

β -glucuronidase (H-300): sc-25827

BACKGROUND

The enzyme β -glucuronidase catalyzes the conversion of β -D-glucuronoside and water to an alcohol and D-glucuronate. Deficiency of β -glucuronidase is the cause of the human lysosomal storage disorder mucopolysaccharidosis type VII (MPS VII). Specifically, two residues appear important for catalytic activity: Glu 451 and Glu 540. Mutations at these sites affect the overall structure of the protein, which normally consists of a homotetramer with each promoter including a jelly roll barrel, an immunoglobulin constant domain and a TIM barrel. Regulation of β -glucuronidase activity may play a role in tumorigenesis and the invasiveness of a number of cancers, and is also an important factor in the development of functional prodrugs that require the cleavage of an active cytostatic by endogenous enzymes for antitumor activity.

REFERENCES

1. Himeno Mnishimura, Y., et al. 1976. Purification and characterization of microsomal and lysosomal β -glucuronidase from rat liver by use of immunofluorescence chromatography. *Eur. J. Biochem.* 70: 349-359.
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3. Varma, R., et al. 1983. β -glucuronidase in sera of patients with epileptic seizure activity, diabetes and some other disease states. *Neurosci. Lett.* 39: 105-111.
4. Guise, K.S., et al. 1985. Isolation and expression in *Escherichia coli* of a cDNA clone encoding human β -glucuronidase. *Gene* 34: 105-110.
5. Watson, G., et al. 1985. Properties of rat and mouse β -glucuronidase mRNA and cDNA, including evidence for sequence polymorphism and genetic regulation of mRNA levels. *Gene* 36: 15-25.
6. Jain, S., et al. 1996. Structure of human β -glucuronidase reveals candidate lysosomal targeting and active-site motifs. *Nat. Struct. Biol.* 3: 375-381.
7. Vervoort, R., et al. 1998. Low β -glucuronidase enzyme activity and mutations in the human β -glucuronidase gene in mild mucopolysaccharidosis type VII, pseudodeficiency and a heterozygote. *Hum. Genet.* 102: 69-78.
8. Kurokawa, H., et al. 2003. Heparanase and tumor invasion patterns in human oral squamous cell carcinomaxenografts. *Cancer Sci.* 94: 277-285.
9. Grube, M., et al. 2003. Verapamil regulates activity and mRNA-expression of human β -glucuronidase in Hep G2 cells. *Naunyn Schmiedebergs Arch. Pharmacol.* 368: 463-469.

CHROMOSOMAL LOCATION

Genetic locus: GUSB (human) mapping to 7q11.21; Gusb (mouse) mapping to 5 G1.3.

SOURCE

β -glucuronidase (H-300) is a rabbit polyclonal antibody raised against amino acids 352-651 mapping at the C-terminus of β -glucuronidase of human origin.

PRODUCT

Each vial contains 200 μ g IgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

APPLICATIONS

β -glucuronidase (H-300) is recommended for detection of β -glucuronidase of mouse, rat and human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2 μ g per 100-500 μ 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 β -glucuronidase siRNA (h): sc-44458, β -glucuronidase siRNA (m): sc-44459, β -glucuronidase shRNA Plasmid (h): sc-44458-SH, β -glucuronidase shRNA Plasmid (m): sc-44459-SH, β -glucuronidase shRNA (h) Lentiviral Particles: sc-44458-V and β -glucuronidase shRNA (m) Lentiviral Particles: sc-44459-V.

Molecular Weight of β -glucuronidase: 82 kDa.

Positive Controls: HL-60 whole cell lysate: sc-2209 or Hep G2 cell lysate: sc-2227.

RECOMMENDED SECONDARY REAGENTS

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use goat anti-rabbit IgG-HRP: sc-2004 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible goat anti-rabbit IgG-HRP: sc-2030 (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 goat anti-rabbit IgG-FITC: sc-2012 (dilution range: 1:100-1:400) or goat anti-rabbit IgG-TR: sc-2780 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941.

SELECT PRODUCT CITATIONS

1. Menendez, C., et al. 2011. Vascular deconjugation of quercetin glucuronide: the flavonoid paradox revealed? *Mol. Nutr. Food Res.* 55: 1780-1790.

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.


 MONOS
Satisfaction
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Try **β -glucuronidase (E-11): sc-374629**, our highly recommended monoclonal alternative to β -glucuronidase (H-300).