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Converting *Pasteurella multocida* α 2–3-sialyltransferase 1 (PmST1) to a regioselective α 2–6-sialyltransferase by saturation mutagenesis and regioselective screening†

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Abstract

A microtiter plate-based screening assay capable of determining the activity and regioselectivity of sialyltransferases was developed. This assay was used to screen two single-site saturation libraries of *Pasteurella multocida* α 2–3-sialyltransferase 1 (PmST1) for α 2–6-sialyltransferase activity and total sialyltransferase activity. PmST1 double mutant P34H/M144L was found to be the most effective α 2–6-sialyltransferase and displayed 50% reduced donor hydrolysis and 50-fold reduced sialidase activity compared to the wild-type PmST1. It retained the donor substrate promiscuity of the wild-type enzyme and was used in an efficient one-pot multienzyme (OPME) system to selectively catalyze the sialylation of the terminal galactose residue in a multigalactose-containing tetrasaccharide lacto-*N*-neotetraoside.

Graphical Abstract

A sialyltransferase mutant for selectively α 2–6-sialylating terminal galactose in polyLacNAc-glycan was identified using a novel microtiter plate-based screening.

Introduction

Sialic acid-terminated glycans and glycoconjugates are critical recognition elements involved in a remarkably wide range of pathological and physiological events in human, including cell-cell recognition, inflammation, cancer metastasis, immune regulation, and bacterial and viral infections.^{1–3} Sialyltransferase-catalyzed reactions with or without *in situ*

†Electronic Supplementary Information (ESI) available: pH profiles and donor substrate promiscuity studies of PmST1 P34H/M144L. ¹H and ¹³C NMR spectra for Neu5Ac2-6Gal β 1-4GlcNAc β 1-3Gal β 1-4Glc β ProN₃ (Neu5Ac2-6LNnT β ProN₃) and Neu5Ac2-6Gal β 1-4GlcNAc β 1-3(Neu5Ac2-6)Gal β 1-4Glc β ProN₃. See DOI: 10.1039/x0xx00000x

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enzymatic generation of cytidine 5'-monophosphate (CMP)-sialic acid have been effective approaches to obtain diverse structurally defined sialosides in high yields.^{2, 3} Of particular note for their low cost and effectiveness are one-pot multienzyme sialylation systems^{4, 5} combining a sialyltransferase (ST) for the formation of a specific sialyl linkage, a CMP-sialic acid synthetase for the generation of a CMP-sialic acid *in situ* from CTP and sialic acid, and optionally a sialic acid aldolase for the formation of the sialic acid from its six-carbon monosaccharide precursor *N*-acetylmannosamine (ManNAc) or derivatives and pyruvate. The flexible substrate specificities of these enzymes have allowed the formation of large libraries of sialosides containing diverse naturally occurring and non-natural modifications at various positions of sialic acid residues. In addition to the diversity of natural sialic acid forms of which more than 50 have been identified,^{6, 7} sialic acid can be found linked to different monosaccharides in various locations and with a range of sialyl linkages.⁸ For example, α -linked sialic acid can be found attached to the C3 of a terminal galactose (Gal) residue; to the C6 of a terminal or an internal Gal or *N*-acetylgalactosamine (GalNAc) residue; to the C6 of an internal glucose (Glc) (e.g. in *Neisseria meningitidis* serogroup Y capsular polysaccharide) or *N*-acetyl glucosamine (GlcNAc) (e.g. in milk oligosaccharide disialyllacto-*N*-tetraose or DSLNT) residue;^{9, 10} or to the C8 or C9 of another sialic acid residue.^{2, 3, 11, 12}

Several bacterial sialyltransferases have been frequently used for preparative and large-scale synthesis of sialosides.² Among these enzymes, those from Carbohydrate Active enZyme (CAZy, www.cazy.org) family^{13, 14} GT80 are of particular interest as several of them have robust heterologous expression in *Escherichia coli*, high catalytic activity, broad promiscuity toward modified donor substrates and various acceptors, and have been widely used for the synthesis of α 2-3- and α 2-6-linked sialosides containing different sialic acid structures.^{15, 16}

While the CAZy GT80 family α 2-3-sialyltransferases (α 2-3STs), such as multifunctional *Pasteurella multocida* sialyltransferase 1 (PmST1),¹⁵ are quite specific in catalyzing α 2-3-sialylation to only the terminal galactose residues in galactosides,^{10, 17-19} the α 2-6-sialyltransferase (α 2-6ST) activity of multifunctional *Photobacterium damsela* α 2-6ST (Pd2,6ST)²⁰ can catalyze α 2-6-sialylation to both terminal and internal galactose or GalNAc residues.^{10, 12, 16, 17, 21}

The lack of selectivity of Pd2,6ST in sialylating terminal or internal Gal was advantageous for one-step sialylation of a tetrasaccharide lacto-*N*-neotetraose (LNnT) for generating disialyllacto-*N*-neotetraose DSLNnT, a hexaose mimicking human milk DSLNT.¹⁰ Both DSLNnT and DSLNT were effective in protecting neonatal rats from necrotizing enterocolitis in an animal model and could be potentially developed into therapeutic agents.^{9, 10} This property of Pd2,6ST also allowed facile synthesis of multisialylated poly-*N*-acetylglucosamine (poly-LacNAc) extensions on O- and N-linked glycans for microarray studies,^{12, 22} as well as the synthesis of glycans with α 2-3-sialylation at the terminal Gal and α 2-6-sialylation at the internal Gal.^{22, 23} However, it generated complications for synthesizing the terminal disialyl tetrasaccharide sequence of gangliosides GD1 α , GT1 α , GQ1 α .²¹ In addition, it would not be suitable for obtaining mono- α 2-6-sialylated longer

glycans such as α 2–6-linked sialyl LNT or sialyl LNnT structures in the milk and on the cell surfaces of human and animals.^{11, 24}

While a human α 2–6ST hST6Gal-I¹² and a ST6Gal-I purified from rat liver²⁵ were able to catalyze the addition of a sialic acid residue selectively to the terminal Gal on poly-LacNAc-type structures, a bacterial sialyltransferase equivalent has not been identified. Other than unselective α 2–6-sialylation of terminal and internal Gal or GalNAc residues by Pd2,6ST,^{10, 12, 16, 17, 21} several other reported bacterial α 2–6STs^{26–28} including those from GT80 family have not been tested for sialylation of poly-LacNAc-type structures. While hST6Gal-I was able to be expressed in *E. coli* cells at a level of 0.266 mg purified enzyme per liter culture by N-terminal truncation with fusing to maltose-binding protein (MBP)²⁹ and the expression level was improved to 2 mg purified protein per liter culture by co-expression of multiple chaperon/foldases in the Origami2(DE3) strain,³⁰ the amount was still limited for large-scale synthetic purposes.

