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Exploring the broad nucleotide triphosphate and sugar-1-phosphate specificity of thymidyltransferase Cps23FL from *Streptococcus pneumoniae* serotype 23F†

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Glucose-1-phosphate thymidyltransferase (Cps23FL) from *Streptococcus pneumoniae* serotype 23F is the initial enzyme that catalyses the thymidyl transfer reaction in prokaryotic deoxythymidine diphosphate-L-rhamnose (dTDP-Rha) biosynthetic pathway. In this study, the broad substrate specificity of Cps23FL towards six glucose-1-phosphates and nine nucleoside triphosphates as substrates was systematically explored, eventually providing access to nineteen sugar nucleotide analogs.

1. Introduction

In past decades, many carbohydrates have been developed as drug targets due to their important pharmaceutical and pharmacological bioactivities.¹ To generate novel therapeutic molecules, glycorandomization is one of the most popular strategies to produce natural and unnatural glycosylated metabolites,² which takes advantage of the glycosyltransferases to generate diverse libraries of glycosylated biomolecules.^{3–5} Although several examples of glycorandomization have been reported,^{6,7} the deficiency of sugar precursors (usually sugar nucleotides, (d)NDP-sugars) greatly limits the application of glycorandomization. Thus, the efficient synthesis of (d)NDP-sugars has been becoming an attractive and hot topic.^{8–15}

Although chemical synthesis of sugar nucleotides has been developed,¹¹ the fastidious manipulations and low yields hindered its broad application. In contrast, enzymatic synthesis of sugar nucleotides is particularly attractive owing to its high synthetic efficiency and high regio-/stereo-specificity.^{14,15} Numerous enzymes that involved into the biosynthetic pathways of sugar nucleotides have been discovered and applied to synthesize sugar nucleotides *in vitro*.^{14,15} Among of them, a thymidyltransferase (RmlA) has been found to catalyse the first

sequential reaction in the biosynthetic pathway of deoxythymidine diphosphate L-rhamnose (dTDP-Rha),¹⁶ in which RmlA converts α -D-glucose-1-phosphate (Glc-1-P) and deoxythymidine triphosphate (dTTP) into dTDP-D-glucose (dTDP-Glc) and pyrophosphate (PPi) (Fig. 1). This enzyme reaction presumably proceeded *via* a bi-bi ordered catalytic mechanism that formed a trigonal bipyrimidal phosphoryl ternary complex.^{17–19} RmlA was attracted more interest because it displayed unusual promiscuity toward both dozens of sugar phosphates and eight natural nucleotide triphosphates.^{20,21} RmlA and its mutant have been applied to prepare up to one hundred of (d)NDP-sugars *in vitro*.^{19,22–24} In our recent report, Cps23FL, a homolog of RmlA that derived from *Streptococcus pneumoniae* serotype 23F, exhibited the excellent activity on conversion of Glc-1-P and dTTP into dTDP-Glc and PPi.²⁵ Thus, its potential to synthesize various sugar nucleotides is worthwhile for exploitation. In this work, the substrate specificity of Cps23FL was systematically investigated with six sugar-1-phosphates (sugar-1-Ps) and nine nucleotide triphosphates (NTPs) in order to explore the potential ability of recombinant wildtype Cps23FL for synthesizing those important and rare sugar nucleotides *in vitro*.

2. Results and discussion

We first investigated the substrate specificity of Cps23FL towards various (d)NTPs using Glc-1-P as the sugar-1-P

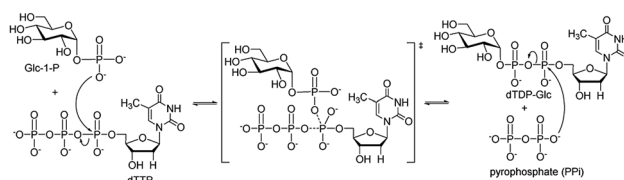


Fig. 1 The presumed mechanism of the reaction catalysed by RmlA.

