

Cite this: *Org. Biomol. Chem.*, 2011, **9**, 1799

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PAPER

Characterization of the TDP-D-ravidosamine biosynthetic pathway: one-pot enzymatic synthesis of TDP-D-ravidosamine from thymidine-5-phosphate and glucose-1-phosphate†

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Received 8th October 2010, Accepted 29th November 2010

DOI: 10.1039/c0ob00854k

Ravidomycin V and related compounds, e.g., FE35A-B, exhibit potent anticancer activities against various cancer cell lines in the presence of visible light. The amino sugar moieties (D-ravidosamine and its analogues, respectively) in these molecules contribute to the higher potencies of ravidomycin and analogues when compared to closely related compounds with neutral or branched sugars. Within the ravidomycin V biosynthetic gene cluster, five putative genes encoding NDP-D-ravidosamine biosynthetic enzymes were identified. Through the activities of the isolated enzymes *in vitro*, it is demonstrated that *ravD*, *ravE*, *ravIM*, *ravAMT* and *ravNMT* encode TDP-D-glucose synthase, TDP-4-keto-6-deoxy-D-glucose-4,6-dehydratase, TDP-4-keto-6-deoxy-D-glucose-3,4-ketoisomerase, TDP-3-keto-6-deoxy-D-galactose-3-aminotransferase, and TDP-3-amino-3,6-dideoxy-D-galactose-*N,N*-dimethyl-transferase, respectively. A protocol for a one-pot enzymatic synthesis of TDP-D-ravidosamine has been developed. The results presented here now set the stage to produce TDP-D-ravidosamine routinely for glycosylation studies.

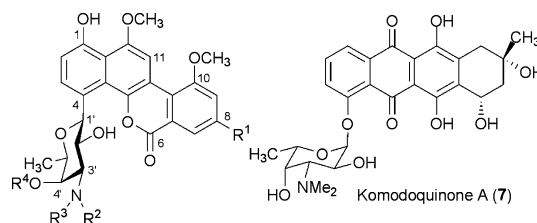
Introduction

Glycosylated microbial natural products possess a wide range of pharmaceutical properties such as anticancer, antiviral, antifungal or antibiotic.¹ The sugar moieties of these molecules are often deoxygenated to various degrees, and are appended to the backbone through *O*- or *C*-, and rarely through *N*- or *S*-glycosidic linkages.^{2–4} Structural and functional studies have demonstrated that the sugar moieties contribute significantly to the biological activities of the parent compounds.

Due to the large structural diversity of deoxysugars, and due to their huge importance for biological activities, much attention has been given to elucidate their biosynthetic pathways.^{5–13} Recent progress in understanding these pathways has proven that an array of glycosylated natural products can be generated through enzymatic glycodiversification.^{14–17} However, the successful application of this concept requires an in-depth knowledge of various deoxysugar biosynthetic pathway enzymes including their catalytic activities, their preferred substrates, and their flexibility towards unnatural substrates. Additionally, glycodiversification of natural products also requires glycosyltransferases (GTs) with sugar-donor substrate flexibility, allowing the attachment of various sugar residues to target aglycones. Although gene

sequences for dozens of GTs are known, only a marginal number of these enzymes have been fully characterized. Unavailability of functional GTs, or their acceptor or sugar donor substrates continue to contribute to the slow progress of GT studies.

Ravidomycin V (**1**) and deacetylavidomycin V (**2**) are potent antitumor antibiotics produced by *Streptomyces ravidus*.^{18,19} Both antibiotics contain a common angucyclinone-derived tetracyclic aromatic benzo[*d*]naphtho[1,2-*b*]pyran-6-one core moiety, which is connected to their respective sugar residues at C-4 position through a C–C bond (Fig. 1). Deacetylavidomycin V (**2**) contains D-ravidosamine (*N,N*-dimethyl-3-amino-3,6-dideoxy-D-galactose) and exhibits superior anticancer/antibacterial activities



Ravidomycin V (**1**) $R^1 = \text{CH}=\text{CH}_2$, $R^2 = R^3 = \text{CH}_3$, $R^4 = (\text{CO})\text{CH}_3$
 Deacetylavidomycin V (**2**) $R^1 = \text{CH}=\text{CH}_2$, $R^2 = R^3 = \text{CH}_3$, $R^4 = \text{H}$
 Ravidomycin M (**3**) $R^1 = \text{CH}_3$, $R^2 = R^3 = \text{CH}_3$, $R^4 = (\text{CO})\text{CH}_3$
 Deacetylavidomycin M (**4**) $R^1 = \text{CH}_3$, $R^2 = R^3 = \text{CH}_3$, $R^4 = \text{H}$
 FE35A (**5**) $R^1 = \text{CH}=\text{CH}_2$, $R^2 = \text{CH}_3$, $R^3 = (\text{CO})\text{CH}_3$, $R^4 = \text{H}$
 FE35B (**6**) $R^1 = \text{CH}=\text{CH}_2$, $R^2 = \text{H}$, $R^3 = R^4 = (\text{CO})\text{CH}_3$

Fig. 1 Structures of natural products comprising ravidosamine sugar moiety.

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† Electronic supplementary information (ESI) available: Full details of *in vitro* experimentation and NMR data. See DOI: 10.1039/c0ob00854k

than ravidomycin V (**1**) itself. The 4'-hydroxyl group of D-ravidosamine is acetylated in ravidomycin V. Ravidomycin M (**3**) and deacetylavidomycin M (**4**) possess a methyl side chain at C-8 position instead of the vinyl side chain present in the major congeners **1** and **2**. The enantiomeric sugar L-ravidosamine was found in the anthracycline antibiotic komodoquinone A (**7**).^{20,21} The substitution pattern specifically at the 3'-amino and 4'-hydroxyl groups of D-ravidosamine are different in the other reported ravidomycin analogues FE35A (**5**) and FE35B (**6**). Gilvocarcin V,²² chrysomycin V,²³ Mer 1020 dA-D,²⁴ BE 12406 A-B,^{25,26} and polycarcin V²⁷ are other closely related analogues, differing from ravidomycin mainly because of their neutral deoxysugar residues. Although the biological activities of these compounds have never been investigated in direct comparison, their biological activity data revealed that amino sugar containing gilvocarcin analogues show increased antibiotic/anticancer potency when compared to compounds with neutral or branched neutral sugar moieties.²⁸ Thus, investigation of the deoxysugar biosynthetic enzymes responsible for the production of D-ravidosamine, and characterization of the glycosyltransferase that attaches the sugar to the polyaromatic backbone are particularly important. An in-depth characterization of these enzymes may provide an opportunity to generate new ravidomycin analogues with altered aminosugar residues for structure-activity relationship (SAR) studies.

To understand the biosynthesis of ravidomycin V, we recently cloned and sequenced its biosynthetic gene cluster from *Streptomyces ravidus*.²⁹ The deoxysugar biosynthetic genes and associated glycosyltransferase were found in the cluster spanning a 6 kb region (Fig. 2). Bioinformatics analysis combined with heterologous expression experiments suggested that five genes, *ravD*, *ravE*, *ravIM*, *ravAMT* and *ravNMT*, encode enzymes involved in the conversion of D-glucose-1-phosphate (**8**) to NDP-D-ravidosamine, the proposed sugar donor substrate for the glycosylation reaction. Similarly, the product of *ravGT* has been proposed to be the C-glycosyltransferase that links D-ravidosamine to the polyketide-derived backbone defuco-gilvocarcin V.