The crystal structures have been reported for several GT80 family sialyltransferases, including those preferring the formation of α 2–3-^{31–35} or α 2–6-sialyl^{36, 37} linkage. Comparing the crystal structures and the sialyl linkage specificities of the products formed by PmST1 (PDB ID 2ILV)^{31, 32} and *Photobacterium* sp. JT-ISH-224 α 2–6ST (Psp2,6ST) (PDB ID 2Z4T)³⁶ indicated that H123 in Psp2,6ST is a key residue determining the α 2–6ST specificity of the enzyme while the corresponding P34 in PmST1 is associated with its α 2–3ST specificity.³⁶ Indeed, this was also predicted by protein sequence alignment of all characterized GT80 family sialyltransferases and was confirmed by another GT80 family *Pasteurella dagmatis* α 2–3ST (Pd2,3ST) (PDB ID 4V2U)³⁸ for which mutating the corresponding P7 residue to a histidine converted the enzyme from an α 2–3ST to an α 2–6ST with 95% selectivity for α 2–6-sialylation.³⁵ The regio-selectivity for the formation of α 2–6-sialyl linkage versus α 2–3-sialyl linkage was further enhanced by mutating M117 to alanine. The Pd2,3ST P7H/M117A double mutant had a greater than 99.5% selectivity for α 2–6-sialylation. However, each of these mutations resulted in decreased sialylation activity with k_{cat}/K_M value dropped from 16 s⁻¹ mM⁻¹ for the wild-type Pd2,3ST to 4.3 s⁻¹ mM⁻¹ and 2.8 s⁻¹ mM⁻¹ for the P7H and P7H/M117A mutants, respectively.

We hypothesized that converting a substrate promiscuous α 2–3ST which catalyzes the selective α 2–3-sialylation of terminal Gal to an α 2–6ST by site-specific mutagenesis might retain its selectivity on sialylating the terminal Gal and substrate promiscuity. PmST1 was chosen as the parent enzyme for engineering. This was because that among all reported GT80 α 2–3STs,^{15, 35, 39–43} PmST1 has the highest expression level in *Escherichia coli* (100 mg per L culture), the highest measured kinetic constants,¹⁵ the largest number of previous mutagenesis studies,^{31, 38–41, 43–45} and the most complete structural and functional characterization.^{15, 31, 32} Additionally, PmST1 has been shown to be promiscuous towards a wide range of CMP-sialic acid donors and galactoside acceptors.^{15, 19, 46–49} PmST1 has multiple functions (Figure 1) including (1) α 2–3ST, (2) α 2–6ST, (3) α 2–3-sialidase, (4) α 2–3-trans-sialidase, (5) donor hydrolysis,^{15, 45} and (6) α 2–6-trans-sialidase⁵⁰ activities activity. As the α 2–6ST activity of PmST1 is much weaker than its α 2–3ST activity especially at pH 7.0,¹⁵ PmST1 has been broadly used in highly efficient synthesis of α 2–3-sialosides with *in situ* generation of CMP-sialic acids by a CMP-sialic acid synthetase with or without

a sialic acid aldolase.^{2, 51} Furthermore, the product sialyl linkage specificity-switching effect of the P7H mutation for Pd2,3ST has already been confirmed for PmST1 with the corresponding P34H mutation in its trans-sialidase reaction capacity for the formation of α 2–6-sialylated products.⁵²

Here we report our strategy of converting PmST1 to an α 2–6ST with the same selectivity in sialylating the terminal Gal in poly-LacNAc-type structures such as a multigalactose-containing tetrasaccharide lacto-*N*-neotetraoside Gal β 1–4GlcNAc β 1–3Gal β 1–4Glc β ProN₃ (LNnT β ProN₃). This was accomplished by structure-guided saturation mutagenesis followed by a novel colorimetric screening method in microtiter plates which can be adapted to automation for a high-throughput platform. Various in situ generated CMP-sialic acids containing naturally occurring and non-natural sialic acid forms were shown to be tolerable donor substrates for the mutant, indicating the retaining of the donor substrate promiscuity of the wild-type PmST1 by the mutant. *N*-Acetylneuraminic acid (Neu5Ac), the most common sialic acid form, was used as a donor precursor for preparative-scale synthesis using a one-pot two-enzyme sialylation system.⁵ The resulting Neu5Ac α 2–6LNnT (LSTc) represents a common structure found in human milk¹¹ and on mammalian cell surface.⁵³

Results and discussion

Initially we sought to reproduce the PmST1 mutant homologs for Pd2,3ST P7H and P7H/M117A. The corresponding PmST1 residues P34 and M144 were mutated to generate PmST1 P34H single mutant and P34H/M144A double mutant. Both mutants catalyzed α 2–6-sialylation but displayed 2.5- and 6.4-fold, respectively, decreased activity relative to the α 2–3ST activity of the wild-type PmST1 (Table 1). A similar loss of activity was reported for the Pd2,3ST P7H single mutant and P7H/M117H double mutant (Table 1).³⁵ Additionally, the PmST1 P34H and P34H/M144A mutants were found to catalyze CMP-Neu5Ac hydrolysis at about 57-fold and nearly 3-fold, respectively, the catalytic efficiency of their α 2–6ST activity, hindering their efficiency for the preparation of sialosides. The α 2–6-sialidase activities of both mutants were more than 150-fold lower than their corresponding α 2–6ST activities, which is a preferable feature for sialoside synthesis.

A closer examination of the sequence homology among GT80 STs revealed that while P34 is conserved in α 2–3STs, and α 2–6STs have a conserved histidine residue at the corresponding site, residues homologous to M144 displayed a greater diversity. Previous mutagenesis studies by our lab and others indicated that M144 of PmST1 and its homologous residues in other GT80 STs are a multifunctional “hotspot,” and that mutations at this position can greatly impact donor hydrolysis activity, sialidase activity, sialyltransferase activity, donor specificity, and acceptor specificity.^{45, 55} Therefore we hypothesized that screening a PmST1 P34H/M144X saturation library might reveal a more reactive α 2–6ST.

