Structure-based Design of β1,4-Galactosyltransferase I (β4Gal-T1) with Equally Efficient N-Acetylgalactosaminyltransferase Activity

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β1,4-Galactosyltransferase I (Gal-T1) normally transfers Gal from UDP-Gal to GlcNac in the presence of Mn2+ ion. In the presence of α-lactalbumin (LA), the Gal acceptor specificity is altered from GlcNac to Glc. Gal-T1 also transfers GalNac from UDP-GalNac to GlcNac, but with only ~0.1% of Gal-T activity. To understand this low GalNAc-transferase activity, we have carried out the crystal structure analysis of the Gal-T1-LA complex with UDP-GalNac at 2.1-Å resolution. The crystal structure reveals that the UDP-GalNac binding to Gal-T1 is similar to the binding of UDP-Gal to Gal-T1, except for an additional hydrogen bond formed between the N-acetyl group of GalNAc moiety with the Tyr-289 side chain hydroxyl group. Elimination of this additional hydrogen bond by mutating Tyr-289 residue to Leu, Ile, or Asn enhances the GalNAc-transferase activity. Although all three mutants exhibit enhanced GalNAc-transferase activity, the mutant Y289L exhibits GalNAc-transferase activity that is nearly 100% of its Gal-T activity, even while completely retaining its Gal-T activity. The steady state kinetic analyses on the Leu-289 mutant indicate that the K_m for GlcNac has increased compared to the wild type. On the other hand, the catalytic constant (k_cat) in the Gal-T reaction is comparable with the wild type, whereas it is 3-5-fold higher in the GalNAc-T reaction. Interestingly, in the presence of LA, these mutants also transfer GalNac to Glc instead of to GlcNac. The present study demonstrates that, in the Gal-T family, the Tyr-289/Phe-289 residue largely determines the sugar donor specificity.

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MATERIALS AND METHODS

Site-directed Mutagenesis—Site-directed mutagenesis was performed using the PCR method. Construction of the mutants was done using plasmid pEGT-d129 as the template; this contains a Cys-342 to Thr mutation. The mutation primers corresponding to the upper DNA strand are: Y289L, 5′-CTCTAAG-3′ and 5′-GATCTTTGAGGCTAAATTTGGA-3′; Y289I, 5′-CTCTAAGGAGGCTAAATTTGGA-3′; and 5′-GATCTTTTGAGGCTAAATTTGGA-3′. The abbreviations used are: β4Gal-T1, β1,4-galactosyltransferase I; LA, α-lactalbumin; LS, lactose synthase; wt, wild-type; r.m.s., root mean square; LacDiNAc, di-N-acetylatedsediamine; GalNAc-T, N-acetylgalactosyltransferase.

β1,4-Galactosyltransferase I (β4Gal-T1, EC 2.4.1.38) catalyzes the transfer of galactose (Gal) from UDP-Gal to the N-acetylglucosamine (GlcNAc) residue present at the nonreducing terminal end of glycans of glycoproteins and glycolipids (1), producing β1,4-linked galactosylated glycan. In addition to GlcNAc as an acceptor, the enzyme can also use other sugars, such as N-acetyl-substituted glucosamines and N-acetyl-mannosamine, as acceptors (2). The enzyme does not have an absolute requirement for the sugar donor UDP-Gal; instead, it exhibits polymorphic donor specificity, in that it also transfers glucose (Glc), α-deoxy-Glc, arabinose, GalNAc, and GlcNAc from their UDP derivatives, albeit at low rates (0.3–5%) compared with Gal transfer (3–7).