Fig. 2 A section of the ravidomycin gene cluster spanning the putative TDP-D-ravidosamine (**17**) biosynthetic genes (*ravD*, *E*, *IM*, *AMT* and *NMT*).

Suzuki and co-workers have developed a synthetic route for the production of D-ravidosamine.³⁰ Multiple synthetic steps involved in this protocol and a low overall yield are major drawbacks of this approach. Activation of the sugar with a suitable nucleosyl-diphosphate (NDP) group poses another challenge towards the synthetic preparation of the donor substrate for RavGT. Recent development in understanding of deoxysugar biosynthetic pathways has made it possible to synthesize a target NDP-activated sugar *in vitro* through the activities of suitable enzymes. This approach is superior compared to the chemical total synthesis as it is concerted, less time-consuming, stereo- and regioselective, and offers high yields of product in an environmentally friendly procedure. We herein, report the preparation of thymidine diphosphate-activated ravidosamine (TDP-D-ravidosamine, **17**) from glucose-1-phosphate **8** utilizing enzymes of the ravidosamine biosynthetic

pathway. We demonstrate through *in vitro* assays and NMR analyses that *ravD*, *ravE*, *ravIM* and *ravAMT* encode TDP-D-glucose synthase, TDP-D-glucose-4,6-dehydratase, TDP-4-keto-6-deoxy-D-glucose-3,4-ketoisomerase and TDP-3-keto-6-deoxy-D-galactose-3-aminotransferase, respectively. Similarly, the product of *ravNMT* was characterized as TDP-3-amino-3,6-dideoxy-D-galactose-*N,N*-dimethyltransferase. Characterization of these five deoxysugar biosynthetic enzymes unambiguously established the biosynthetic pathway of **17**. In addition, we report a one-pot enzymatic synthetic protocol for the production of **17**. The results presented in this study lay the foundation for the *in vitro* characterization of the glycosyltransferase RavGT involved in ravidomycin biosynthesis.

Results and discussion

Expression and characterization of RavD and RavE

Basic local sequence alignment search tool (BLAST) analysis of RavD revealed high sequence similarity to several NDP-D-glucose synthases from deoxysugar biosynthetic pathways. RavD has the highest sequence homology to the proposed NDP-glucose synthase ChryD (83% identity/90% similarity) from the chrysomycin biosynthetic pathway.²⁹ The other proteins that displayed high amino acid homology with RavD were NDP-D-glucose synthases GilD (75%/86%),³¹ MtmD (76%/86%),³² SsfS1 (74%/85%)³³ and StrD (71%/83%)³⁴ from the gilvocarcin, mithramycin, SF2575 and streptomycin biosynthetic pathways, respectively. These proteins are proposed/proven to catalyze the attachment of TDP at C-1 of glucose utilizing **8** and TTP (**25**). These similarity results indicated that RavD catalyzes the conversion of **8** → **9**. To validate this proposed function, the enzyme was expressed in *E. coli*. SDS-PAGE analysis revealed a dominant band for His₆-RavD (Fig. S1†). Incubation of the purified enzyme with **8** and **25** resulted in the production **9** (calculated 98% product yield). The conversion was confirmed through HPLC analysis of the products with the addition of commercially available **9** and also through comparison of the ¹H NMR data of the product with data reported in the literature (Fig. S6, S11†).³⁵ Compared to TTP, there was less than 5% turnover observed when using adenosine-5'-triphosphate (ATP), uracil-5'-triphosphate (UTP), guanosine-5'-triphosphate (GTP) or cytidine-5'-triphosphate (CTP) as substrates.

The amino acid sequence of the product of *ravE* displayed strong similarities (70%–75% amino acid identities) to NDP-D-glucose-4,6-dehydratases involved in deoxysugar biosynthetic pathways from various secondary metabolites such as RmbB (accession no ACR46364) from the doxorubicin, ChryE from the chrysomycin, Gile from the gilvocarcin, MtmE from the mithramycin, StrE from the streptomycin, and SsfS2 from the SF2575 biosynthetic pathway. These similarity results indicated that RavE could serve as a TDP-D-glucose-4,6-dehydratase, which catalyzes the deoxygenation and oxidation at C-6 and C-4 positions of **9**, respectively. To test this assumption, RavE was heterologously produced in *E. coli* and purified. The size of the protein band observed on SDS-PAGE was in agreement with the calculated value for His₆-RavE. Incubation of **9** in the presence of NAD⁺ did not reveal any noticeable change in the HPLC profile when compared to the control without enzyme except for the broadening of the TDP-D-glucose peak (Fig. 3, trace b). The preparative scale

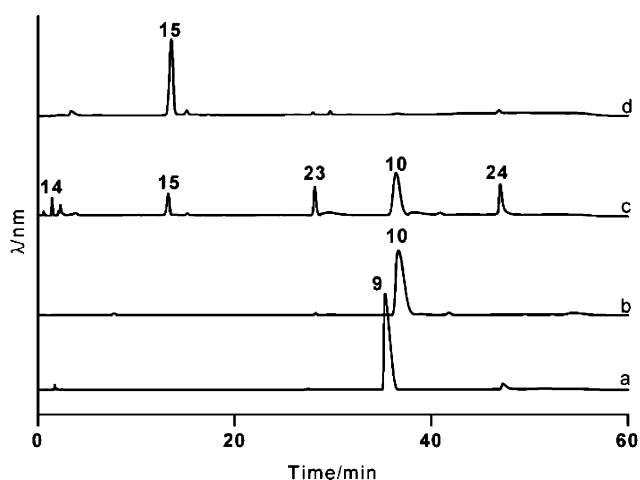


Fig. 3 HPLC traces demonstrating the conversion of TDP-D-glucose (**9**) to TDP-3-amino-3,6-dideoxy-D-galactose (**15**). Trace a: **9** generated from TTP (**25**) and glucose-1-phosphate (**8**) using RavD; trace b: reaction mixture following the incubation of **9** with RavE; trace c: reaction mixture including **10**, PLP, L-glutamate, RavIM and RavAMT; trace d: TDP-3-amino-3,6-dideoxy-D-galactose (**15**) purified through gel filtration. The compounds for **14**, **23** and **24** represents yet uncharacterized compound (preferably (2*R*,3*S*)-2-methyl-3,5-dihydroxy-4-keto-2,3-dihydropyran), TMP and TDP, respectively.

reaction was carried out by incubating **8**, **25** and NAD^+ with RavD and RavE. Separation of the crude products through gel filtration led to the isolation of **10** with 90% product yield. The structure of **10** was confirmed through comparison of its ^1H NMR data with those reported in the literature (see ESI for the ^1H NMR spectrum†). These results demonstrated that RavD and RavE catalyze the conversion of **8**→**9** and **9**→**10**, respectively, during TDP-D-ravidosamine (**17**) biosynthesis.

Functional assignment of RavIM and RavAMT as isomerase and aminotransferase

BLAST analysis of the amino acid sequence of RavIM revealed a modest sequence similarity (40%–50% amino acid identities) with putative enzymes belonging to cupin2-superfamily. These include hypothetical proteins from *Beggiatoa* sp. PS (accession ZP_01998770), *Limnobacter* sp. MED105 (ZP_01913949) and C-terminal domains of WxcM domain-containing proteins from various organisms. WxcM from *Xanthomonas campestris* was proposed to catalyze two steps, i) isomerization of **10**→**12** by its C-terminal region and ii) *N*-acetylation of **15** by the N-terminal region, during the biosynthesis of the TDP-*N*-acetyl-3-amino-3,6-dideoxy-D-galactose (**18**) subunit of surface lipopolysaccharide (LPS)³⁶ RavIM also showed considerable amino acid sequence identities (42% identity/62% similarity) to the recently characterized 3,4-ketoisomerase FdtA from *Aneurinibacillus thermoaerophilus*.³⁷ FdtA catalyzes the conversion of **10**→**12** during the biosynthesis of the *S*-layer glycan chain. Closer analysis of the RavIM amino acid sequence revealed the presence of the characteristic signature [RGXHAH(K/R)X(L/I)XQXGS] identified in FdtA (Fig. S3†). Analyses of active-site residues in the FdtA crystal structure and subsequent site-directed mutagenesis studies identified several active-site residues in this signature region.³⁸

These results support the aforementioned FdtA-like role of RavIM in the biosynthesis of **17**.