In order to identify mutants with improved sialyltransferase activities and with regioselectivity for the formation of either α 2–3- or α 2–6-sialyl linkage, a novel high-throughput screening assay was developed and used to screen PmST1 P34X and P34H/M144X saturation libraries. To screen each saturation library, a 96-deep well culture plate

was inoculated with 88 mutant colonies, 4 wild-type colonies, and 4 empty vector colonies. The libraries were checked by sequencing several colonies from each library as is typical for saturation mutagenesis. No nucleotide sequence bias was detected during this analysis. TopLib analysis⁵⁶ shows that the likelihood of discovering the best variant from screening 88 wells of an NNK single site saturation library is 96.24%. The lysates from these cultures were used for activity screen in 384-well plates. The screen relied on using sialic acid transferred by the sialyltransferase mutants as a protective group for *para*-nitrophenyl β -D-galactopyranoside (Gal β pNP, **1**), a substrate for β -galactosidases which produces a more intense chromophore upon hydrolysis (Figure 2A). On a 384-well microtiter plate, the PmST1 mutants were allowed to sialylate Gal β pNP (**1**), then two aliquots of each reaction were transferred to empty 384-well microtiter plate wells. One of these wells (well A) was treated with β -galactosidase only, while the other (well B) was treated with both β -galactosidase and *Streptococcus pneumoniae* sialidase SpNanB,^{57–59} an α 2–3-sialyl linkage specific sialidase. In well A, the remaining **1** in the sialylation reaction was hydrolyzed by the excess β -galactosidase added to form *para*-nitrophenol (pNP). In well B, both remaining **1** and any α 2–3-sialoside (**2**) formed would be hydrolyzed by the combined actions of β -galactosidase and α 2–3-specific sialidase SpNanB to form pNP and only the α 2–6-sialoside (**3**) resisted hydrolysis catalyzed by the exoglycosidases and remained in the mixture. Upon quenching with *N*-cyclohexyl-3-aminopropanesulfonic acid (CAPS) buffer (pH 9.6) to form *para*-nitrophenolate from pNP and quantification at 405 nm using a plate reader, the extents of total sialylation (from well A) and α 2–6-sialylation (from well B) were determined and the α 2–3-sialylation could be calculated by the difference between the total and the α 2–6-sialylation.

The assay was validated using synthetic Neu5Aca2–3Gal β pNP (**2**) and Neu5Aca2–6Gal β pNP (**3**) as product standards (Figure 2B). Indeed, treating compound **2** with both β -galactosidase and α 2–3-sialidase SpNanB (well B) led to its hydrolysis and a large increase in the A_{405 nm} relative to **2** treated with only β -galactosidase (well A). In comparison, compound **3** was resistant to degradation under both conditions and low signal was observed for both wells. Compound **1** released *para*-nitrophenolate under both conditions resulting in high A_{405 nm} values for both.

Sialyltransferase activity was tested at pH 6.5. This pH was chosen with the idea that mutants with altered sialidase activity or linkage specificity would be more noticeable, as wild-type PmST1 shows enhanced rates of α 2–3-sialidase and α 2–6ST activity at low pH. Additionally, this pH was found to be optimal for the degradation of compound **2** in well B. Wells of interest from each plate were re-expressed and re-screened from a single microtiter plate for validation and easy comparison. The combined rescreen data of PmST1 P34X and P34H/M144X libraries are shown in Figure 3. In the P34X library, P34H was identified from two wells as the sole mutant displaying detectable α 2–6ST activity. In the P34H/M144X library, enzymes with higher yields in producing α 2–6-sialosides than the parent P34H mutant were sequenced and were identified to be mutants with M144 mutated to leucine (L), isoleucine (I), or valine (V). The two wells that displayed the greatest change in absorbance signal were found to harbor P34H/M144L.

The two best variants, PmST1 mutants P34H/M144L and P34H/M144V, were purified and their apparent kinetic parameters were measured for sialyltransferase, sialidase, and donor hydrolysis activities. In order to minimize the influence of donor hydrolysis, the rate of which varies between mutants, sialyltransferase kinetics were measured with a constant concentration of CMP-Neu5Ac and varied concentration of an acceptor substrate. To allow comparison to previous kinetic analysis, 4-methylumbelliferyl- β -D-lactoside (Lac β MU) was chosen as the acceptor substrate. The choice of an acceptor other than the one used for screening was preferred as it prevented potentially unintentional identification of enzymes that work only specifically for a given acceptor used in the screening.^{60, 61} The P34H/M144L mutant displayed significantly higher activity (8.7- and 5.4-fold, respectively) than Pd2,6ST or Psp2,6ST,^{54, 62, 63} the two most characterized and widely used GT80 α 2-6STs. Its α 2-6ST activity was 76% of the wild-type PmST1 α 2-3ST activity. P34H/M144L also displayed 1/3 of the donor hydrolysis activity and 1/53 of the sialidase activity of the wild-type PmST1. In contrast, P34H and P34H/M144A displayed reduced activity consistent with Pd2,3ST P7H and P7H/M117A, although accurate measurement of the P34H ST activity was made impossible by the surprising 43-fold increased donor hydrolysis activity. Using high-performance liquid chromatography (HPLC) to separate α 2-3- and α 2-6-regioisomers, it was possible to establish the linkage specificity of the mutants. The product α 2-6-sialyl linkage specificity of PmST1 P34H was enhanced from $94.0 \pm 0.4\%$ to $97.9 \pm 0.5\%$ by additional M144A mutation, similarly to the reported shift from 94.7% (P7H single mutant) to 99.6% (P7H/M117A double mutant) for Pd2,3ST (Table 1). To our delight, similar to P34H/M144A, P34H/M144L displayed a high preference for α 2-6-sialylation. The P34H/M144V mutant was found to be similarly selective for α 2-6-sialylation, though it also displayed decreased α 2-6ST activity in a manner similar to P34H/M144A but not P34H/M144L. Nevertheless, P34H/M144V displayed lower donor hydrolysis and sialidase activities than P34H/M144A.