α-Lactalbumin (LA), a mammary gland-specific calcium-binding protein, alters the sugar acceptor specificity of Gal-T1 toward Glc (8). This activity is described as the lactose synthase (LS) activity of β4Gal-T1 (EC 2.4.1.22). LA has been shown to alter not only the sugar acceptor specificity of the enzyme, but also its sugar donor specificity (7, 9), because it stimulates the transfer of Glc from UDP-Glc to GlcNAc. Similarly, it has been shown previously that the transfer of GalNAc from UDP-GalNAc to GlcNAc by Gal-T1 is also enhanced in the presence of LA (5). To better understand the modulation of substrate specificity, we had carried out a series of studies on the crystal structure of LA-Gal-T1 complex with various substrates (7, 10). Analysis of the molecular interactions between LA-Gal-T1 and Glc from the crystal structure suggested an explanation for the way in which LA modulates the acceptor specificity (10). The structural studies of the LA-Gal-T1 complex with UDP-Glc, together with the kinetic studies of Glc transfer from UDP-Glc to an acceptor sugar, also offered an explanation for the way in which LA modulates the donor specificity of Gal-T1 (7). In our continuing efforts to understand the structure and function of Gal-T1, we examined in the present study the interaction of UDP-GalNAc with Gal-T1. The results suggest that the formation of a hydrogen bond between GalNAc and Tyr-289 is responsible for the poor GalNAc-transferase activity that Gal-T1 exhibits. On the other hand, mutation of Tyr-289 to leucine, isoleucine, or asparagine enhances the GalNAc-transferase activity. Among these mutants, Y289L has GalNAc-transferase activity that is nearly equal to Gal-T activity.
results and discussion

Crystal Structure of Gal-T1-LA-UDP-GalNAc Complex and Molecular Interactions of UDP-GalNAc Molecule—The overall molecular structure and the interactions between Gal-T1 and LA molecules in the Gal-T1-LA-UDP-GalNAcMn 

The overall structure of the Gal-T1-LA-UDP-GalNAcMn 

free N-acetyl group in UDP-GalNAc, which may better trap the water molecule.

Although the interactions of Gal-T1 with the GalNAc moiety of UDP-GalNAc are similar to the interactions of the Gal moiety of UDP-Gal (17), additional interactions are observed between the N-acetyl group of UDP-GalNAc with Gal-T1 (Fig. 1B). In the Gal-T1-UDP-Gal complex, the O2 hydroxyl group forms only one hydrogen bond with Asp-252, but in the present crystal structure the N2 nitrogen atom is hydrogen-bonded to the Asp-252 side chain carbonyl oxygen atom, whereas the carbonyl oxygen atom is hydrogen-bonded to the Tyr-289 side chain hydroxyl group.

The presence of the N-acetyl group seems to have perturbed the conformation of both the GalNAc and the Tyr-289 side chain. The orientation of the N-acetyl group with respect to the sugar ring is represented by the 

H2-C2-N2-H2, which in the case of GalNAc, GlcNAc, and ManNAc was observed to be nearly 180° (18). In the present crystal structure, there are two Gal-T1-LA-UDP-GalNAcMn 

molecules in the asymmetric unit; in one molecule the torsion angle is 150° and in the other it is 120°, suggesting the existence of the N-acetyl group in an unfavorable orientation. This unfavorable orientation is a result of the
lack of space between the side chain hydroxyl group of Tyr-289 and the carbonyl oxygen atom of the \( \text{N}^-\text{acetyl} \) group. In the two molecules of the asymmetric unit, because of this slight difference in the torsion angle of H2-C2-N2-HN, orientation of the side chain of Tyr-289 is slightly different to accommodate the \( \text{N}^-\text{acetyl} \) group.