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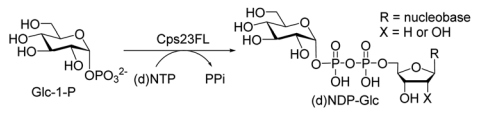
† Electronic supplementary information (ESI) available: Experimental procedures, MS and NMR data and spectra of the synthetic (d)NDP-sugars. See DOI: 10.1039/d0ra05799a

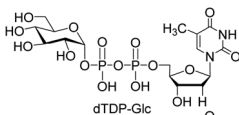
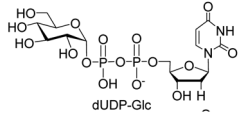
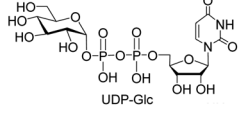
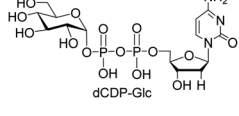
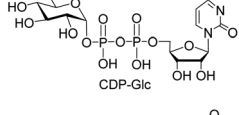
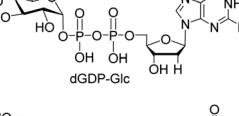
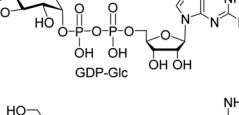
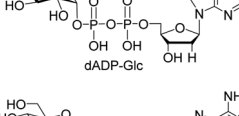
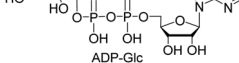
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substrate. Nine (d)NTPs were examined into the reaction systems, respectively (Table 1). Briefly, enzymatic reactions were commenced with the addition of purified Cps23FL (6 or 60 μM) to a solution containing HEPES buffer (30 mM, pH 7.5), MgCl_2 (5 mM), YIPP (2 U mL^{-1}), (d)NTPs (3 mM) and Glc-1-P (5 mM) in

Table 1 The profiles of the enzymatic reactions of various (d)NTPs and Glc-1-P catalysed by Cps23FL



Entry	(d)NTP	(d)NDP-Glc	Yield ^a (yield) ^b
1	dTTP		86% (93%)
2	dUTP		88% (95%)
3	UTP		40% (94%)
4	dCTP		75% (93%)
5	CTP		35% (87%)
6	dGTP		53% (89%)
7	GTP		NP ^c (91%) ^d
8	dATP		27% (73%) ^d
9	ATP		13% ^d (78%) ^d

^a Yields obtained in 6 μM of Cps23FL. ^b Yields obtained in 60 μM of Cps23FL. ^c No product detected by HPLC within 48 h. ^d Enzymatic reaction time was 6 h.

a final volume of 100 μL . Then, the resulting reactions were incubated at 37 $^{\circ}\text{C}$ for 1 h and quenched in boiling water bath for 30 s followed by centrifugation at 12 300 rpm for 10 min. The (d)NDP-Glc product of each enzymatic reaction was detected and isolated by HPLC (DionexCarboPacTM PA-100, 0–1.0 M ammonium acetate eluent) (Table S1†). The yields of (d)NDP-Glcs were depicted in Table 1. In 6 μM enzymatic reaction system, Cps23FL exhibited broad substrate specificities towards deoxy-nucleoside triphosphate (dTTP, dUTP, dCTP, and dGTP) (Table 1, entries 1, 2, 4, 6) and afforded good to high isolate yields of dNDP-Glcs (>50%) except that of dATP (only 27%) (Table 1, entry 8). However, Cps23FL showed only low or none activities toward UTP, CTP, GTP and ATP substrates (Table 1, entries 3, 5, 7 and 9) under this enzyme concentration even with prolonging reaction time to 48 h (for GTP). Accordingly, the influence of different concentration of NTP (0, 1.5, 3.0, 6.0, 12.0, 24.0 mM) and Csp23FL (0, 0.06, 0.6, 6.0, 30, 60 μM) on the product conversion was investigated using ATP as a model substrate (Fig. 2). It was disclosed that the high conversion of ADP-Glc within 6 h was achieved at 3.0 mM concentration of ATP (Fig. 2A), and also efficiently improved with the increased Cps23FL concentration (Fig. 2B). As shown in Table 1, under 60 μM enzyme concentration, all tested nucleoside triphosphates could be commendably recognized by Cps23FL and completely converted into their corresponding (d)NDP-Glcs in high yields (73–95%). Our results indicated Cps23FL had a notable substrate tolerance and the changes on both nucleobase and substituent group on the C-2 position of ribose residue could not influence its enzymatic activity. Therefore, Cps23FL preferred both pyrimidine nucleoside and purine nucleoside whether in nucleotide form or their deoxy form under high enzyme concentration, which was consistent with those results reported by RmlA.^{20,21} All the prepared (d)NDP-Glcs were well characterized by MS and NMR spectrometry, and the characterization data of dTDP-Glc,²⁵ dUDP-Glc,^{26,27} UDP-Glc,^{10,26} CDP-Glc,^{10,26} GDP-Glc,²⁶ and ADP-Glc^{26,27} were in good agreement with those previously reported. More remarkably, dCDP-Glc, dGDP-Glc, and dADP-Glc were characterized first time by NMR spectrometry in this study.²⁸