To assign the exact functional role of RavIM, the enzyme was heterologously produced in *E. coli*, and the protein was purified using immobilized metal affinity chromatography (IMAC) (Fig. S1†). Size exclusion chromatography analyses of RavIM with known proteins suggested that the enzyme exists in dimeric form. The enzyme was then subjected to a series of enzyme assays. No significant change was observed in the HPLC chromatogram of the assay mixture, when RavIM was incubated with **10** for 30 min. However, the incubation of **10** with RavIM for 24 h resulted in the consumption of **10** and the production of a new unknown product observable in the HPLC at 2.9 min (presumably (2*R*,3*S*)-2-methyl-3,5-dihydroxy-4-keto-2,3-dihydropyran **14**), TDP (at 51 min) and a minor product (at 43 min). The UV spectrum of the minor compound observed at 43 min indicated a thymine chromophore. Thus, the unknown compound peak at 43 min could be the product (**12**) of RavIM. Despite several attempts we were unable to isolate and characterize **12** due to its poor yield (< 5%) and instability. However, these results indicate RavIM might be catalyzing the isomerization reaction (**10**→**12**), and that product **12** might be spontaneously decomposing to TDP (**24**) and dihydromaltol derivative **14** in the absence of the follow-up enzyme, aminotransferase RavAMT.

BLAST analyses revealed that the product of *ravAMT* shows strong sequence similarity to aminotransferases involved in the biosynthesis of D-desosamine such as EryC1³⁹ (60%/75%), OleN2⁴⁰ (61%/73%) and DesV⁴¹ (59%/71%) from the erythromycin A, oleandomycin and pikromycin biosynthetic pathways, respectively. All of these homologous enzymes were pyridoxal-5-phosphate (PLP)-dependent and were proposed/characterized to catalyze amino transfer from an amino acid, preferably L-glutamate, to the carbonyl carbon of a deoxysugar intermediate.^{41,42} Therefore, we assumed that RavAMT might serve as an aminotransferase which converts the product of RavIM (**12**) to TDP-3-amino-3,6-dideoxy-D-galactose (**15**).

To validate the function of RavAMT, the enzyme was over-expressed in *E. coli* with a His₆-tag and was purified using IMAC. Analysis of the structure by size exclusion chromatography indicated that the enzyme exists as a dimer. Incubation of **10** with RavIM and RavAMT in the presence of sodium glutamate and pyridoxal-5-phosphate (PLP) resulted in the formation of a new compound showing a peak (calculated 20% yield) at 12.8 min in the HPLC analysis. Production of TDP (**24**) in minor amounts was also noticed (Fig. 3, trace c and d). The new peak was collected from the HPLC and desalted through gel filtration. Low resolution ESI mass analyses revealed a molecular ion peak at 546 Daltons, which matched with the molecular weight of **15** (molecular formula $\text{C}_{16}\text{H}_{27}\text{N}_3\text{O}_{14}\text{P}_2$, calcd. $[\text{M} - \text{H}]^- = 546.0968$, Fig. S12†).

The negative-mode high-resolution ESI-MS revealed a molecular ion peak at 546.0922 which further confirmed the calculated molecular formula of **15**. The ^1H NMR of this compound revealed C-5 methyl protons as a doublet at δ 1.16. The proton at C-5 position appeared as a quartet at δ 4.25. The small coupling constant ($J = 3.0$ Hz) observed for the C-4 proton (δ 3.93) indicated its equatorial orientation. The proton at C-3 position appeared as a double doublet at δ 3.63 with the coupling constants $J = 11.0$ Hz and 3.0 Hz. This further confirmed the presence

Table 1 ^1H NMR (500 MHz) data of TDP-3-amino-3,6-dideoxy-D-galactose (**15**) and TDP-D-ravidosamine (**17**) in D_2O

Position	15		17	
	δ_{H} [ppm]	Multiplicity (J/Hz)	δ_{H} [ppm]	Multiplicity (J/Hz)
1	5.54	dd (6.9, 3.5)	5.64	dd (6.9, 3.6)
2	3.88	td (11.1, 3.5)	4.21	m
3	3.63	dd (11.1, 3.1)	3.60	dd (11.4, 2.7)
4	3.93	br, d (3.1)	4.21	m
5	4.25	q (6.4)	4.25	q (6.6)
5-CH ₃	1.16	d (6.4)	1.19	d (6.6)
1'	6.28	t (6.8)	6.33	t (6.8)
2'	2.36–2.28	m	2.29–2.38	m
3'	4.55	br, m	4.60	br, m
4'	4.10–4.16	m	4.10–4.16	m
5'	4.10–4.16	m	4.10–4.16	m
5''-CH ₃	1.87	s	1.91	d (1.1)
6''	7.66	br, s	7.70	d (1.1)
3-N(CH ₃) ₂	—	—	2.94	s

of an equatorial proton at C-4 and of an axial hydrogen atom at C-2. As observed for **10**, the proton at C-2 and C-1 of **15** appeared as a double triplet at δ 3.88 and a double doublet at δ 5.54, respectively. These assignments were made by analyzing the coupling constants of the ^1H NMR in addition to ^1H - ^1H COSY correlations (Table 1). As a positive control, FdtA and FdtB were also expressed in *E. coli* and the purified enzymes were used to replace RavIM and RavAMT, respectively. Incubation of **10** with FdtA resulted in the gradual consumption of substrate with the concomitant production of a new compound, likely **12** (eluted at 43 min) (see Figs. S2 and S4†). This peak continuously grew for 60 min and then remained unchanged from 60 through 120 min. However, a continuous production of TDP (**24**) and a new peak (1.8 min) was observed throughout the assay period indicating the decomposition of **12** (Fig. S5†). The specific activity of FdtA ($201 \mu\text{mol mg}^{-1} \text{min}^{-1}$) was found to be ~6-fold higher than that of RavIM ($31.3 \mu\text{mol mg}^{-1} \text{min}^{-1}$). Co-incubation of FdtA and FdtB with TDP-4-keto-6-deoxy-D-glucose, pyridoxal-5-phosphate and L-glutamate resulted in the production of **15** as observed earlier through the co-assay of RavIM and RavAMT. These results also clearly established that RavIM and RavAMT act as 3,4-keto-isomerase and aminotransferase, respectively, which catalyze the conversion of **10**→**12** and **12**→**15**, respectively.