**Effect of the \( \text{N}^-\text{Acetyl} \) Group of GalNAc on GalNAc-T Activity**—Among the interactions of the \( \text{N}^-\text{acetyl} \) group of UDP-GalNAc with the Gal-T1 molecule, the hydrogen bond between the carbonyl oxygen atom of the \( \text{N}^-\text{acetyl} \) group and the Tyr-289 side chain hydroxyl group seems to cause steric hindrance for catalysis. Studies using positional isotope exchange, secondary deuterium isotope effects, and inhibition of catalytic activity by acceptor analogs have suggested a plausible catalytic mechanism for the Gal-T1 (19–21). According to this mechanism, upon a nucleophilic attack from the O4 hydroxyl group of the acceptor molecules, Gal moiety, during its separation from UDP-Gal in the transition state complex, exits with a substantial sp\(^2\) character at the anomeric center C1. Such an oxocarbonium ion intermediate forms a covalent linkage with the O4 oxygen atom, forming a \( \beta1-4 \) glycosidic linkage between galactose and the acceptor molecule. Because it is not feasible to obtain the crystal structure of Gal-T1 in the presence of both donor and acceptor molecules to determine the nature of interactions between the substrates and Gal-T1, we have carried out modeling studies based on the individual donor and acceptor bound to Gal-T1 structures (10, 17). These studies indicated that the O4 hydroxyl group of the acceptor lines up against the anomeric center C1 of the bound UDP-Gal molecule at 4.5 Å (Fig. 2A). During catalysis, this assembly of the substrates in the catalytic pocket requires the displacement of the substrates toward each other. Because any displacement of the acceptor GlcNAc toward the donor is prevented by Trp-314, it is most likely that the donor Gal moiety moves toward the acceptor
during catalysis. In the Gal-T reaction, when the Gal moiety is involved during catalysis, there is enough space between the C2 atom of Gal and the side chain of Tyr-289, as the O2 hydroxyl group of Gal is facing away from the Tyr-289 residue (Fig. 2A). However, in the crystal structure of Gal-T1/UDP-GalNAc, the bulky N-acetyl group of GalNAc occupies this space with its carbonyl oxygen atom hydrogen-bonded to the hydroxyl group of Tyr-289 (Fig. 2B). We hypothesized from this observation that, during catalysis, this hydrogen bond is expected to hinder the reaction by not allowing the GalNAc moiety to move toward the acceptor molecule. Second, because of lack of available space between the N-acetyl group and Tyr-289, the former exists in a less favorable orientation. To create the space in the Gal-T1 molecule in the vicinity of the N-acetyl group binding site of UDP-GalNAc, we mutated the Tyr-289 residue to Leu, Ile, or Asn (Fig. 2C). A phenylalanine mutation was not considered, because it does not create enough space when the N-acetyl group adopts its favorable conformation. Although Leu or Asn residue substitution creates a similar pocket because of the similarity of their side chain length (Fig. 2C), Leu is expected to create a hydrophobic pocket whereas Asn creates a hydrophilic one. Additionally, because of its shorter side chain length, an Ile substitution would be expected to create even a larger pocket size than the Leu substitution. Thus, investigations of these three mutants were expected to identify the required size and the nature of the pocket in the vicinity of the N-acetyl group-binding site of UDP-GalNAc.

Comparison of Gal-T and GalNAc-T Catalytic Activities of Wild-type and Gal-T1 Mutants—The mutants Y289L, Y289N, and Y289I exhibit both Gal-T and GalNAc-T activities at the saturation substrate concentrations (Table II and Fig. 3). The mutants Y289L and Y289N exhibit equally strong Gal-T and GalNAc-T activities, suggesting that a Leu or Asn substitution of Tyr-289 seems to create the required optimal space. The specific GalNAc-T activity of the mutant Y289L at the saturation concentration is half that of Y289L (Table II); it exhibits a slightly weaker affinity to UDP-GalNAc (Fig. 3). Therefore, an Ile substitution seems to create more than the required space, thus reducing the specific activities. Although Asn substitution exhibits the GalNAc-T activity as well as the Leu substitution, the protein seems to be less stable and tends to denature rapidly compared with the Leu mutant. It has been shown that wt-Gal-T1 exhibits a very low glucosyltransferase activity (Glc-T), but no N-acetylgalactosaminyltransferase (GlcNAc-T) activity (3, 6, 7). On the contrary, the mutants exhibit reasonable GlcNAc-T activity (Table II) where they transfer GlcNAc from UDP-GlcNAc to the acceptor GlcNAc but do not exhibit Glc-T activity. It is interesting to note that in this GlcNAc-T activity the initial product, the disaccharide GlcNAc β1,4-GlcNAc, itself is an acceptor for the enzyme, thus producing tri- and longer

| Residue | Atom | Distance in Angstroms |
|---------|------|----------------------|
| His-347 | Nε2  | 2.43                 |
| UDP-GalNAc | O[Pα] | 2.24                 |
| UDP-GalNAc | O[Pβ] | 2.31                 |
| Met-344 | Sδ   | 2.84                 |
| Asp-252 | Oε2  | 2.22                 |
| Water   | O    | 2.57                 |