The pH-dependence of the product sialyl linkage specificity was investigated using an HPLC assay in a pH range of 5.0 to 10.5 for both PmST1 P34H/M144L mutant and wild-type PmST1. This assay provided quantitative data instead of qualitative results observed by previous thin-layer chromatography (TLC) assay.¹⁵ In addition, instead of using 1:1 ratio of donor versus acceptor in the previous TLC assay,¹⁵ a 5:1 ration of donor versus acceptor was used. This was to minimize the interferences by donor hydrolysis and sialidase activities. As shown in Figure S1 (ESI[†]), PmST1 was found to be a predominant α 2-3ST across the entire pH range, although its α 2-3-selectivity decreased when the pH was below 7.0. Previously observed linkage specificity switch from α 2-3- to α 2-6 for PmST1 major sialyl product when the pH changed from a higher (>6.5) to a lower (<6.0) range¹⁵ could be explained by the enhanced α 2-3-sialidase activity at the lower pH range which caused the hydrolysis of the α 2-3-sialyl product rather than a change in sialyltransferase linkage specificity. For the experiments described here, the influence of the sialidase activity of PmST1 was minimized by controlling the amount of enzyme used and using a short reaction time. A similar product linkage selectivity shift observed for PmST1 trans-sialidase reactions was also due to the

[†]Electronic Supplementary Information (ESI) available: pH profiles and donor substrate promiscuity studies of PmST1 P34H/M144L. ¹H and ¹³C NMR spectra for Neu5Ac α 2-6Gal β 1-4GlcNAc β 1-3Gal β 1-4Glc β ProN₃ (Neu5Ac α 2-6LNnT β ProN₃) and Neu5Ac α 2-6Gal β 1-4GlcNAc β 1-3(Neu5Ac α 2-6)Gal β 1-4Glc β ProN₃. See DOI: 10.1039/x0xx00000x

enhanced α 2–3-sialidase activity at low pH which reduced α 2–3-sialoside concentration over time.⁵⁰ In comparison, PmST1 P34H/M144L mutant Figure S2 (ESI[†]) exhibited a much smaller shift in selectivity across the pH range. The selectivity of α 2–6-sialyl product rose from 98.0% at pH 5.0 to 98.9% at pH 10.5. This could be due to the much lower sialidase activity of PmST1 P34H/M144L mutant compared to wild-type PmST1.

It is worth mentioning that the PmST1 mutants studied including P34H, P34H/M144A, P34H/M144V, and P34H/M144L retained a similar expression level as the wild-type enzyme (100 mg L⁻¹ culture) under the same expression conditions. All mutants were routinely expressed at 90–115 mg L⁻¹ culture at 37 °C for 3 h after induction with 100 μ M of isopropyl β -D-1-thiogalactopyranoside (IPTG).

The donor substrate specificity of PmST1 P34H/M144L was tested using cytidine 5'-monophosphate-sialic acid (CMP-Sia) and derivatives generated *in situ* from sialic acids and derivatives or their six-carbon precursors using *Neisseria meningitidis* CMP-Sia synthetase (NmCSS)⁶⁴ with or without *Pasteurella multocida* sialic acid aldolase (PmAldolase)⁶⁵ in PmST1 P34H/M144L-containing one-pot multienzyme (OPME) systems. Sialic acids and derivatives as well as their precursors used were *N*-methylglycolylmannosamine (ManNGcOMe, A), mannose (B), 6-azido-6-deoxy-*N*-acetylmannosamine (ManNAc6N₃, C), 4-azido-4-deoxy-*N*-acetylmannosamine (ManNAc4N₃, D), 6-*O*-methyl-*N*-acetylmannosamine (ManNAc6OMe, E), and *N*-azidoacetylmannosamine (ManNAz, F), Neu5Ac (G), *N*-glycolylneuraminic acid (Neu5Gc, H), 8-*O*-methyl-*N*-acetylneuraminic acid (Neu5Ac8OMe, I), and 9-*O*-acetyl-*N*-acetylneuraminic acid (Neu5,9Ac₂, J). A longer (24 h) reaction time was used for one-pot three-enzyme reactions from sialic acid precursors due to the slower processes by the sialic acid aldolase-catalyzed reactions. In comparison, a shorter (1 h) reaction time was used for one-pot two-enzyme reactions from sialic acid and derivatives. Other than the reaction containing a substrate (Neu5,9Ac₂, J) with a labile 9-*O*-acetyl group which needed to be carried out at a lower pH (7.1) condition, all other reactions were carried out at pH 8.5 which was the optimal for the NmCSS activity without affecting the activities of PmAldolase⁶⁵ and PmST1 P34H/M144L significantly. Propyl azido β -lactoside (Lac β ProN₃) was used as the acceptor substrate. The formation of the product in each reaction was clearly demonstrated by the electrospray ionization high resolution mass spectrometry (ESI-HRMS) results (Figure S3, ESI[†]). Therefore, PmST1 P34H/M144L retained the donor substrate promiscuity of the wild-type PmST1 in tolerating CMP-sialic acid and derivatives containing diverse sialic acid forms.

The PmST1 P34H/M144L double mutant was further tested for sialylating LNNt β ProN₃, a compound containing both terminal and internal Gal residues, using CMP-Neu5Ac as the donor substrate. To our delight, mono-sialylated compound was the major product when 1.0 equivalent of CMP-Neu5Ac was used. In order to confirm the location of the mono-sialylation, a preparative-scale synthesis of monosialylated LNNt β ProN₃ from LNNt β ProN₃, 1.0 equivalent of Neu5Ac, and 1.5 equivalent of cytidine 5'-triphosphate (CTP) was carried out using an efficient one-pot two-enzyme sialylation system⁵ containing *Neisseria meningitidis* CMP-sialic acid synthetase (NmCSS)⁶⁴ and PmST1 P34H/M144L mutant. As shown in Figure 4, NmCSS was responsible for converting CTP and Neu5Ac to generate CMP-Neu5Ac *in situ* which was used as the donor substrate by the PmST1 P34H/

M144L mutant. Controlling the amount of enzyme used and the timing of the reaction by monitoring with high-resolution mass spectrometry (HRMS) was important to achieve high-yield production of mono-sialylated product as the major product. Reaction at 37 °C for 1 hour was found to be optimal for the formation of mono-sialylated product. The formation of disialylated product (5% yield) was also observed which was separated from monosialylated product (90% yield) by high performance liquid chromatography (HPLC) purification using a C18 column.