Heterologous production and characterization of RavNMT

The encoded amino acid sequence of *ravNMT* showed strong similarity to various *S*-adenosylmethionine (SAM)-dependent methyltransferases (AdoMet-MTase). These homologous enzymes include *N*-methyltransferases such as SnogX (56%/71%) and SnogA (57%/70%) from the TDP-L-nogalamine biosynthetic pathway.⁴³ Similarly, *N,N*-dimethyltransferases such as DesVI (55%/71%) and OleM1 (53%/68%) from the TDP-D-desosamine pathway^{42,44} represent the other homologous enzymes in the database. Both enzymes are involved in the biosynthesis of TDP-D-desosamine, and transfer two methyl groups in an iterative fashion from SAM to the amino nitrogen. Since the nitrogen at C-3' position of ravidosamine contains two methyl groups, and there was no other deoxysugar-*N*-methyltransferase encoding candidate gene identified in the ravidomycin V gene cluster, we

reasoned that RavNMT serves as a dimethyltransferase catalyzing the conversion of **15**→**17**.

To validate the proposed function of RavNMT, the enzyme was heterologously produced in *E. coli*, and purified through IMAC (Fig. S1†). Size exclusion chromatography analysis of RavNMT along with standard proteins suggested that RavNMT exists as a dimer. Incubation of **15** with RavNMT in the presence of SAM resulted in the complete consumption of **15** with concomitant production of a new peak (calculated ~90% product yield) at 4.8 min in the HPLC (Fig. 4). The compound pertaining to the new peak was isolated and purified through HPLC and gel-filtration. High resolution ESI mass of the isolated compound revealed a *quasi* molecular ion peak of 574.1227 Dalton, which is in agreement with the molecular formula $\text{C}_{18}\text{H}_{31}\text{N}_3\text{O}_{14}\text{P}_2$ of the proposed product **17** (calcd. MW, $[\text{M} - \text{H}]^- = 574.1281$). Moreover, the ^1H NMR analysis of the compound revealed a new singlet peak at δ 2.94 indicating the presence of two *N*-methyl groups. To prove that RavNMT catalyzes the transfer of two methyl groups sequentially, the reaction was stopped after 10 min and the products subjected to MS analyses. The negative-mode high-resolution ESI mass spectra of the reaction mixture revealed molecular ion peaks of 574.1227, 560.1073 and 546.0922 Daltons, which corresponded to the molecular ions of **17**, **16** and **15**, respectively. These results showed that RavNMT catalyzes the transfer of methyl groups from SAM to the amino nitrogen of **15** in a sequential manner.

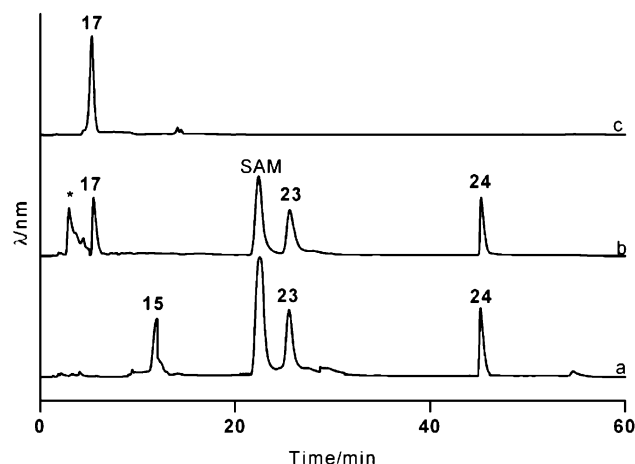


Fig. 4 HPLC traces demonstrating the conversion of TDP-3-amino-3,6-dideoxy-D-galactose (**15**) to TDP-D-ravidosamine (**17**). Trace a: control assay mixture including **15** and *S*-adenosylmethionine (SAM); trace b: products formed after addition of RavNMT with the mixture shown in trace a; trace c: TDP-D-ravidosamine (**17**) purified through gel filtration. Peaks **23** and **24** represent TMP and TDP produced through the spontaneous decomposition of **15**, respectively. The peak marked with an asterisk was not characterized, as its UV spectrum did not indicate a thymidine chromophore.

One-pot enzymatic synthesis of TDP-D-ravidosamine

One of the major goals of this study was to generate the anticipated sugar donor substrate **17** of RavGT, necessary for future *in vitro* glycosylation assays. Such assays require an easy, reproducible and an efficient way to generate **17**. Having all of the biosynthetic enzymes in hand, we attempted to set up a protocol

to generate **17** in a large-scale one-pot setup. However, the fairly expensive starting material TTP (**25**) would limit a larger-scale production of **17**. Therefore, we cloned the enzymes necessary for the biosynthesis of TTP from *E. coli* following a reported protocol.⁴⁵ Thymidine monophosphate kinase (TMK) generates **24** from **23**, and acetate kinase (ACK) further phosphorylates **24** to TTP (**25**). ATP serves as a phosphate donor thereby converting into ADP, acetate kinase also regenerates ATP by transferring the phosphate group of acetyl phosphate to ADP (Scheme 2).

Both proteins were overexpressed in *E. coli* and purified through IMAC (Fig. S1†). Incubation of **23** with acetyl phosphate and catalytic amount of ATP resulted in the production of **24** and **25** in quantitative yields. Since these reactions were reversible, the reaction was driven forward by adding TDP-D-glucose synthase and **8** to the assay mixture (Fig. 5). TDP-D-glucose synthase from *Salmonella typhimurium* (RfbA) was heterologously produced in *E. coli* and was used in place of relatively unstable RavD. Considering the fact that **24** can serve as an inhibitor for the downstream pathway enzymes, overall TDP-ravidosamine (**17**) biosynthesis was divided into two stages in the same pot. In the first stage, **23**, **8** and ATP were incubated with ACK, TMK, RfbA and RavE/RfbB/RmlB that resulted in the production of **10** efficiently (~95% yield) in 5 min (for

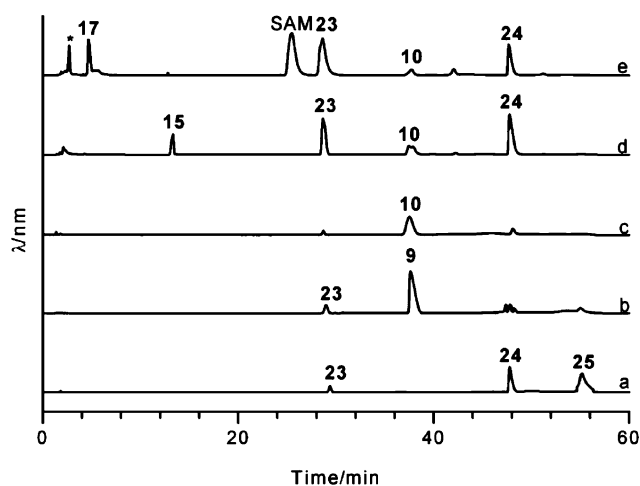


Fig. 5 Enzymatic total synthesis of TDP-D-ravidosamine. Trace a: Thymidine monophosphate (TMP, **23**) incubated with acetate kinase (ACK), thymidine monophosphate kinase (TMK) and adenosine triphosphate (ATP), **24** = TDP, **25** = TTP; trace b: assay mixture of trace a incubated with glucose-1-phosphate (G-1-P) and RavD, **9** = TDP-D-glucose; trace c: assay mixture of trace b supplemented with RavE, **10** = TDP-4-keto-6-deoxy-D-glucose; trace d: assay mixture of trace c supplemented with RavIM, RavAMT, L-glutamate and pyridoxal-phosphate (PLP), **15** = TDP-3-amino-3,6-dideoxy-D-galactose; trace e: assay mixture of trace d supplemented with S-adenosyl methionine (SAM) and RavNMT, **17** = TDP-D-ravidosamine. Reactions were carried out for 1 h prior to the HPLC analyses. The compounds on the HPLC traces for a, b, c and d remained unchanged when the commercial TTP was used in place of the TTP generated *in situ* by ACK, TMK, acetyl phosphate, TMP (**23**) and ATP. The peaks with numbers **9**, **10**, **15**, **17**, **23**, **24** and **25** represent TDP-D-glucose, TDP-4-keto-6-deoxy-D-glucose, TDP-3-amino-3,6-dideoxy-D-galactose, TDP-D-ravidosamine; TMP, TDP and TTP, respectively. The peak marked with an asterisk was not characterized, as its UV spectrum did not indicate a thymidine chromophore. SAM represents S-adenosyl methionine.