FIG. 2. A, modeling of the GlcNAc in its binding site (based on the crystal structure of Gal-T1/UDP-GlcNAc complex) on the Gal-T/UDP-GalNAc crystal structure (17). As can be seen, the acceptor atom O4 of GlcNAc is in the proper orientation with the anemic center (C1 atom) of the Gal moiety of UDP-Gal. The distance between these atoms is ~4 Å. It has been hypothesized that the enzyme catalysis follows the separation principle, in which the Gal moiety is expected to move toward the acceptor while forming the disaccharide. Because there is enough space between Tyr-289 and the Gal moiety, Tyr-289 is not expected to hinder the Gal-T activity. This is illustrated by the dotted surface showing the van der Waals sphere for the Tyr-289 side chain hydroxyl group and the C2 and O2 atoms of the Gal moiety. B, modeling of the GlcNAc in its binding site on the Gal-T1/UDP-GalNAc-Mn²⁺ crystal structure. Because LA does not interact with the donor molecule, it is not shown in the figure. Similar to UDP-Gal binding (Fig. 2A), the O4 atom of GlcNAc is located ~4 Å away from the anemic center of GalNAc moiety of UDP-GalNAc. The major difference between the binding of UDP-Gal and UDP-GalNAc seems to be an additional hydrogen bond observed between the N-acetyl group and Gal T1, particularly with Tyr-289. The side chain hydroxyl group of Tyr-289 is hydrogen-bonded with the carbonyl oxygen atom of the N-acetyl group of GalNAc (shown by double arrow). If the catalytic mechanism still follows the separation principle, this hydrogen bond would hinder the required displacement of the GalNAc moiety during the disaccharide linkage formation. As can be seen from the van der Waals sphere (dotted surface), little space exists between the Tyr side chain hydroxyl group and the N-acetyl group. Therefore, based on this observation, it was hypothesized that lack of space is responsible for the poor GalNAc-T activity. C, based on the hypothesis that more space is needed between the N-acetyl group of UDP-GalNAc and Tyr-289 residue to facilitate the reaction, the substitution of Tyr-289 by Leu is modeled. The van der Waals sphere shown in dotted surface for the side chain atoms of Leu-289 and the N-acetyl group clearly indicates that such substitution creates additional space.
chain saccharides. A separate study on this GlcNAc-T property is being carried out.

Because the Y289L variant of Gal-T1 exhibits both Gal-T and GalNAc-T enzyme activities with equal efficiencies, double substrate kinetic studies were carried out to determine the kinetic constants for both the donor and acceptor molecules. The kinetic data from both reactions fit best to Equation 2, with a zero $K_i$ value, describing an asymmetric initial velocity pattern for a double-displacement or "ping-pong" mechanism. The Y289L mutant in the Gal-T reaction shows for the acceptor nearly 20-fold higher $K_m$ than the wild-type Gal-T1 (Table III). The catalytic constant ($k_{cat}$) in the Gal-T1 reaction is comparable with the wild type, but it is nearly 3–5-fold higher in the GalNAc-T reaction (Table III). It is possible that this high $k_{cat}$ was achieved either by stabilizing the transition state or by the increase in the free energy of the ternary complex. Although the mutation was designed to eliminate the steric hindrance caused by Tyr-289 during the transition state, the latter possibility cannot be ruled out because of the observed high $K_m$ value for the acceptor GlcNAc. In the crystal structure of Gal-T1, the residues around Tyr-289 are involved in the acceptor binding. The lack of an aromatic side chain at position 289 may destabilize these residues, thus affecting the binding of the acceptor substrate. This is substantiated by the earlier mutational studies on human Gal-T1, where substitution by Phe for the corresponding residue Tyr-287 does not affect the acceptor binding, whereas substitution with Gly totally abolished the catalytic activity (22). Currently, mutational studies are being