Next, the substrate tolerance of Cps23FL for sugar-1-Ps was investigated with dTTP as pyrophosphate donor. The sugar-1-Ps tested in this study were the native substrate Glc-1-P, galactose-

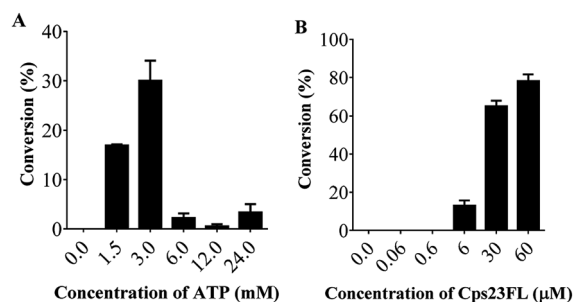
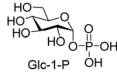
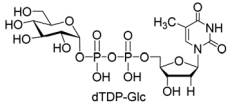
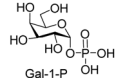
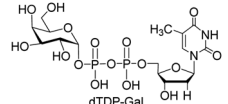
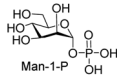
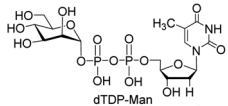
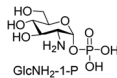
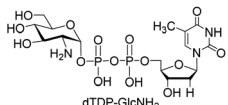
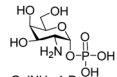
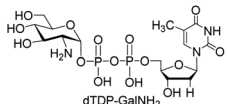
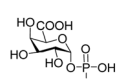
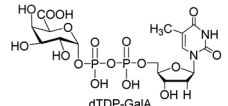


Fig. 2 The influence of different concentration of (A) ATP and (B) Cps23FL on the enzymatic reaction of Cps23FL with Glc-1-P and ATP within 6 h.



