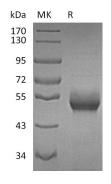
Catalog #: PHH2132



## **Summary**

Name	B4GALT1			
Purity	Greater than 95% as determined by reducing SDS-PAGE			
Endotoxin level	<1 EU/µg as determined by LAL test.			
Construction Accession #	Recombinant Human Beta-1,4-galactosyltransferase 1 is produced by our Mammalian expression system and the target gene encoding Gly44- Ser398(Tyr285Leu) is expressed with a 6His tag at the C-terminus. P15291			
ACCESSION #	P13231			
Host	Human Cells			
Species	Human			
opecies	Human			
Predicted Molecular Mass	40.1 KDa			
-				
Predicted Molecular Mass	40.1 KDa Supplied as a 0.2 μm filtered solution of 20mM Tris-HCl, 150mM NaCl, 2mM			
Predicted Molecular Mass Formulation	40.1 KDa Supplied as a 0.2 μm filtered solution of 20mM Tris-HCl, 150mM NaCl, 2mM EDTA, 20% Glycerol, pH 8.0. The product is shipped on dry ice/polar packs. Upon receipt, store it immediately			

## **SDS-PAGE** image



## Background

Alternative Names	1,4- galactosyltransferase, po galactosyltransferase 1; Beta-1,4- GT1: GTB					
Background	Beta1,4-Galactosyltransferase-I	(B4GALT1),	one o	of seven	beta1,4-	

galactosyltransferases, is an enzyme commonly found in the trans-Golgi complex that adds galactose to oligosaccharides. By sequence similarity, the beta 4GalTs form four groups: beta 4GalT1 and beta 4GalT2, beta 4GalT3 and beta 4GalT4, beta 4GalT5 and beta 4GalT6, and beta 4GalT7. beta 4GalT1 is unique among the seven enzymes because it can be expressed either as membrane associated form or secreted form. The secreted form is restricted to lactating mammary tissues where the enzyme forms a heterodimer with alpha -lactalbumin to catalyze the synthesis of lactose. The Golgi complex form catalyzes the production of lactose in the lactating mammary gland and could also be responsible for the synthesis of complex-type N-linked oligosaccharides in many glycoproteins as well as the carbohydrate moieties of glycolipids. The cell surface form functions as a recognition molecule during a variety of cell to cell and cell to matrix interactions, as those occurring during development and egg fertilization, by binding to specific oligosaccharide ligands on opposing cells or in the extracellular matrix. Defects in beta 4GalT1 are the cause of congenital disorder of glycosylation type 2D (CDG2D)

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## Note

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