details, see Experimental section). Because of its instability RavE was substituted by the TDP-D-glucose-4,6-dehydratase (RfbB) from *Salmonella typhimurium*. RfbB has been routinely used for an efficient conversion of **9**→**10**.^{8,35,46} However, following several batches of protein production, the recombinant RfbB was expressed mostly as insoluble protein. Therefore, an additional TDP-D-glucose-4,6-dehydratase (encoded by the gene *rmlB*) was cloned from *E. coli*, and the protein was expressed and purified. MS analysis of the products revealed that addition of RmlB in the assay mixture resulted in the complete conversion of **9**→**10** within five minutes. Following the complete conversion of **8**→**10**, the enzymes were removed from the assay mixture and the second stage enzymes (RavIM, RavAMT, and RavNMT) along with their co-factors (PLP, L-glutamate, and S-adenosyl-methionine (SAM)) were then added to the assay mixture to complete the synthesis of **17**. More than 95% of **10** was consumed in 65 min (Fig. 6) with the concomitant productions of **15** and a mixture of **16** and **17**. The presence of **16** and **17** in the assay mixture was confirmed through HR-MS analysis. However, we were unable to determine their exact quantities, as both compounds were eluted at identical retention times in the HPLC. The production of these compounds remained unchanged from 60 to 180 min. This was a clear indication of the lack of SAM in the assay mixture. Addition of excess SAM resulted in a dramatic increase in production of **17** with the consumption of **15** as well as of **16** in additional 60 min, as evident from ¹H NMR analyses of the products. The overall yield of **17** generated was estimated to be 85%.

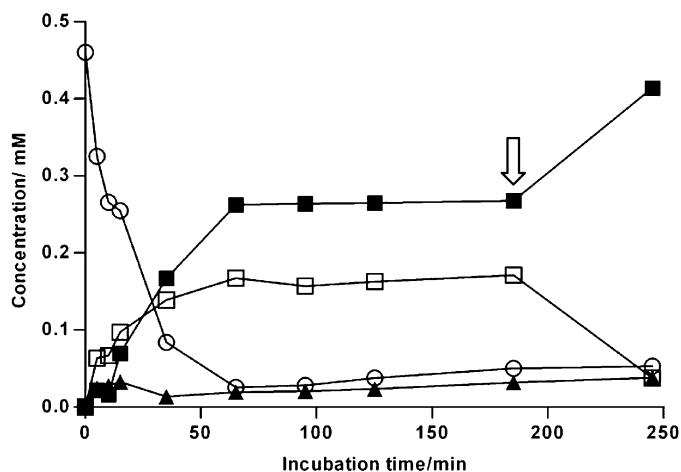
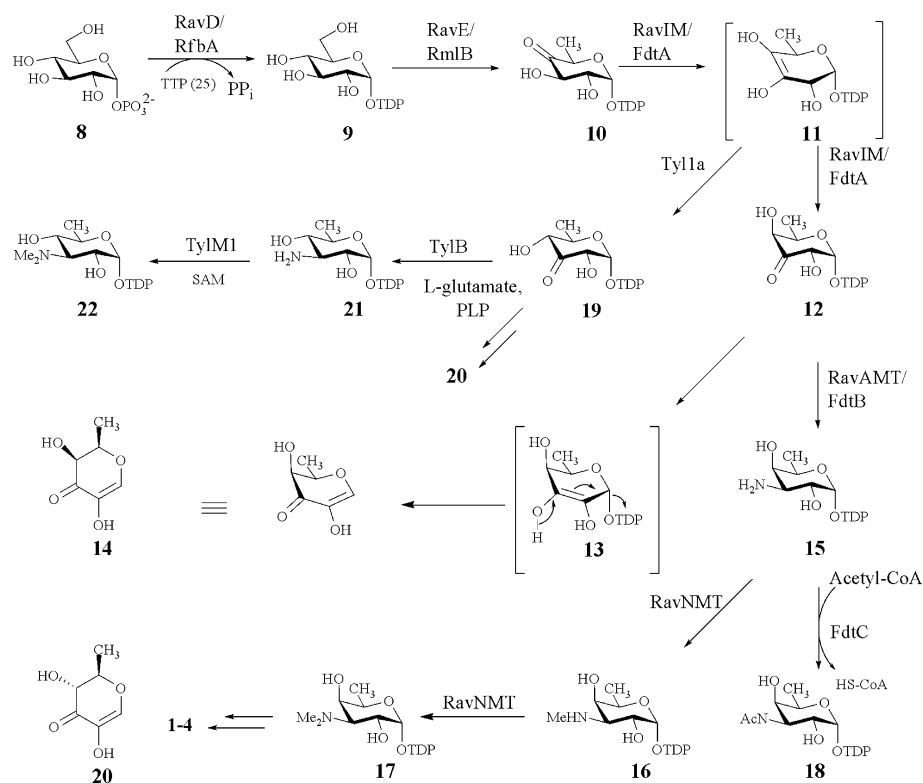


Fig. 6 Conversion profile of TDP-4-keto-6-deoxy-D-glucose (**10**) to TDP-D-ravidosamine (**17**) at the second stage. Lines with empty circles, solid triangles and empty squares represent the concentration profile of TDP-4-keto-6-deoxy-D-glucose (**10**), TDP-3-keto-6-deoxy-D-galactose (**12**), and TDP-3-amino-3,6-dideoxy-D-galactose (**15**), respectively. The line with filled squares from 0 through 180 min represents concentrations of the mixture of TDP-D-ravidosamine (**17**) and TDP-3-N-methyl-3-amino-3,6-dideoxy-D-galactose (**16**). Further addition of SAM at 180 min (arrow) resulted in the complete conversion of **15** and **16**→**17** in 240 min.

Discussion

Bioactive glycosylated natural products containing amino sugars have the advantage that they can be solubilized through



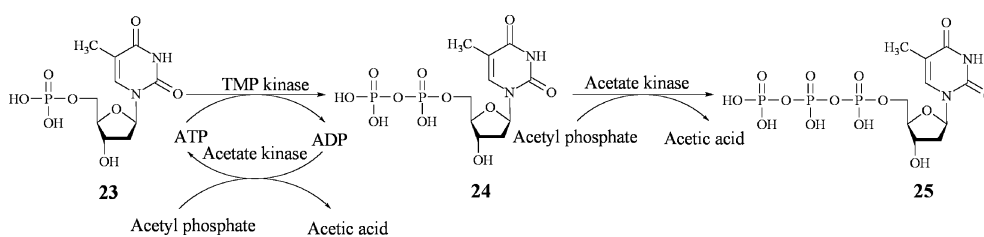
Scheme 1 Biosynthetic pathways to TDP-D-ravidosamine (17), TDP-D-mycaminose (22), and TDP-3-N-acetamido-3,6-dideoxy-D-galactose (18).

formulation as salts, or packaged into nano-liposome delivery vehicles.^{47–49} Salt forms of such products often exhibit better bioavailability in addition to their increased hydrophilicity. Therefore, the study of dedicated glycosyltransferases which utilize amino sugars is promising for the generation of new pharmaceutically useful natural products through glycodiversification. C-GTs which establish stable C–C bonds between sugar and aglycone are considered special, since C-glycosylated compounds cannot be destroyed by hydrolysis through glycosidases. Although dozens of C-GTs have been identified from secondary metabolite biosynthetic pathways, none of those that utilize deoxyaminosugars has been characterized *in vitro*. Deacetylavidomycin V (2) is of particular interest as it contains a C-glycosidically linked aminosugar and is one of the most potent drugs among the gilvocarcin family of anticancer compounds. It was our hope that by producing the anticipated sugar donor substrate for RavGT, presumably TDP-D-ravidosamine (17), we would be able to further investigate the glycosylation activity of RavGT *in vitro*.