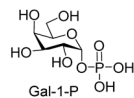
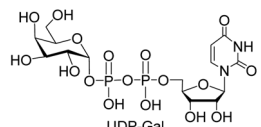
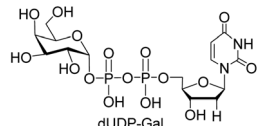
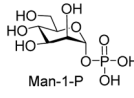
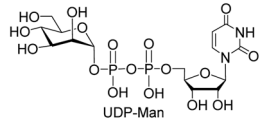
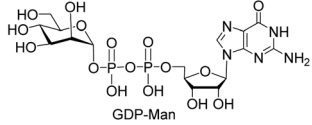
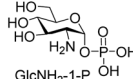
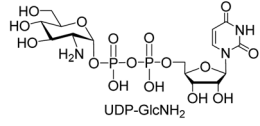
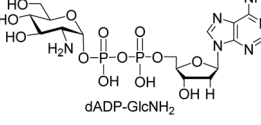
1-phosphate (Gal-1-P), mannose-1-phosphate (Man-1-P), glucosamine-1-phosphate (GlcNH₂-1-P), galactosamine-1-phosphate (GalNH₂-1-P), and galacturonic acid-1-phosphate (GalA-1-P), respectively. The enzymatic reactions were performed in 60 μM of Cps23FL with the same reaction condition described above. The reaction results shown in Table 2 indicated that Cps23FL exerted different activities for sugar-1-Ps in the following order: Glc-1-P ≈ Gal-1-P > Man-1-P > GlcNH₂-1-P ≫ GalNH₂-1-P. Cps23FL could efficiently catalyse both Glc-1-P and Gal-1-P to form dTDP-Glc/Gal in excellent yields (Table 2, entries 1 and 2). Furthermore, Man-1-P could be also well tolerated by Cps23FL to yield dTDP-Man in a high yield of 88% (Table 2, entry 3). Nevertheless, GlcNH₂-1-P was transformed into dTDP-GlcNH₂ by Cps23FL in good conversion rate (71%) whereas only 5% for GalNH₂-1-P and no activity for GalA-1-P (Table 2, entries 4, 5 and 6). All these results indicated that the configuration change at C-2 and C-4 positions (Man-1-P, Gal-1-P) or amino group substitution at C-2 position (GlcNH₂-1-P) could not affect their recognition by Cps23FL, while the configuration change at both the configuration change at C-4 position and substitution at C-2 or C-6 positions (GalNH₂-1-P, GalA-1-P) might greatly decrease their enzymatic activities. Our findings were commendably supported by profile of the crystal structure of RmlA and Glc-1-P complex (PDB entry 1G23), in

Table 2 Yields for the conversion of different sugar-1-Ps and dTTP by 60 μM of Cps23FL

Entry	Sugar-1-P	dTDP-sugar	Yield
1			93%
2			91%
3			88%
4			71% ^a
5			5% ^b
6			NP ^b

^a Enzymatic reaction time was 6 h. ^b Detection by HPLC with prolonged reaction time to 48 h.

Table 3 Yields of various hybrid (d)NDP-sugars catalysed by 60 μM of Cps23FL within 48 h

Entry	Sugar-1-P	(d)NTP	(d)NDP-sugar	Yield
1		UTP		86%
2		dUTP		83%
3		UTP		78%
4		GTP		13% ^a
5		UTP		12% ^a
6		dATP		21% ^a

^a Detection by HPLC.

which the hydroxyl groups on C-2, C-4 and C-6 positions of glucose residue formed hydrogen bonds and van der Waals interacting with RmlA.^{6,17} Similarly, all the synthetic dTDP-sugars were characterized by MS and NMR spectrometry. The spectra data of dTDP-Gal, and dTDP-Man were well agreed with those previously reported.¹⁰ dTDP-GlcNH₂ was characterized by NMR spectrometry,²⁸ whilst dTDP-GalNH₂ was only acquired by ESI-HRMS spectrum due to only trace amount of product obtained in reaction.

We further investigated the potential synthetic ability of Cps23FL for preparation of other (d)NDP-sugars with both non-native sugar-1-P analogues and (d)NTPs as substrates. Six (d)NDP-sugar analogues were synthesized with diverse yields (Table 3). Surprisingly, Cps23FL exhibited good activity toward Gal-1-P and (d)UTP substrates within 48 h, providing UDP-Gal and dUDP-Gal in an 86% and 83% isolated yield, respectively (Table 3, entries 1 and 2). For Man-1-P and UTP or GTP substrate, however, different conversion yields of UDP-Man (78%) and GDP-Man (13%) were observed after incubation reaction time of 48 h (Table 3, entries 3 and 4). Previous study has suggested that only RmlA mutant (Q24S) could catalyse the