TABLE II
Specific activities of the catalytic domain of Gal-T1 (d129-Gal-T1, residues 130–402) with C342T mutation and the Tyr-289 mutants
Reactions were performed under saturating conditions of all substrates (see "Materials and Methods" for assay conditions). In these reactions, the concentration for the donors was at 500 μM and the acceptor α-benzyl-GlcNAc was at 25 mM. In the Gal-T reaction of C342T, the acceptor concentration was 10 mM, because at 25 mM it showed inhibition. Because Y289N rapidly undergoes denaturation, reliable data could not be obtained.

| Enzyme     | UDP-galactose → GlcNAc Gal-T activity | UDP-GalNAc → GlcNAc GalNAc-T activity | UDP-GlcNAc → GlcNAc GlcNAc-T activity |
|------------|--------------------------------------|---------------------------------------|--------------------------------------|
| C342T      | 11.8                                 | 0.1                                   | 0                                    |
| Y289L_C342T| 16.2                                 | 27.3                                  | 8.2                                  |
| Y289I_C342T| 3.8                                  | 14.7                                  | 6.2                                  |

TABLE III
Kinetic parameters for the donor and the acceptor substrates by Y289L_C342T mutant in the Gal-T and GalNAc-T catalytic reactions (see text for assay conditions)

| Mutant    | Gal-T UDP-galactose → GlcNAc | GalNAc-T UDP-N-acetylgalactosamine → GlcNAc |
|-----------|------------------------------|---------------------------------------------|
|           | $K_a$ $K_b$ $k_{cat}$        | $K_a$ $K_b$ $k_{cat}$                      |
| C342T     | 93 (6) 11 (1) 14             | ND ND ND                                   |
| Y289L_C342T| 75 (1) 198 (1) 8.5           | 354 (3) 51 (1) 40                         |

ND, not determined.

Fig. 3. The catalytic activity of the wt-Gal-T1 and the mutants Y289L, Y289I, and Y289N. A, Gal-T activity of wt-Gal-T1 and of mutants at the saturating concentrations of α-benzyl-GlcNAc as the acceptor (see "Materials and Methods"). The mutant Y289L seems to be as active as the wt-Gal-T1, whereas Y289I seems to be weaker. B, GalNAc-T activity of the wt-Gal-T1 and of mutants. The wt-Gal-T1 exhibits very poor GalNAc-T activity, whereas Y289L exhibits more GalNAc-T activity than its Gal-T activity. Y289I exhibits a somewhat weaker GalNAc-T activity, compared with that of Y289L. The activities are measured with 50 mM α-benzyl-GlcNAc as acceptors at different concentrations of the donor. The Gal-T activity of the wt-Gal-T1 was measured with only 10 mM α-benzyl-GlcNAc, because it showed inhibitions at 50 mM concentration.
carried out to enhance the acceptor affinity of Y289L mutant to determine which mechanism is responsible for the high catalytic efficiency of the enzyme.

Although the weak GalNAc-T activity of the wild-type Gal-T1 had been demonstrated in the earlier studies and the product disaccharide, di-N-acetyllactosperidiamine (LacdiNAc), was characterized by 1H NMR technique, no details were reported (6). Because the Y289L mutant exhibits equally high GalNAc-T activity as it does Gal-T activity, the disaccharide product from this reaction was purified and analyzed by 1H NMR spectroscopy. Previously, the same product LacdiNAc, from N-acetylgalactosaminyltransferase isolated from bovine mammary gland, had been fully characterized by 1H NMR spectroscopy (23). The present NMR spectrum is quite similar to that of the already characterized LacdiNAc, demonstrating that the mutant Y289L transfers GalNAc from UDP-GalNAc to GlcNAc, forming a β1–4 linkage between GalNAc and GlcNAc.

