

# $\beta$ -1,4-Gal-T7 (P-12): sc-160112

## BACKGROUND

$\beta$ -1,4-galactosyltransferases ( $\beta$ -1,4-Gal-T) are type II membrane-bound glycoproteins that are substrate-specific and function to transfer galactose in a  $\beta$ -1,4 linkage to an acceptor sugar. There are seven members of the  $\beta$ -1,4-Gal-T family, all of which are directed to the Golgi apparatus through a hydrophobic sequence at the N-terminus.  $\beta$ -1,4-Gal-T7, also known as B4GALT7 or XGALT1, is a 327 amino acid single-pass type II membrane protein that is expressed at high levels in heart, pancreas and liver.  $\beta$ -1,4-Gal-T7 uses manganese to catalyze the UDP-dependent biosynthesis of glycosphingolipids. The gene encoding  $\beta$ -1,4-Gal-T7 is mutated in Ehlers-Danlos syndrome progeroid type (EDSP), a variant form of Ehlers-Danlos syndrome characterized by progeroid facies, mild mental retardation, short stature, skin hyperextensibility, moderate skin fragility, joint hypermobility principally in digits.

## REFERENCES

- Pal, R., Hoke, G.M. and Sarngadharan, M.G. 1989. Role of oligosaccharides in the processing and maturation of envelope glycoproteins of human immunodeficiency virus type 1. *Proc. Natl. Acad. Sci. USA* 86: 3384-3388.
- Lo, N.W., Shaper, J.H., Pevsner, J. and Shaper, N.L. 1998. The expanding  $\beta$  4-galactosyltransferase gene family: messages from the databanks. *Glycobiology* 8: 517-526.
- Amado, M., Almeida, R., Schwientek, T. and Clausen, H. 1999. Identification and characterization of large galactosyltransferase gene families: galactosyltransferases for all functions. *Biochim. Biophys. Acta* 1473: 35-53.
- Kuroiwa, A., Matsuda, Y., Okajima, T. and Furukawa, K. 2000. Assignment of human xylosylprotein  $\beta$ -1,4-galactosyltransferase gene (B4GALT7) to human chromosome 5q35.2 $\rightarrow$ q35.3 by *in situ* hybridization. *Cytogenet. Cell Genet.* 89: 8-9.
- Faiyaz-UI-Haque, M., Zaidi, S.H., Al-Ali, M., Al-Mureikhi, M.S., Kennedy, S., Al-Thani, G., Tsui, L.C. and Teebi, A.S. 2004. A novel missense mutation in the galactosyltransferase-I (B4GALT7) gene in a family exhibiting facioskeletal anomalies and Ehlers-Danlos syndrome resembling the progeroid type. *Am. J. Med. Genet. A* 128A: 39-45.
- Götte, M. and Kresse, H. 2005. Defective glycosaminoglycan substitution of decorin in a patient with progeroid syndrome is a direct consequence of two point mutations in the galactosyltransferase I ( $\beta$ 4GalT-7) gene. *Biochem. Genet.* 43: 65-77.
- Seidler, D.G., Faiyaz-UI-Haque, M., Hansen, U., Yip, G.W., Zaidi, S.H., Teebi, A.S., Kiesel, L. and Götte, M. 2006. Defective glycosylation of decorin and biglycan, altered collagen structure, and abnormal phenotype of the skin fibroblasts of an Ehlers-Danlos syndrome patient carrying the novel Arg270Cys substitution in galactosyltransferase I ( $\beta$ 4GalT-7). *J. Mol. Med.* 84: 583-594.
- Daligault, F., Rahuel-Clermont, S., Gulberti, S., Cung, M.T., Branlant, G., Netter, P., Magdalou, J. and Lattard, V. 2009. Thermodynamic insights into the structural basis governing the donor substrate recognition by human  $\beta$ 1,4-galactosyltransferase 7. *Biochem. J.* 418: 605-614.

## RESEARCH USE

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

## CHROMOSOMAL LOCATION

Genetic locus: B4GALT7 (human) mapping to 5q35.3; B4galt7 (mouse) mapping to 13 B1.

## SOURCE

$\beta$ -1,4-Gal-T7 (P-12) is an affinity purified goat polyclonal antibody raised against a peptide mapping within an internal region of  $\beta$ -1,4-Gal-T7 of human origin.

## PRODUCT

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

Blocking peptide available for competition studies, sc-160112 P, (100  $\mu$ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

## APPLICATIONS

$\beta$ -1,4-Gal-T7 (P-12) is recommended for detection of  $\beta$ -1,4-Gal-T7 of mouse, rat and human origin by Western Blotting (starting dilution 1:200, 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); non cross-reactive with other  $\beta$ -1,4-Gal-T family members.

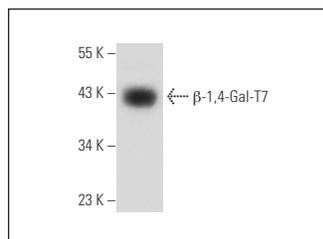
$\beta$ -1,4-Gal-T7 (P-12) is also recommended for detection of  $\beta$ -1,4-Gal-T7 in additional species, including equine, canine, bovine, porcine and avian.

Suitable for use as control antibody for  $\beta$ -1,4-Gal-T7 siRNA (h): sc-91927,  $\beta$ -1,4-Gal-T7 siRNA (m): sc-108227,  $\beta$ -1,4-Gal-T7 shRNA Plasmid (h): sc-91927-SH,  $\beta$ -1,4-Gal-T7 shRNA Plasmid (m): sc-108227-SH,  $\beta$ -1,4-Gal-T7 shRNA (h) Lentiviral Particles: sc-91927-V and  $\beta$ -1,4-Gal-T7 shRNA (m) Lentiviral Particles: sc-108227-V.

Molecular Weight of  $\beta$ -1,4-Gal-T7: 37 kDa.

Positive Controls: mouse skeletal muscle tissue extract: sc-364250.

## DATA



$\beta$ -1,4-Gal-T7 (P-12): sc-160112. Western blot analysis of  $\beta$ -1,4-Gal-T7 expression in mouse skeletal muscle tissue extract.

## STORAGE

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