¹³C Nuclear magnetic resonance (NMR) data (Table S1, Figure S4–S6 ESI[†]) confirmed that the attachment of Neu5Ac in the mono-sialylated product was to the C-6 of the terminal Gal residue in LNnTβProN₃. Sialylation resulted in a significant downfield shift of the substituted carbon (a downfield shift of 2.31 ppm from 60.91 ppm in the starting tetrasaccharide to 63.22 ppm in the sialylated product) (Table S1, ESI[†]). This chemical shift change seen in Neu5Acα2–6Gal was consistent with that observed for other glycans containing the same structural component.^{66, 67} In addition, sialylation led to a significant downfield shift ($\delta = +2.31$ ppm) of C-4 in GlcNAc in the ¹³C spectra due to the steric interaction between Neu5Ac and GlcNAc in the product.

Conclusions

Regiospecificity is one of the most valued properties of glycosyltransferases, the primary catalysts for the synthesis of carbohydrates. However, changes to this property could not be screened for in a high-throughput fashion with previous glycosyltransferase screening assays. Engineering active, well expressed, and donor promiscuous enzymes for altered linkage regiospecificity is a promising new way to expand the enzyme toolbox for oligosaccharide synthesis. The microtiter plate screening assay demonstrated in this work can be adapted to work for a large variety of glycosyltransferases. With access to suitable exoglycosidases, changes to regioselectivity or stereoselectivity can be determined. The relative simplicity, low cost, and broad applicability of this assay make it a powerful new tool for the field of glycobiology. Using this screening method, a highly active α2–6ST PmST1 mutant P34H/M144L was identified to display 8.7-fold and 5.4-fold higher α2–6ST activity, respectively, than commonly used α2–6STs Pd2,6ST and Psp2,6ST. The identified PmST1 mutant P34H/M144L retained the donor substrate promiscuity of the wild-type enzyme. It allowed highly efficient selectively sialylation of the terminal Gal residue in a multigalactose-containing tetrasaccharide lacto-*N*-neotetraoside using a one-pot two-enzyme system.

Experimental Section

Materials and methods

Chemicals were purchased and used as received. NMR spectra were recorded in the NMR facility of University of California, Davis on a Bruker Avance-800 NMR spectrometer (800 MHz for ¹H, 200 MHz for ¹³C). Chemical shifts are reported in parts per million (ppm) on the δ scale. High resolution (HR) electrospray ionization (ESI) mass spectra were obtained using a Thermo Electron LTQ-Orbitrap Hybrid MS at the Mass Spectrometry Facility in the University of California, Davis. Gel filtration chromatography was performed with a column

(100 cm × 2.5 cm) packed with Bio-Gel P-2 Fine resins (Bio-Rad). Gal β pNP and sodium pyruvate were from Sigma. *N*-Acetylneuraminic acid (Neu5Ac) was from Inalco (Italy). Cytosine 5'-triphosphate (CTP) was purchased from Hangzhou Meiya Pharmaceutical Co. Ltd. Recombinant enzymes *Neisseria meningitidis* CMP-sialic acid synthetase (NmCSS),⁶⁴ *Pasteurella multocida* multifunctional α 2-3ST 1 (PmST1),¹⁵ *Photobacterium damsela* α 2-6ST (Pd2,6ST),^{16, 54} and *Photobacterium* species α 2-6ST (Psp2,6ST)⁶² were expressed and purified as described previously. Neu5Ac α 2-3Gal β pNP (2),⁴⁶ Neu5Ac α 2-6Gal β pNP (3),⁴⁶ 4-methylumbelliferyl- β -D-lactoside (Lac β MU),¹⁵ and tetrasaccharide LNnT β ProN₃⁶⁸ were prepared as described previously.

Site-directed saturation mutagenesis

PmST1 point mutant P34H and single-site saturation library P34X were generated with the QuikChange II XL Site-Directed Mutagenesis kit (Agilent Technologies) according to the manufacturer's instructions using primers P34H_f: ATCACGCTGTATTTAGATCATGCCTCCTTACCGGC, P34H_r: GCCGGTAAGGAGGCATGATCTAAATACAGCGTGAT, P34X_f: GGTCAAAAACAATCACGCTGTATTTAGATNNKGCCTCCTTACCGGCA, and P34X_r: TGCCGGTAAGGAGGCMNNATCTAAATACAGCGTGATTGTTTTTGGACC. Single-site saturation library P34H/M144X was generated via the Phusion Site-Directed Mutagenesis Kit (Thermo Scientific) using 5' phosphorylated primers M144X_f: NNKGAATATGTTGATTTAGAAAAAGAAGAAATAAAGATATTTCCGC and M144X_r: TGAGCCATCGTCATAAAGATTAACTGC. The assembled DNA was transformed into *E. coli* 10 G Electrocompetent Cells (Lucigen). Ten percent of the transformed cells were plated on LB agar plates supplemented with ampicillin in order to determine the number of total transformants. The remaining transformed cells were diluted into fresh LB media (10 g L⁻¹ tryptone, 5 g L⁻¹ yeast extract, and 10 g L⁻¹ NaCl) supplemented with ampicillin, grown overnight at 37 °C 250 rpm, and the plasmid DNA was isolated. This DNA was transformed into homemade chemically competent *E. coli* BL21(DE3) cells.

Library expression

To each well of a 96-well deep well plate was added LB media (1 mL). For each library, 88 wells were inoculated with colonies from agar plates, 4 wells were inoculated with cells expressing wild-type PmST1, and 4 wells were inoculated with cells containing empty pET22b vector. These plates were cultured for overnight at 37 °C with shaking (350 rpm), and the following day these cultures (20 μ L each) were used to inoculate cultures (1 mL each) in another 96-deep well plate. 50% glycerol solution (200 μ L) was added to each well on the original plate, and the plate was sealed and stored at -80 °C. The freshly inoculated cells were grown for four hours at 37 °C at 350 rpm before being induced with isopropyl β -D-1-thiogalactopyranoside (IPTG, 100 μ M) and grown for another four hours at 37 °C at 350 rpm. The cells were pelleted by centrifugation, resuspended in Tris-HCl (50 mM, pH 8.0) supplemented with 4 mg mL⁻¹ lysozyme, and frozen at -80 °C. After thawing at room temperature, cell debris was pelleted by centrifugation and the lysate was used for the sialyltransferase and sialidase assays.

