

# Glucosidase II $\beta$ (A-7): sc-514870

## BACKGROUND

Trimming of glucoses from N-linked core glycans on newly synthesized glycoproteins occurs sequentially through the action of Glucosidases I and II in the endoplasmic reticulum (ER). Glucosidase II is an ER-localized enzyme that contains  $\alpha$  and  $\beta$  subunits (Glucosidase II $\alpha$  and Glucosidase II $\beta$ ) which form a defined heterodimeric complex. Glucosidase II $\alpha$  is the catalytic core of the enzyme and can function independently of the  $\beta$  subunit. The sequence of Glucosidase II $\beta$  encodes protein rich in glutamic and aspartic acid with a putative ER retention signal (HDEL) at the C-terminus. The phosphorylated form of Glucosidase II $\beta$  is localized in the plasma membrane and is highly expressed in FGF-stimulated fibroblasts and epidermal carcinoma cells. Glucosidase II $\beta$  was first purified from a human carcinoma cell line as a potential substrate for protein kinase C. Through the HDEL signal at the C-terminus, Glucosidase II $\beta$  retains the complete complex in the ER.

## REFERENCES

1. Shailubhai, K., Pratta M.A. and Vijay, I.K. 1987. Purification and characterization of Glucosidase I involved in N-linked glycoprotein processing in bovine mammary gland. *Biochem. J.* 247: 555-562.
2. Saxena, S., Shailubhai, K., Dong-Yu, B. and Vijay, I.K. 1987. Purification and characterization of Glucosidase II involved in N-linked glycoprotein processing in bovine mammary gland. *Biochem. J.* 247: 563-570.
3. Trombetta, E.S., Simons, J.F. and Helenius, A. 1996. Endoplasmic reticulum Glucosidase II is composed of a catalytic subunit, conserved from yeast to mammals, and a tightly bound noncatalytic HDEL-containing subunit. *J. Biol. Chem.* 271: 27509-27516.
4. Trembl, K., Meimaroglou, D., Hentges, A. and Bause, E. 2000. The  $\alpha$ - and  $\beta$ -subunits are required for expression of catalytic activity in the heterodimeric Glucosidase II complex from human liver. *Glycobiology* 10: 493-502.
5. Trombetta, E.S., Fleming, K.G. and Helenius, A. 2001. Quaternary and domain structure of glycoprotein processing Glucosidase II. *Biochemistry* 40: 10717-10722.

## CHROMOSOMAL LOCATION

Genetic locus: PRKCSH (human) mapping to 19p13.2; PrkcsH (mouse) mapping to 9 A3.

## SOURCE

Glucosidase II $\beta$  (A-7) is a mouse monoclonal antibody specific for an epitope mapping between amino acids 383-420 within an internal region of Glucosidase II $\beta$  of human origin.

## PRODUCT

Each vial contains 200  $\mu$ g IgG<sub>2a</sub> kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-514870 P, (100  $\mu$ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% stabilizer protein).

## APPLICATIONS

Glucosidase II $\beta$  (A-7) is recommended for detection of Glucosidase II $\beta$  of mouse, rat and human origin by Western Blotting (starting dilution 1:100, dilution range 1:100-1:1000), immunoprecipitation [1-2  $\mu$ g per 100-500  $\mu$ g of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

Suitable for use as control antibody for Glucosidase II $\beta$  siRNA (h): sc-29598, Glucosidase II $\beta$  siRNA (m): sc-29599, Glucosidase II $\beta$  shRNA Plasmid (h): sc-29598-SH, Glucosidase II $\beta$  shRNA Plasmid (m): sc-29599-SH, Glucosidase II $\beta$  shRNA (h) Lentiviral Particles: sc-29598-V and Glucosidase II $\beta$  shRNA (m) Lentiviral Particles: sc-29599-V.

Molecular Weight of Glucosidase II $\beta$ : 80-90 kDa.

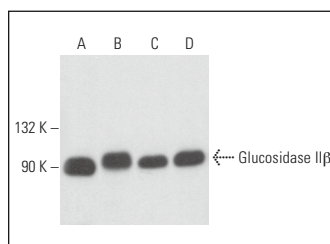
Positive Controls: A-10 cell lysate: sc-3806, K-562 whole cell lysate: sc-2203 or NIH/3T3 whole cell lysate: sc-2210.

## RECOMMENDED SUPPORT REAGENTS

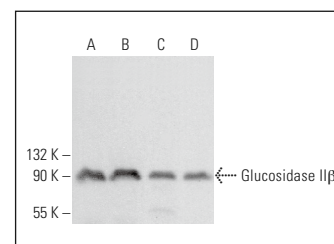
To ensure optimal results, the following support reagents are recommended:

- 1) Western Blotting: use m-IgG $\kappa$  BP-HRP: sc-516102 or m-IgG $\kappa$  BP-HRP (Cruz Marker): sc-516102-CM (dilution range: 1:1000-1:10000), Cruz Marker™ Molecular Weight Standards: sc-2035, UltraCruz® Blocking Reagent: sc-516214 and Western Blotting Luminol Reagent: sc-2048.
- 2) Immunoprecipitation: use Protein A/G PLUS-Agarose: sc-2003 (0.5 ml agarose/2.0 ml).
- 3) Immunofluorescence: use m-IgG $\kappa$  BP-FITC: sc-516140 or m-IgG $\kappa$  BP-PE: sc-516141 (dilution range: 1:50-1:200) with UltraCruz® Mounting Medium: sc-24941 or UltraCruz® Hard-set Mounting Medium: sc-359850.

## DATA



Glucosidase II $\beta$  (A-7): sc-514870. Western blot analysis of Glucosidase II $\beta$  expression in K-562 (A), NIH/3T3 (B), RAW 264.7 (C) and A-10 (D) whole cell lysates.



Glucosidase II $\beta$  (A-7): sc-514870. Western blot analysis of Glucosidase II $\beta$  expression in K-562 (A), HEL 92.1.7 (B), M1 (C) and Daudi (D) whole cell lysates.

## STORAGE

Store at 4° C, \*\*DO NOT FREEZE\*\*. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

## RESEARCH USE

For research use only, not for use in diagnostic procedures.

## PROTOCOLS

See our web site at [www.scbt.com](http://www.scbt.com) for detailed protocols and support products.