reactions of Man-1-P and GTP to synthesize GDP-Man.¹⁹ This study was the first time to report the synthesis of GDP-Man by recombinant wildtype Cps23FL. Similarly, Cps23FL could also catalyse GlcNH₂-1-P with UTP or dATP to produce UDP-GlcNH₂ or dADP-GlcNH₂ in low yields (<21%) with the incubation time to 48 h (Table 3, entries 5 and 6). These results indicated its great potential on synthesis of diverse (d)NDP-sugars using non-native substrates. The synthetic UDP-Gal and UDP-Man were well confirmed by NMR and ESI-HRMS spectrometry,^{23,26} and dUDP-Gal was characterized first time by NMR spectrometry.²⁸

3. Conclusion

In summary, the broad substrate specificity of the recombinant thymidyltransferase Cps23FL cloned from *S. pneumonia* serotype 23F against six sugar-1-Ps and nine (d)NTPs was investigated in detail in this study. It was disclosed that Cps23FL exhibited much better recognition towards (d)NTP substrate than sugar-1-P substrate. As a result, nineteen sugar nucleotides were eventually constructed by Cps23FL-catalysed transformation reaction, and fifteen of them were available isolated by semi-preparation HPLC technique in milligram scales in high yields of 70–95%. However, four of them including GDP-Man, dTDP-GalNH₂, UDP-GlcNH₂, and dADP-GlcNH₂ were obtained in very low yields of less 22%. This can presumably be attributed to the significant structural alternation of both nucleoside triphosphate and sugar 1-phosphate, which is beyond the recognizable substrate tolerance of Cps23FL enzyme and thus likely weakens their binding ability at the active site.¹⁹ Moreover, the synthetic (d)NDP-sugars were proved to be stable enough within 48 h that evidenced by the HPLC observation of no more than 5% degradation of (d)NDP-sugars in reaction system. Furthermore, it was reported first time that wildtype Cps23FL could catalyse transformation of GDP-Man (even in low yield of 13%), which could only be achieved by RmlA mutant in previous study.¹⁹ This finding would allow us to broaden Cps23FL activity through enzyme engineering technique in case of improving synthetic efficacy of GDP-Man. Overall, this study provides a convenient route for the efficient preparation of a library of sugar nucleotides by wildtype Cps23FL, which will greatly facilitate glycosyltransferase-based glycorandomization synthesis of complex glycosylated metabolites.

Conflicts of interest

The authors declare no conflict of interest.