Sialyltransferase screening

Into every other column and every other row of a clear bottom 384-well microtiter plate was added 20 μ L lysate and 20 μ L of reaction master mix to final concentrations of Gal β pNP (0.4 mM), CMP-Neu5Ac (1.2 mM), Tris-HCl (100 mM, pH 7.5), and MgCl₂ (10 mM). After one hour at 37 °C, 15 μ L of each reaction was added into two adjacent empty wells. To one of these wells was added β -galactosidase (5 μ L, 8 mU μ L⁻¹), and to the other of these wells was added β -galactosidase (5 μ L, 8 mU μ L⁻¹) and NanB (70 μ g mL⁻¹). After two hours at 37 °C, CAPS buffer (40 μ L, 500 mM, pH 9.6) was added to each well and the absorbances at 405 nm of the samples in the plate were measured using a Synergy HT microplate reader (BioTek).

Overexpression and purification

For the expression of PmST1 and its mutants, 1 L of LB media containing 100 μ g mL⁻¹ ampicillin was inoculated with 1 mL of an overnight culture of *Escherichia coli* BL21(DE3) harboring the appropriate plasmid. This culture was grown at 37 °C with vigorous shaking at 250 rpm until reaching an OD₆₀₀ of 0.8–1.0, then expression was induced with IPTG to a final concentration of 100 μ M and the protein was expressed at 37 °C with vigorous shaking at 250 rpm for another 3 hours. The cells were harvested by centrifugation at 4 °C in a Sorvall Legend RT centrifuge at 4000 rpm for 30 min. The cell pellets were resuspended in Tris-HCl (20 mL, 100 mM, pH 7.5). The cells were broken by sonication with the following method: amplitude at 65%, 10's pulse on and 20's pulse off for 18 cycles. Then the cell lysate was centrifuged at 4 °C, 8,000 rpm for 30 min, the supernatant was loaded onto a Ni²⁺-NTA affinity column at 4 °C that was pre-equilibrated with 6 column volumes of binding buffer (50 mM Tris-HCl buffer, pH 7.5, 10 mM imidazole, 0.5 M NaCl). The column was washed with 10 column volumes of binding buffer and 10 column volumes of washing buffer (50 mM of Tris-HCl buffer, pH 7.5, 50 mM of imidazole, 0.5 M of NaCl) sequentially to wash away the nonspecific binding protein. The target protein was eluted using Tris-HCl buffer (50 mM, pH 7.5) containing 200 mM of imidazole and 0.5 M NaCl. Fractions containing the purified protein were combined and dialyzed against Tris-HCl buffer (20 mM, pH 7.5) supplemented with 10% glycerol. The enzyme solutions were aliquoted, flash frozen in liquid N₂, and stored at -20 °C.

Sialyltransferase kinetics

Reactions were performed in duplicate at 37 °C for 10 to 30 minutes with Tris-HCl (100 mM, pH 8.5), MgCl₂ (10 mM), CMP-Neu5Ac (5 mM), enzyme (0.0988 μ M P34H, 0.06343 μ M P34H/M144A, 0.0339 μ M P34H/M144V, 0.0169 μ M P34H/M144L), and varying concentrations (0.5, 1.0, 2.0, and 5.0 mM) of Lac β MU as the acceptor substrate. Reactions were stopped by adding an equal volume of pre-chilled methanol. The mixtures were incubated on ice for 30 min and centrifuged at 13,000 rpm for 5 min. Supernatants were diluted with 25% acetonitrile and analyzed with a Shimadzu LC-6AD system equipped with a membrane online degasser, a temperature control unit, and a fluorescence detector (Shimadzu RF-10AXL). A reverse-phase Premier C18 column (250 mm $\times$ 4.6 mm i.d., 5 μ m particle size, Shimadzu) protected with a C18 guard column cartridge was used. The fluorophore (MU)-labeled compounds were separated with an isocratic flow of 20%

acetonitrile and were detected by excitation at 325 nm and emission at 372 nm. The apparent kinetic parameters were obtained by fitting the experimental data (the average values of duplicate assay results) into the Michaelis–Menten equation using Grafit 5.0.

Donor hydrolysis kinetics

Reactions were performed in duplicate at 37 °C for 10–30 minutes with Tris-HCl (100 mM, pH 8.5), MgCl₂ (10 mM), enzyme (0.0011 μM P34H, 0.0339 μM P34H/M144V, 0.07 μM P34H/M144A, 0.0718 μM P34H/M144L, 1.71 μM Psp2,6ST, 0.402 μM Pd2,6ST), and varying concentrations (2.0, 5.0, 10.0, and 20.0 mM) of CMP-Neu5Ac. Reactions were stopped by adding an equal volume of pre-chilled methanol. The mixtures were incubated on ice for 30 min and centrifuged at 13,000 rpm for 5 min. Supernatants were analyzed with a P/ACE™ MDQ capillary electrophoresis (CE) system equipped with a UV-Vis detector (Beckman Coulter, Fullerton, CA). The CE procedure utilized a 75 μm i.d. capillary, 25 KV/80 μA, 5 s vacuum injections, was monitored at 254 nm, and used sodium tetraborate (25 mM, pH 9.4) buffer as the running buffer. The apparent kinetic parameters were obtained by fitting the experimental data (the average values of duplicate assay results) into the Michaelis–Menten equation using Grafit 5.0.

Sialidase kinetics

Reactions were performed in duplicate at 37 °C for 10 to 30 minutes with Tris-HCl (100 mM, pH 5.5), MgCl₂ (10 mM), CMP (0.5 mM), enzyme (3.53 μM P34H/M144A, 1.1 μM P34H, 0.364 μM P34H/M144L, 207.6 μM Psp2,6ST, 33.9 μM P34H/M144V), and varying concentrations (0.5, 1.0, 2.0, and 5.0 mM) of Neu5Acα2–6LacβMU. Reactions were stopped by adding an equal volume of pre-chilled methanol. The mixtures were incubated on ice for 30 min and centrifuged at 13,000 rpm for 5 min. Supernatants were analyzed with a P/ACE™ MDQ capillary electrophoresis (CE) system equipped with a UV-Vis detector (Beckman Coulter, Fullerton, CA). The CE procedure utilized a 75 μm i.d. capillary, 25 KV/80 μA, 5 s vacuum injections, was monitored at 315 nm, and used sodium tetraborate (25 mM, pH 9.4) buffer as the running buffer. The apparent kinetic parameters were obtained by fitting the experimental data (the average values of duplicate assay results) into the Michaelis–Menten equation using Grafit 5.0.