We had previously demonstrated that expression of the cosmid cosRav32 in *S. lividans* TK64 accumulates 2 and deacetylavidomycin M (4). This indicated that the hypothesized enzymes RavD, RavE, RavIM, RavAMT and RavNMT are sufficient to biosynthesize 17. Biosynthesis of all deoxysugars begins with the activation of hexose-1-phosphate utilizing a particular NTP. Although TTP is most commonly used by NDP-hexose synthases, there are also examples where GTP or CTP are utilized for activation.^{50–53} Therefore, various NTPs were tested for the activation of 8. *In vitro* assays clearly demonstrated that RavD preferentially utilizes TTP (25) to generate 9 which converts into 10 through the activity of RavE. These two steps are

common to a variety of deoxysugar biosynthetic pathways, such as TDP-D-olivose,⁵⁴ TDP-L-olivose,⁵⁵ TDP-D-mycarose,¹⁴ TDP-L-mycarose,⁴⁶ TDP-D-fucose⁵⁶ and TDP-L-rhamnose.⁵⁷ Isomerization of 10→12, catalyzed by RavIM, was the unique step of the TDP-ravidosamine (17) biosynthetic pathway. Such a conversion was further confirmed by the formation of an identical product in a parallel assay involving RavIM's homologue FdtA. To the best of our knowledge, no such isomerase has been previously reported from secondary metabolite biosynthetic pathways. FdtA from *Aneurinibacillus thermoaerophilus* L420-91 was the only enzyme characterized to catalyze such a transformation during the biosynthesis of 3-acetamido-3,6-dideoxy-D-galactose (18) during cell wall biosynthesis.^{37,38} The TDP-D-mycaminose (22) biosynthetic pathway is identical to the ravidosamine (17) pathway except for the isomerization step where Tyll1a converts 10→19 with an equatorial hydroxyl group at the C-4 position (Scheme 1).³⁵ Incorporation of ravidosamine in place of the mycaminose moiety of a pikromycin derivative through the expression of FdtA in the Tyll1a-deletion mutant has demonstrated a close resemblance of these pathways.⁵⁸ Comparative studies of RavIM and FdtA revealed that the former enzyme is less active compared to the latter (specific activity of RavIM appeared to be ~6.4-fold less than that of FdtA). Such a small turnover of the recombinant RavIM hindered us from studying the isomerization step in detail. Therefore, FdtA was used to monitor the 3,4-ketoisomerization reaction.

Isolation and characterization of 12 would provide direct evidence for the activity of FdtA and RavIM. However, the newly formed compound was spontaneously decomposed to TDP (24) and presumably (2R,3S)-2-methyl-3,5-dihydroxy-4-keto-2,3-dihydropyran (14); however, the structure of the latter is yet to be



Scheme 2 Enzymatic synthesis of thymidine-triphosphate (TTP).

confirmed. In a time course study, a continuous production of **24** with the consumption of **10** was observed, but the concentration of a new peak pertaining to **12** remained unchanged after 55 min. This suggested an existence of two parallel reactions: (i) the enzyme-catalyzed isomerization $10 \rightarrow 12$ and (ii) the spontaneous decomposition of $12 \rightarrow 24$ and **14**. The product **12** could easily undergo keto–enol tautomerism leading to the spontaneous decomposition as shown in Scheme 1. There is precedence for such a transformation, observed in the D-mycaminose pathway where the Tyl1a product **19** spontaneously degraded to TDP and (2*R*,3*R*)-2-methyl-3,5-dihydroxy-4-keto-2,3-dihydropyran (**20**).³⁵

Comparable conversions of $10 \rightarrow 15$ using RavIM–RavAMT or FdtA–FdtB were in agreement with previous reports using Tyl1a–TylB, where the isomerization reaction is reversible and transamination drives the equilibrium forward.³⁵ The conversion of **15** to **17** through the activity of RavNMT unambiguously proved the enzyme to be an *N,N*-dimethyltransferase. Identification of both **17** and **16** in the assay mixture suggested a sequential transfer of methyl groups from SAM to the amino nitrogen. DesVI and TylM1 from desosamine and mycaminose biosynthetic pathways, respectively, are other examples of such *N,N*-dimethyltransferases which catalyze dimethylation of a sugar amino group in a sequential manner.^{44,59}

Along with a high demand of NDP-activated sugars for *in vitro* glycosylation studies, reports on one-pot enzymatic syntheses of NDP-sugars are rapidly growing.^{46,57,60} With all of the functional biosynthetic enzymes in hand, we also devised a one-pot enzymatic synthesis protocol for the quick and efficient production of TDP-D-ravidosamine (**17**) in a cost-efficient manner. Instead of using expensive commercial TTP (**25**), the compound was generated *in situ* from TMP and acetyl phosphate using acetate kinase (ACK) and thymidine monophosphate kinase (TMK) from *E. coli*. An identical strategy was applied earlier for the enzymatic synthesis of TDP-L-rhamnose,⁵⁷ TDP-4-keto-6-deoxy-D-glucose (**10**),⁴⁵ and TDP-4-amino-4,6-dideoxy-D-glucose.⁶⁰ In the first stage, intermediate **10** was produced in satisfactory yields (~95%). Further incubation of the first-stage product with the remaining biosynthetic enzymes resulted in the efficient production of TDP-D-ravidosamine (**17**), with ~85% yield for the second-stage process. Suzuki *et al.* had developed an elegant chemical synthesis of ravidosamine and some of its derivatives.³⁰ However, like the syntheses of other deoxysugars, particularly when TDP-activated, the reported procedure involved lengthy synthetic steps with overall poor yields. In addition, the published ravidosamine synthesis lacks the TDP-activation, and is therefore unsuited to provide **17** for glycosyltransferase studies. The enzymatic total

synthesis described here is also likely the better method to mass-produce ravidosamine derivatives in general.

In summary, all of the proposed ravidosamine biosynthetic enzymes were characterized *in vitro*. The results unambiguously established a pathway for the biosynthesis of the sugar moiety of deacetylavidomycin V (**2**) and deacetylavidomycin M (**4**). The one-pot enzymatic synthesis protocol developed in this study is a valuable tool for producing TDP-D-ravidosamine (**17**) in large scale for *in vitro* glycosylation studies involving RavGT as well as other suitable GTs.

