β-glucosidase (N-17): sc-30844



The Power to Question

BACKGROUND

β-glucosidase is a predominantly liver enzyme which efficiently hydrolyzes β-D-glucoside and β-D-galactoside. Defects in β-glucosidase cause Gaucher disease, an inherited condition distinguished by the accumulation of glucosylceramide within the cells of the reticuloendothelial system. β-glucosidase is used in enzyme replacement treatment aimed at treating Gaucher disease. The absorption of dietary flavonoid glycosides in humans involves a critical deglycosylation step that is mediated by epithelial β -glucosidases.

REFERENCES

- Overkleeft, H.S., et al. 1998. Generation of specific deoxynojirimycin-type inhibitors of the non-lysosomal glucosylceramidase. J. Biol. Chem. 273: 26522-26527.
- de Graaf, M., et al. 2001. Cloning and characterization of human liver cytosolic β-glycosidase. Biochem. J. 356: 907-910.

CHROMOSOMAL LOCATION

Genetic locus: GBA (human) mapping to 1q22; Gba (mouse) mapping to 3 F1.

SOURCE

 $\beta\text{-glucosidase}$ (N-17) is an affinity purified goat polyclonal antibody raised against a peptide mapping near the N-terminus of $\beta\text{-glucosidase}$ of human origin.

PRODUCT

Each vial contains 200 μg lgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-30844 P, (100 μ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

APPLICATIONS

β-glucosidase (N-17) is recommended for detection of β-glucosidase of mouse, rat and human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1–2 μg per 100–500 μg 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).

 β -glucosidase (N-17) is also recommended for detection of β -glucosidase in additional species, including equine, canine, bovine and porcine.

Suitable for use as control antibody for β -glucosidase siRNA (h): sc-44904, β -glucosidase siRNA (m): sc-44905, β -glucosidase shRNA Plasmid (h): sc-44904-SH, β -glucosidase shRNA Plasmid (m): sc-44905-SH, β -glucosidase shRNA (h) Lentiviral Particles: sc-44904-V and β -glucosidase shRNA (m) Lentiviral Particles: sc-44905-V.

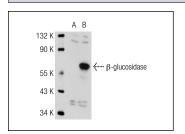
Molecular Weight of β-glucosidase: 57 kDa.

Positive Controls: β -glucosidase (h): 293T Lysate: sc-110483, MCF7 whole cell lysate: sc-2206 or mouse liver extract: sc-2256.

RECOMMENDED SECONDARY REAGENTS

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use donkey anti-goat IgG-HRP: sc-2020 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible donkey anti-goat IgG-HRP: sc-2033 (dilution range: 1:2000-1:5000), Cruz Marker™ Molecular Weight Standards: sc-2035, TBS Blotto A Blocking Reagent: sc-2333 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 donkey anti-goat IgG-FITC: sc-2024 (dilution range: 1:100-1:400) or donkey anti-goat IgG-TR: sc-2783 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941.

DATA



β-glucosidase (N-17): sc-30844. Western blot analysis of β-glucosidase expression in non-transfected: sc-117752 (\mathbf{A}) and human β-glucosidase transfected sc-110483 (\mathbf{B}) 293T 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 or our catalog for detailed protocols and support products.



Try β -glucosidase (B-6): sc-166407 or β -glucosidase (C-2): sc-365745, our highly recommended monoclonal alternatives to β -glucosidase (N-17).

Santa Cruz Biotechnology, Inc. 1.800.457.3801 831.457.3800 fax 831.457.3801 Europe +00800 4573 8000 49 6221 4503 0 www.scbt.com