A Point Mutation in the Codon for Amino Acid 289 Can Convert the Enzyme with Dual Property—In humans, the β4Gal-T family has seven members (Gal-T1 to -T7), each with high sequence identity within their catalytic domain (24, 25). These family members are known to transfer Gal from UDP-Gal to different sugar acceptors. For example, Gal-T6 transfers Gal to Glc of Glc-ceramide, whereas Gal-T7 transfers Gal to xylose (25). Among these seven members, four members, Gal-T1 to Gal-T4, have a Tyr residue at position 287 (which corresponds to Tyr-289 in bovine Gal-T1), whereas Gal-T5 and Gal-T6 have a Phe residue. Mutational studies in human Gal-T1 have shown that the Y287F mutant exhibits the same enzyme characteristics with kinetic constants as that of the wild-type Gal-T1 (22). Because the Phe residue may also create a similar steric problem for UDP-GalNAc as the Tyr residue, these enzymes are expected to exhibit low GalNAc-T activity. It is interesting to note that the family members show variation in their codon nucleotide sequence for residue 287 (Table IV), particularly at the second and third nucleotide positions of the codon. A mutation of the first nucleotide base thymidine (T) to either adenine (A) or cytidine (C) would result in a mutation of the Tyr/Phe residue to either Leu or Asn. Such a spontaneous point mutation at the first nucleotide position of the codon for the residue 287 would generate an enhanced equal GalNAc-transferase activity that might have adverse effects. Incorporation of GalNAc instead of Gal might lead to 1) termination of the total synthesis of a glycoconjugate, or 2) glycans with different oligosaccharide sequence. It is worth noting, however, that, although mutants may possess dual enzyme activities, the substitution of Gal by GalNAc is expected to be highly
irregular and will depend on whether or not UDP-GalNAc is available in the Golgi apparatus. 

Role of LA in the GalNAc-T Activity of the wt-Gal-T1 and the Y289L Mutant—Although Gal-T1 transfers GalNAc from UDP-GalNAc to GlcNAc with $-0.1\%$ of its Gal-T efficiency, it has been shown that LA enhances this transfer (5), a situation similar to glucosyltransferase activity (Glc-T) of Gal-T1, where LA increases this activity from 0.3 to 10\% (7). We have shown previously through crystal structure and enzyme kinetic studies that, in the Glc-T reaction, LA plays a kinetics role in stimulating Gal-T1 to transfer Glc from the UDP-Glc to GlcNAc (7, 9). Although we have not carried out detailed kinetic studies on the GalNAc-T reaction by the wild-type Gal-T1 in the presence of LA, the similarity between Glc-T and GalNAc-T catalytic activities indicates that LA probably plays a similar role in the GalNAc-T reaction. It has been stated that LA enhances this GalNAc-T activity to 55\% of its Gal-T activity (5); however, we find that the activity is enhanced only to nearly 1\% (data not shown). This may have been because of the differences in the assay methods. For example, in the present study and the studies reported by others (12), the donor concentration is not shown. This may have been because of the differences in the donor specificity of the enzyme.

Unlike wt-Gal-T1, where LA stimulates the transfer of GalNAc only to GlcNAc and not to Glc (5), the Tyr-289 mutants (Y289L, Y289N, and Y289I) in the presence of LA transfer GalNAc preferably to Glc rather than to GlcNAc. This property of the Tyr-289 mutants of Gal-T1, which have an optimal space for the binding of the N-acetyl group of UDP-GalNAc to Gal-T1, is required, and such a space need not be hydrophobic nor hydrophilic in nature. In the present study, we have shown that the Tyr-289 mutants of Gal-T1, which have an optimal space for the binding of N-acetyl group, exhibit as high GalNAc-T activity as Gal-T activity, as well as GlcNAc-T activity. In the human Gal-T family members, the Tyr/Phe residue at 287 (or in bovine Tyr-289) seems to be important for determining the donor sugar specificity of these enzymes, and mutation of this residue broadens the donor specificity of the enzyme. The present study suggests that a single point mutation can result in glycan moieties substituting Gal with GalNAc or GlcNAc, which may have significant effects on cellular functions.

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