Acknowledgements

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- 28 Spectroscopic data of **dCDP-Glc**: ^1H NMR (600 MHz, D_2O): δ 7.91 (d, $J = 7.8$ Hz, 1H, $-\text{CH}=\text{}$), 6.13 (t, $J = 6.6$ Hz, 1H, H-1), 6.03 (d, $J = 7.8$ Hz, 1H, $-\text{CH}=\text{}$), 5.42 (dd, $J = 7.2$, 3.0 Hz, 1H, H-1'), 4.43 (br s, H-3), 4.09–3.97 (m, 3H, H-4, H-5a,b), 3.72 (br d, $J = 9.6$ Hz, 1H, H-5'), 3.68 (d, $J = 12.6$ Hz, 1H, H-6a'), 3.63–3.56 (m, 2H, H-3', H-6b'), 3.35 (br d, $J = 9.6$ Hz, 1H, H-2'), 3.29 (t, $J = 9.6$ Hz, 1H, H-4'), 2.32–2.25 (m, 1H, H-2), 2.20–2.13 (m, 1H, H-2); ^{13}C NMR (150 MHz, D_2O): δ 162.83, 142.76, 95.77, 95.46 (d, $J = 6.6$ Hz, C-1'), 86.19 (C-1), 85.64 (d, $J = 9.0$ Hz, C-4), 72.75 (C-3'), 72.68 (C-5'), 71.51 (d, $J = 8.4$ Hz, C-2'), 70.59 (C-3), 69.05 (C-4'), 65.11 (C-5), 60.18 (C-6'), 39.41 (C-2); ^{31}P NMR (243 MHz, D_2O): δ -11.34 (d, $J = 20.8$ Hz), -12.99 (d, $J = 20.8$ Hz); ESI-TOF HRMS m/z calcd for $\text{C}_{15}\text{H}_{24}\text{N}_3\text{O}_{15}\text{P}_2$ $[\text{M} - \text{H}]^{-1}$ 548.0688, found 548.0616. **dGDP-Glc**: ^1H NMR (600 MHz, D_2O): δ 7.92 (s, 1H, $-\text{CH}=\text{}$), 6.15 (t, $J = 7.2$ Hz, 1H, H-1), 5.40 (dd, $J = 6.6$, 3.0 Hz, 1H, H-1'), 4.58 (br s, 1H, H-3), 4.08 (br s, 1H, H-4), 4.01–3.96 (m, 2H, H-5a,b), 3.74–3.69 (m, 1H, H-5'), 3.67 (d, $J = 12.6$ Hz, 1H, H-6a'), 3.62–3.54 (m, 2H, H-3', H-6b'), 3.36–3.31 (m, 1H, H-2'), 3.27 (t, $J = 9.6$ Hz, 1H, H-4'), 2.68–2.60 (m, 1H, H-2a), 2.38–2.30 (m, 1H, H-2b); ^{13}C NMR (150 MHz, D_2O): δ 158.89, 153.75, 151.34, 137.52, 116.11, 95.45 (d, $J = 7.5$ Hz, C-1'), 85.46 (d, $J = 9.0$ Hz, C-4), 83.49 (C-1), 72.70 (C-3'), 72.66 (C-5'), 71.49 (d, $J = 9.0$ Hz, C-2'), 71.22 (C-3), 69.07 (C-4'), 65.37 (C-5), 60.19 (C-6'), 38.37 (C-2); ^{31}P NMR (243 MHz, D_2O): δ -11.22 (d, $J = 20.6$ Hz), -12.99 (d, $J = 20.6$ Hz); ESI-TOF HRMS m/z calcd for $\text{C}_{16}\text{H}_{24}\text{N}_5\text{O}_{15}\text{P}_2$ $[\text{M} - \text{H}]^{-1}$ 588.0750, found 588.0680. **dADP-Glc**: ^1H NMR (600 MHz, D_2O): δ 8.32 (s, 1H), 8.09 (s, 1H), 6.36 (t, $J = 6.6$ Hz, 1H, H-1), 5.42–5.39 (m, 1H, H-1'), 4.59 (s, 1H, H-3), 4.14 (s, 1H, H-4), 4.05–3.95 (m, 2H, H-5a,b), 3.71 (d, $J = 9.6$ Hz, 1H, H-5'), 3.66 (d, $J = 12.6$ Hz, 1H, H-6a'), 3.60 (t, $J = 9.6$ Hz, 1H, H-3'), 3.56 (dd, $J = 12.6$, 4.2 Hz, 1H, H-6b'), 3.33 (d, $J = 9.6$ Hz, 1H, H-2'), 3.26 (t, $J = 9.6$ Hz, 1H, H-4'), 2.70–2.64 (m, 1H, H-2), 