abcam

Product datasheet

Glycogen synthase 1/GYS1 peptide ab204860

Description

Product name Glycogen synthase 1/GYS1 peptide

Purity > 95 % HPLC.

Animal free No

Nature Synthetic

Sequence YRRAAVPPSPSLSRHSSPHQ(pS)EDEEE

Amino acids 636 to 661

Specifications

Our Abpromise guarantee covers the use of ab204860 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Applications Functional Studies

HPLC

Form Lyophilized

Additional notes ab204860 (Glycogen synthase 1/GYS1 peptide) can be utilized as a substrate for the following

active protein kinases:

<u>ab60863</u> (Active human GSK3 beta full length protein) <u>ab63193</u> (Active human GSK3 beta full length protein)

This product was previously labelled as Glycogen synthase 1

Preparation and Storage

Stability and Storage Shipped at 4°C. Store at -20°C. Avoid freeze / thaw cycle.

ReconstitutionDilute peptide in 25mM Tris-HCl, pH 7.5 solution to a final concentration of 1 mg/ml. For optimal

storage, aliquot diluted product into smaller quantities and store at recommended temperature.

General Info

Function Transfers the glycosyl residue from UDP-Glc to the non-reducing end of alpha-1,4-glucan.

Pathway Glycan biosynthesis; glycogen biosynthesis.

Involvement in disease Defects in GYS1 are the cause of muscle glycogen storage disease type 0 (GSD0b)

[MIM:611556]; also known as muscle glycogen synthase deficiency. GSD0b is a metabolic disorder characterized by fasting hypoglycemia presenting in infancy or early childhood. The role of muscle glycogen is to provide critical energy during bursts of activity and sustained muscle

work.

Sequence similarities Belongs to the glycosyltransferase 3 family.

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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