Product sialyl linkage specificity studies

Reactions were performed in duplicate at 37 °C for 10 minutes in Tris-HCl buffer (100 mM, pH 8.5) containing MgCl₂ (10 mM), CMP-Neu5Ac (10 mM), LacβMU (2 mM), and enzyme (0.036 μM PmST1, 0.104 μM PmST1 P34H, 0.104 μM PmST1 P34H/M144A, 0.010 μM PmST1 P34H/M144L, 0.104 μM PmST1 P34H/M144V, 0.082 μM Psp2,6ST, 0.085 μM Pd2,6ST). pH Profiles for sialylation yields and product sialyl linkage percentages were also carried out for PmST1 and PmST1 P34H/M144L mutant with varied pH. Buffers used were 2-(*N*-morpholino)ethanesulfonic acid (MES) for pH 5.0–6.5, Tris-HCl for pH 7.0–9.0, and CAPS for pH 9.5–10.5. Reactions were stopped by adding an equal volume of pre-chilled methanol. The mixtures were incubated on ice for 30 min and centrifuged at 13,000 rpm for 5 min. Supernatants were analyzed with an Agilent 1290 Infinity II liquid chromatography (LC) system equipped with a UV-Vis detector. A reverse-phase Zorbax Eclipse Plus C18 Rapid Resolution HD column (50 mm × 2.1 mm i.d., 1.8 μm particle size,

Agilent) was used. The fluorophore (MU)-labeled compounds were separated with a 2 minute isocratic flow of 9% acetonitrile/91% H₂O supplemented with 0.1% TFA and were detected by absorbance at 315 nm.

High resolution mass spectrometry (HRMS)-based donor substrate specificity studies of PmST1 P34H/M144L using one-pot multienzyme sialylation systems

One-pot three-enzyme reactions were carried out at 37 °C for 24 hours in a total volume of 10 µL in Tris-HCl buffer (100 mM, pH 8.5) containing LacβProN₃ (1 mM), *N*-acetylmannosamine (ManNAc), mannose or derivatives as sialic acid precursors (1.5 mM), sodium pyruvate (5 mM), CTP (2 mM), PmAldolase (0.2 µg), NmCSS (0.15 µg), and PmST1 P34H/M144L (0.05 µg). Sialic acid precursors used were ManNGcOMe (A), mannose (B), ManNAc6N₃ (C), ManNAc4N₃ (D), ManNAc6OMe (E), and ManNAz (F).

One-pot two-enzyme reactions were carried out at 37 °C for 1 hour in a total volume of 10 µL in Tris-HCl buffer (100 mM, pH 8.5; pH 7.5 was used for the reaction containing 9-*O*-acetyl Neu5Ac) containing LacβProN₃ (1 mM), sialic acid or derivatives (1.5 mM), CTP (2 mM), NmCSS (0.15 µg), and PmST1 P34H/M144L (0.05 µg). Sialic acids and derivatives used were Neu5Ac (G), Neu5Gc (H), Neu5Ac8OMe (I), and Neu5,9Ac₂ (J).

Samples were analyzed by electrospray ionization (ESI)-HRMS using a Thermo Electron LTQ-Orbitrap Hybrid MS in a negative mode.

One-pot two-enzyme preparative-scale synthesis of sialylated products using PmST1 P34H/M144L in a one-pot two-enzyme system

To prepare sialylated L_NTβProN₃, a reaction mixture in a total volume of 4 mL containing Tris-HCl buffer (100 mM, pH 8.5), Galβ1-4GlcNAcβ1-3Galβ1-4GlcβProN₃ (L_NTβProN₃, 30 mg, 0.038 mmol), Neu5Ac (11 mg, 0.038 mmol), CTP (30 mg, 0.057 mmol), MgCl₂ (20 mM), NmCSS (1.2 mg), and PmST1 P34H/M144L (0.02 mg) were incubated in a shaker at 37 °C for 1 h. The reaction was stopped by adding 4 mL of ethanol followed by incubation at 4 °C for 30 min. After centrifugation, the supernatant was concentrated and passed through a Bio-Gel P-2 gel filtration column (water was used as an eluant). Further purification by semi-preparative HPLC using a C18 column provided monosialylated product Neu5Acα2-6Galβ1-4GlcNAcβ1-3Galβ1-4GlcβProN₃ as a white powder (37.7 mg, 90%) and disialylated product Neu5Acα2-6Galβ1-4GlcNAcβ1-3(Neu5Acα2-6)Galβ1-4GlcβProN₃ (2.7 mg, 5%), also as a white powder.

Neu5Acα2-6Galβ1-4GlcNAcβ1-3Galβ1-4GlcβProN₃. ¹H NMR (800 MHz, D₂O) δ 4.70 (d, *J* = 8.0 Hz, 1H), 4.46 (d, *J* = 8.0 Hz, 1H), 4.43 (d, *J* = 8.0 Hz, 1H), 4.42 (d, *J* = 8.0 Hz, 1H), 4.14 (d, *J* = 3.2 Hz, 1H), 3.99–3.52 (m, 25H), 3.44 (t, *J* = 6.4 Hz, 2H), 3.30 (t, *J* = 8.8 Hz, 1H), 2.65 (d, *J* = 12.0 and 4.0 Hz, 1H), 2.04 (s, 3H), 2.01 (s, 3H), 1.89 (m, 2H), 1.70 (t, *J* = 12.0 Hz, 1H). ¹³C NMR (200 MHz, D₂O) δ 174.81, 174.78, 173.43, 103.33, 102.82, 102.47, 101.98, 100.00, 81.89, 80.30, 78.22, 74.76, 74.65, 74.23, 74.13, 73.56, 72.66, 72.41, 72.29, 72.11, 71.59, 70.60, 69.84, 68.27, 68.23, 68.19, 68.08, 67.24, 63.22, 62.52, 60.84, 59.99, 59.90, 54.80, 51.76, 47.73, 39.94, 28.10, 22.16, 21.91. HRMS (ESI): *m/z* calculated for C₄₀H₆₇N₅O₂₉, *m/z* [M - H]⁻, calculated 1080.3843, found 1080.3833.