Experimental

Cloning and preparation of expression constructs

Oligonucleotide primers used in this study are summarized in Table S1†. Strain *E. coli* XL1 Blue MRF was used for the routine amplification of plasmid DNA. Genomic DNA of *Salmonella enterica* serovar. typhimurium LT2, *E. coli* XL1 Blue and *Aneurini-bacillus thermoaerophilus* L420-91 were prepared following a standard protocol.⁶¹ Nucleotide sequences for the genes encoding TDP-D-glucose-4,6-dehydratase (*rmlB*), acetate kinase (*ack*) and thymidine monophosphate kinase (*tmk*) were amplified from the genomic DNA of *E. coli*. Similarly, the genes encoding TDP-D-glucose thymidyltransferase (*rfbA*), TDP-D-glucose-4,6-dehydratase (*rfbB*), and TDP-4-keto-6-deoxy-D-glucose-3,4-keto-isomerase (*fdtA*) and aminotransferase *fdtB* were amplified from the genomic DNA of *Salmonella typhimurium* LT2 and *Aneurini-bacillus thermoaerophilus* L420-91, respectively. All of the TDP-D-ravidosamine biosynthetic genes were amplified from cosmid cosRav32²⁹

Expression and purification of proteins

Pfu polymerase (Stratagene) was used for PCR amplifications, and the obtained products were cloned into the ZeroBlunt-TOPO vector (Invitrogen). Sequencing of the TOPO clones were carried out to ascertain the fidelity of the PCR amplifications. Nucleotide fragments covering *ack*, *ravD*, *ravE*, *ravIM*, *ravAMT*, *ravNMT* and *rfbA* genes were isolated from their respective TOPO-clones using *Nde*I and *Eco*RI restriction enzymes and subsequently ligated to identical sites of the pET28a(+) expression vector (Novagen). Similarly, gene sequences for *tmk*, *rfbB* and *rmlB* were ligated at *Eco*RI/*Hind*III, *Bam*HI/*Hind*III and *Bam*HI/*Xho*I sites of pET28a(+), respectively. The generated constructs were then transformed into *E. coli* BL21(DE3). A single colony was inoculated into 20 mL of LB liquid supplemented with kanamycin

sulfate and was grown for 4 h to prepare a seed culture. One liter of LB (100 mL $\times$ 10 flasks) was inoculated with 10 mL of the seed culture and grown at 37 °C until OD₆₀₀ reached ~0.5. The culture was then grown at 18 °C for 12 h following addition of IPTG (0.5 mmol final concentration). The cell pellets were collected by centrifugation (4000 $\times$ g, 15 min) and were washed twice with 20 mL of lysis buffer (50 mmol KH₂PO₄, 300 mmol NaCl, 10 mmol imidazole, pH 7.6). Disruption of the pellets was carried out using a French Press (Thermo electron corporation), and the crude soluble enzyme fraction was collected through centrifugation (17 000 $\times$ g). N-terminal His-tagged enzymes were purified through immobilized metal affinity chromatography (IMAC). The purified proteins were further desalted using a Profinia protein purification system (Bio-Rad). The Bradford protein assay method was used to determine the concentration of the enzymes.⁶² The purities of the enzymes were evaluated using dodecylsulfate–polyacrylamide gel electrophoresis (SDS-PAGE). The yields of purified proteins (mg per L culture) which slightly varied from batch to batch are as follows: ACK, 8–10; TMK, 18–20; RfbA, 18–20; RmlB, 21–24; FdtA, 6–8; FdtB, 10–13; RavD, 18–20; RavE, 8–10; RavNMT, 12–15; RavAMT, 12–15 and RavIM, 4–6.

Substrates and co-factors

Thymidine-5'-phosphate (TMP, **23**), thymidine-5'-diphosphate (TDP, **24**), thymidine-5'-triphosphate (TTP, **25**), uridine-5'-triphosphate (UTP), guanosine-5'-triphosphate (GTP), cytidine-5'-triphosphate (CTP), adenosine-5'-triphosphate (ATP), glucose-1-phosphate (**8**), TDP-D-glucose (**9**), acetyl phosphate, pyridoxal-5'-phosphate (PLP), L-glutamic acid and S-adenosyl methionine (SAM) were purchased from Sigma–Aldrich.

Enzyme assay of RavD and preparation of TDP-D-glucose

Enzyme assays for TDP-D-glucose synthases (RavD or RfbA) were carried out in 200 μ L mixture composed of 50 mmol phosphate buffer (pH 7.5), 20 mmol MgCl₂, 5 mmol glucose-1-P (**8**), 5 mmol of each NTP (ATP, UTP, TTP, GTP and CTP, respectively) and 3 μ mol enzyme. The mixture was incubated for 30 min in a 37 °C water bath. The reaction was stopped by heating the mixture at 90 °C for 3 min. The precipitated enzymes were removed by centrifugation and the clear supernatant fractions were passed through an amicon filter (MW cut-off 3000 Da). HPLC analyses of the products were carried out at 25 °C using Waters HPLC system equipped with photodiode array detector. Compounds were monitored at 267 nm wavelength. The filtrate (50 μ L) was injected in the HPLC. CarboPackTM PA1 (4 $\times$ 250 mm) column (Dionex) was used to separate compounds following the previously reported procedure.³⁵ A gradient of ammonium acetate (500 mmol, solvent B) and water (solvent A) was used to separate the assay products: 5–20% solvent B from 0–15 min, 20–60% solvent B from 15–35 min, 60–100% solvent B from 35–37 min, 100% solvent B from 37–40 min, 100–5% solvent B from 40–45 min and 5% solvent B from 45–60 min. A control sample without enzyme and standard TDP-D-glucose (**9**) was also injected for the identification of the product peak. For large-scale production of **9**, TMP (5 mmol), **8** (5.1 mmol), acetyl phosphate (20 mmol) and ATP (5 μ mol) were mixed in a 20 mL solution composed of 50 mmol phosphate buffer (pH 7.5), 20 mmol MgCl₂, 4 μ mol ACK, 6 μ mol TMK

and 5 μ mol RavD/RfbA. The mixture was incubated for 1 h, and the reaction was quenched by heating the mixture at 90 °C for 5 min. The mixture was centrifuged at 10 000 $\times$ g for 10 min and the supernatant was lyophilized. The dry powder was dissolved in 5 mL of water and the solution was loaded into the pre-equilibrated Bio-Gel P2 gel filtration column 100 cm $\times$ 25 mm. Separation was carried out at 4 °C with water at flow rate of 0.2 mL min⁻¹. Aliquots of 4 mL were collected and the purity of compounds was monitored by HPLC at 267 nm. Pure fractions were combined, lyophilized and analyzed by NMR. About 80 mg of pure **9** was obtained along with an additional 125 mg of **9** with minor TMP (**23**) and TDP (**24**) impurities. The purity **9** was assayed through HPLC and ¹H NMR (Fig. S6†). ¹H NMR of **9** (500 MHz, D₂O): δ 1.90 (3H, s, 5''-Me), 2.31–2.38 (2H, m, 2'-H), 3.43 (1H, t, *J* 9.8 Hz, 3-H), 3.51 (1H, td, *J* 9.8, 3.5 Hz, 2-H), 3.72–3.78 (2H, m, 4-H, 5-H), 3.81–3.90 (2H, m, 6-H₂), 4.12–4.16 (1H, m, 4'-H), 4.12–4.16 (2H, m, 5'-H), 4.62 (1H, m, 3'-H), 5.58 (1H, dd, *J* 7.4, 3.5 Hz, 1-H), 6.33 (1H, t, *J* 6.98 Hz, 1'-H), 7.72 (1H, s, br, 6''-H).