2.47–2.41 (m, 1H, H-2); ^{13}C NMR (150 MHz, D_2O): δ 155.11, 152.05, 148.63, 139.97, 118.55, 95.45 (d, $J = 6.6$ Hz, C-1'), 85.65 (d, $J = 9.0$ Hz, C-4), 83.67 (C-1), 72.73 (C-5'), 72.67 (C-3'), 71.52 (d, $J = 8.7$ Hz, C-2'), 71.14 (C-3), 69.10 (C-4'), 65.35 (d, $J = 6.0$ Hz, C-5), 60.22 (C-6'), 38.97 (C-2); ^{31}P NMR (243 MHz, D_2O): δ -11.22 (d, $J = 20.7$ Hz), -12.97 (d, $J = 20.7$ Hz); ESI-TOF HRMS m/z calcd for $\text{C}_{16}\text{H}_{24}\text{N}_5\text{O}_{14}\text{P}_2$ $[\text{M} - \text{H}]^{-1}$ 572.0800, found 572.0728. **dTDP-GlcNH₂**: ^1H NMR (600 MHz, D_2O): δ 7.55 (s, 1H, $-\text{CH}=\text{}$), 6.17 (t, $J = 7.2$ Hz, 1H, H-1), 5.66 (dd, $J = 6.6$, 3.0 Hz, 1H, H-1'), 4.44 (br s, 1H, H-3), 4.05–3.98 (m, 3H, H-4, H-5a,b), 3.78–3.72 (m, 2H, H-3', H-5'), 3.70 (d, $J = 12.6$ Hz, 1H, H-6a'), 3.64 (dd, $J = 12.6$, 3.6 Hz, 1H, H-6b'), 3.38 (t, $J = 9.6$ Hz, 1H, H-4'), 3.18 (br d, $J = 10.2$ Hz, 1H, H-2'), 2.25–2.16 (m, 2H, H-2a,b), 1.75 (s, 3H, $-\text{CH}_3$); ^{13}C NMR (150 MHz, D_2O): δ 166.49, 151.64, 137.21, 111.62, 92.72 (d, $J = 4.5$ Hz, C-1'), 85.13 (d, $J = 9.0$ Hz, C-4), 84.94 (C-1), 73.11 (C-5'), 70.82 (C-3), 69.74 (C-3'), 68.94 (C-4'), 65.51 (d, $J = 4.5$ Hz, C-5), 59.86 (C-6), 54.04 (d, $J = 9.0$ Hz, C-2'), 38.49 (C-2), 11.54 ($-\text{CH}_3$); ^{31}P NMR (243 MHz, D_2O): δ -11.34 (d, $J = 21.0$ Hz), -13.75 (d, $J = 21.0$ Hz); ESI-TOF HRMS m/z calcd for $\text{C}_{16}\text{H}_{26}\text{N}_3\text{O}_{15}\text{P}_2$ $[\text{M} - \text{H}]^{-1}$ 562.0850, found 562.0776. **dUDP-Gal**: δ 7.80 (d, $J = 7.8$ Hz, 1H, $-\text{CH}=\text{}$), 6.18 (t, $J = 6.6$ Hz, 1H, H-1), 5.81 (d, $J = 7.8$ Hz, 1H, $-\text{CH}=\text{}$), 5.45 (dd, $J = 7.2$, 3.6 Hz, 1H, H-1'), 4.50–4.44 (m, 1H, H-3), 4.08–3.98 (m, 4H, H-4, H-5a,b, H-5'), 3.88 (d, $J = 3.0$ Hz, 1H, H-4'), 3.76 (dd, $J = 10.2$, 3.0 Hz, 1H, H-3'), 3.67–3.63 (m, 1H, H-2'), 3.63–3.55 (m, 2H, H-6a,b'), 2.28–2.19 (m, 2H, H-2a,b); ^{13}C NMR (150 MHz, D_2O): δ 166.23, 151.55, 141.81, 102.40, 95.74 (d, $J = 7.5$ Hz, C-1'), 85.42 (d, $J = 9.0$ Hz, H-4), 85.32 (C-1), 71.81 (C-5'), 70.79 (C-3), 69.23 (C-3'), 69.01 (C-4'), 68.33 (d, $J = 9.0$ Hz, C-2'), 65.29 (d, $J = 6.0$ Hz, C-5), 60.91 (C-6'), 38.76 (C-2); ^{31}P NMR (243 MHz, D_2O): δ -11.20 (d, $J = 19.4$ Hz), -12.77 (d, $J = 19.4$ Hz); ESI-TOF HRMS m/z calcd for $\text{C}_{15}\text{H}_{24}\text{N}_2\text{O}_{16}\text{P}_2$ $[\text{M} - \text{H}]^{-1}$ 549.0528, found 565.0449.