Neu5Ac α 2-6Gal β 1-4GlcNAc β 1-3(Neu5Ac α 2-6)Gal β 1-4Glc β ProN₃. ¹H NMR (800 MHz, D₂O) δ 4.70 (d, J = 8.0 Hz, 1H), 4.48 (d, J = 8.0 Hz, 1H), 4.43 (d, J = 8.0 Hz, 1H), 4.41 (d, J = 8.0 Hz, 1H), 4.16 (d, J = 3.2 Hz, 1H), 3.99–3.51 (m, 32H), 3.45 (t, J = 6.4 Hz, 2H), 3.32 (t, J = 8.8 Hz, 1H), 2.69 (d, J = 12.8 and 4.8 Hz, 1H), 2.65 (d, J = 12.8 and 4.8 Hz, 1H), 2.03 (s, 3H), 2.01 (s, 6H), 1.90 (m, 2H), 1.72 (t, J = 12.0 Hz, 1H), 1.70 (t, J = 12.0 Hz, 1H). ¹³C NMR (200 MHz, D₂O) δ 174.86 (2C), 174.84, 173.52, 173.44, 103.52, 103.20, 102.60, 101.94, 100.25, 100.09, 82.16, 80.57, 79.63, 74.61, 74.58, 74.22, 73.64, 73.22, 72.66, 72.50, 72.49, 72.37, 72.30, 71.71, 71.67, 70.69, 69.62, 68.37, 68.34, 68.32(2C), 68.18, 68.11, 67.28, 63.42, 63.30, 62.61, 62.58, 60.19, 60.11, 54.89, 51.85, 51.74, 47.84, 40.09, 40.04, 28.20, 22.25, 22.01, 21.99. HRMS (ESI): C₅₁H₈₄N₆O₃₇, m/z [M/2 - 1]⁻, calculated 635.2358, found 635.2335.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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Figure 1. Multiple functions of wild-type PmST1 including (1) $\alpha 2-3$ ST, (2) $\alpha 2-6$ ST, (3) $\alpha 2-3$ -sialidase, (4) $\alpha 2-3$ -trans-sialidase, (5) donor hydrolysis, and (6) $\alpha 2-6$ -trans-sialidase activities.

Figure 2.

A high-throughput screening assay for identifying PmST1 mutants with improved sialyltransferase activities and with regioselectivity for the formation of either α 2-3- or α 2-6-sialyl linkage. A) Reaction schemes for the assay and B) Assay validation with starting material Gal β pNP (1) and product standards Neu5Aca2-3Gal β pNP (2) and Neu5Aca2-6Gal β pNP (3) by the addition of β -galactosidase without (in well A, black columns) or with (in well B, white columns) α 2-3-sialidase SpNanB.

Figure 3.

$A_{405\text{ nm}}$ data from rescreening PmST1 P34X and P34H/M144X libraries. Empty vector controls are shown in red, wild-type control wells are shown in green, and wells identified to contain the P34H single mutant are shown in blue. Well A was treated with β -galactosidase, and lower absorbance indicates total (both $\alpha 2-3$ - and $\alpha 2-6$ -) sialylation of the acceptor. Well B was treated with both β -galactosidase and $\alpha 2-3$ -sialidase SpNanB, and lower absorbance in both well A and well B indicates $\alpha 2-6$ -sialylation of the acceptor.

Figure 4.

One-pot two-enzyme synthesis of monosialylated Neu5Ac α 2-6LNnT β ProN₃.

Table 1

Apparent kinetic parameters for PmST1 and its mutants. Data from previous reports include PmST1 (^aRef¹⁵, ^bRef⁴⁵), ^cPd2,3ST,³⁵ Pd2,6ST (^dRef⁵⁴, ^eRef⁶²), ND, not determined. nd, not detected.

Parent Enzyme	Mutant	Product sialyl linkage	Sialyltransferase				CMP-Neu5Ac Hydrolysis			Sialidase Activity		
			k_{cat} (s ⁻¹)	K_M (mM)	k_{cat}/K_M (s ⁻¹ mM ⁻¹)	Specificity (% 6')	k_{cat} (s ⁻¹)	K_M (mM)	k_{cat}/K_M (s ⁻¹ mM ⁻¹)	k_{cat} (s ⁻¹)	K_M (mM)	k_{cat}/K_M (s ⁻¹ mM ⁻¹)
Pd2,3ST	WT	α2–3	^c 24	^c 1.5	^c 16	^c < 0.6	ND	ND	ND	ND	ND	ND
	P7H	α2–6	^c 16	^c 3.8	^c 4.3	^c 94.7	ND	ND	ND	ND	ND	ND
	P7H/M117A	α2–6	^c 18	^c 6.4	^c 2.8	^c > 99.6	ND	ND	ND	ND	ND	ND
PmST1	WT	α2–3	^a 47	^a 1.4	^a 34	3.9 ± 0.8	^b 27 ± 1	^b 1.5 ± 0.2	^b 18	^a 230	^a 24	^a 9.5
	P34H	α2–6	8.6 ± 1.6	0.6 ± 0.4	13.5	94.0 ± 0.4	925.5 ± 66.7	1.2 ± 0.5	771.3	1.2 ± 0.1	13.7 ± 1.5	0.088
	P34H/M144A		9.0 ± 1.3	1.7 ± 0.5	5.3	97.9 ± 0.5	34.5 ± 0.3	2.3 ± 0.1	15.1	0.51 ± 0.03	15.1 ± 1.1	0.034
	P34H/M144L		20.0 ± 4.4	0.8 ± 0.2	26	98.7 ± 0.1	30.5 ± 0.8	3.4 ± 0.1	8.9	1.14 ± 0.1	6.4 ± 0.6	0.178
	P34H/M144V		5.5 ± 0.4	1.4 ± 0.2	3.9	98.5 ± 0.1	62.4 ± 7.6	10.5 ± 2.7	6.0	0.27 ± 0.019	14.9 ± 1.3	0.018
Pd2,6ST	WT		^d 2.3 ± 0.1	^d 0.8 ± 0.1	^d 3	nd	6.7 ± 1.6	44.9 ± 14.2	0.15	^e 0.14 ± 0.01	^e 7.6 ± 0.5	^e 0.018
Psp2,6ST	WT		^f 1.7 ± 0.2	^f 0.4 ± 0.1	^f 4.8	nd	^g 4.8 ± 0.5	^g 17 ± 4	^g 0.28	(7.9 ± 0.3) × 10 ⁻³	4.7 ± 0.3	0.0017