ECO:0000269 PubMed:24569478,   |
|                   | ECO:0000269 PubMed:25860611, ECO:0000269 PubMed:26738595}.  |
| Molecular Weight: | 46.6 kDa Including tag.   |

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| Target Details      |   |
|---------------------|---|
| UniProt:            | Q9R0N7  |
| Pathways:           | Synaptic Vesicle Exocytosis   |
| Application Details |   |
| Application Notes:  | In addition to the applications listed above we expect the protein to work for functional studies<br>as well. As the protein has not been tested for functional studies yet we cannot offer a gurantee<br>though.   |
| Comment:            | Protein has not been tested for activity yet. 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            |   |
| Format:             | Liquid  |
| 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.   |

Images

Expiry Date:



Unlimited (if stored properly)

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

protein-specific purification process

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