Enzyme assay of RavE and preparation of TDP-4-keto-6-deoxy-D-glucose

To carry out TDP-D-glucose-4,6-dehydratase assays, the enzymes (4 μ mol RavE or 3 μ mol RfbB or 3 μ mol RmlB) were added to 200 μ L solution composed of phosphate buffer (50 mmol, pH. 7.5), MgCl₂ (20 mmol), NAD⁺ (2 μ mol) and **9** (5 mmol). The mixture was incubated for 30 min and the reaction was stopped. An enzyme-free solution was analyzed by HPLC. For large-scale production, a one-pot enzymatic synthesis was carried out in 50 mL volume composed of phosphate buffer (50 mmol, pH 7.5), MgCl₂ (20 mmol), **8** (150 mmol), TMP (**23**, 98.3 mmol), acetyl phosphate (400 mmol), ATP (1 mmol), ACK (0.8 μ mol), TMK (2.1 μ mol) and RfbA (2.0 μ mol). The mixture was incubated for 1 h at 37 °C, and then RavE (final concentration of 2.73 μ mol) or RfbB (2.3 μ mol) and NAD⁺ (2 μ mol) were added and incubation was continued for an additional 2 h. Surprisingly, when RavE and RfbB was expressed for additional reactions they appeared exclusively in the form of inclusion bodies. Therefore, these enzymes were replaced by RmlB (6 μ mol) in the following batches. The reaction mixture was heated to denature the enzymes. The precipitated fractions were removed by centrifugation. The crude product was lyophilized to powder and dissolved in 10 mL water. Separation of the products was carried out in Bio-gel P2 following the protocol described above. The purity of each fraction was analyzed by HPLC. Pure fractions were combined and lyophilized to powder. This afforded 700 mg of TDP-4-keto-6-deoxy-D-glucose (**10**) with ~95% purity and an additional 630 mg of **10** with >70% purity. The purity and identity of the compound was further confirmed through the comparison of ¹H NMR data reported in the literature (Fig. S7†). ¹H NMR of **10** (500 MHz, D₂O): δ 1.06 (3H, d, *J* 6.50 Hz, 5-Me), 1.77 (3H, s, *J* 0.97 Hz, 5''-Me), 2.23 (2H, dd, *J* 6.90, 4.85 Hz, 2'-H₂), 3.51 (1H, td, *J* 10.07, 3.3 Hz, 2-H), 3.63 (1H, d, *J* 10.07 Hz, 3-H), 3.93 (1H, q, *J* 6.5, 5-H), 3.99–4.06 (3H, m, 4'-H, 5'-H), 4.43–4.47 (1H, m, 3'-H), 5.40 (1H, dd, *J* 6.8, 3.3 Hz, 1-H), 6.19 (1H, t, *J* 6.90 Hz, 1'-H), 7.57 (1H, d, 6''-H).

Assay of RavIM alone or coupled with RavAMT

RavIM (20 μ mol final concentration) was incubated at 37 °C with **10** (1 mmol) in a 500 μ L solution composed of phosphate

buffer (50 mmol, pH 7.5) and MgCl_2 (20 mmol) for 60 min. An aliquot (60 μL) was drawn from the incubation mixture and flash-frozen in dry ice–acetone. The enzyme was removed by further heating the mixture at 90 °C for 3 min. The solution was filtered through a centricon (MW cut-off 3000 Da) and the filtrate (50 μL) was analyzed by HPLC. In the coupled enzyme assays, RavAMT (30 μmol), L-glutamate (10 mmol) and PLP (60 μmol) were added in the RavIM assay mixture and progress of the reaction was monitored until 60 min. A new peak appeared in the coupled assays at 12.8 min which was collected and analyzed by high-resolution ESI-MS. Similarly, FdtA and FdtA–FdtB coupled assays were also carried out in parallel as positive controls. For the preparative scale production of TDP-3-amino-3,6-dideoxy-D-galactose (**15**), the substrate and enzyme concentrations were doubled and the assay volume was increased to 20 mL. After 6 h of incubation, enzymes were removed and the volume was reduced to 1 mL. Separation of the compounds was carried out in analytical-scale HPLC. The compound eluted at 12.8 min as before and was collected and separated from the ammonium acetate buffer using a Bio-Gel P2 column. The pure fractions were combined and lyophilized. Overall, 12 mg powder of **15** was purified. The structure of **15** was confirmed through MS and NMR data analyses.

HPLC analyses of FdtA-catalysed isomerization reaction

TDP-4-keto-6-deoxy-D-glucose (**10**, 1 mmol) was incubated at 37 °C with 24.9 μmol enzyme in 400 μL solution containing 50 mmol phosphate buffer and 20 mmol magnesium chloride. Aliquots (60 μL) were drawn from the assay mixture at various time points and quenched in a dry ice–acetone bath. The samples were boiled for 4 min and centrifuged to remove the precipitated proteins. The clear supernatant (50 μL) was injected into the HPLC and the peak areas under the curve were normalized (Fig. S4 and S5†).

In vitro assay of RavNMT

For an analytical scale assay, a 200 μL assay mixture containing 50 mmol phosphate buffer, 20 mmol MgCl_2 , 1 mmol of **15**, 4.0 mmol SAM and 10 μmol RavNMT were incubated for two hours. An enzyme-free solution was prepared as discussed earlier and injected in an HPLC. A new peak appeared at 4.8 min and was collected and analyzed by HR-ESI mass spectrometry. For large-scale production, a coupled assay was carried out in 20 mL using **10** (2 mmol), L-glutamate (14 mmol), PLP (80 μmol), SAM (4.0 mmol) RavIM (4.8 μmol), RavAMT (6.5 μmol) and RavNMT (8.0 μmol). After two hours of incubation, the enzymes were removed and the solution was lyophilized. The dried powder was dissolved in 1 mL water and separated by HPLC. The compound eluted at 4.8 min and was collected and desalted using a Bio-gel-P2 column. Lyophilization of the pure fractions afforded 5 mg of TDP-D-ravidosamine (**17**) and 8 mg of a 1 : 1 mixture of TDP-3-N-methyl-3-amino-3,6-dideoxy-D-galactose (**16**) and **17**, respectively. The structure of **17** was determined through NMR and HR-MS analyses.

Two-stage one-pot enzymatic synthesis of TDP-D-ravidosamine

Overall enzymatic synthesis of **17** was divided into two stages. For the first stage, a 20 mL solution containing **8** (0.5 mmol), **23**

(0.48 mmol), ATP (45 μmol), acetyl phosphate (2 mmol), ACK (5 μmol), TMK (10 μmol), RfbA (8 μmol), RmlB (5.8 μmol), phosphate buffer (50 mmol, pH 7.5) and MgCl_2 (20 mmol) was incubated at 37 °C for 30 min. HPLC analyses of the assay mixture revealed the production of **10** (0.47 mmol) in the assay mixture. Enzymes were precipitated by heating the solution to 95 °C and removed through centrifugation. In the second stage, RavIM (2.3 μmol), AMT (3.9 μmol), RavNMT (4.5 μmol), L-glutamate (7 mmol), PLP (42 μmol) and SAM (0.7 mmol) were then added to the assay mixture and incubation was continued for 5 h. Considering the fact that SAM degrades in water, an additional 0.7 mmol (final concentration) of SAM was supplied after 3 h. The reaction was monitored using HPLC as discussed above. A calibration curve was generated varying the concentration of **23** from 62 nM to 15 mmol. The concentrations of TDP-D-hexoses were determined using this calibration curve. The reaction was stopped by heating at 90 °C for 4 min. The denatured protein was removed through centrifugation and the liquid fraction was lyophilized. The peak corresponding to **17** was separated through gel filtration. The fractions including **17** were lyophilized and then analyzed by NMR spectroscopy.

Acknowledgements

This work was supported by the US National Institutes of Health (grant CA 102102 to JR). The NMR and mass spectrometry core facilities of the University of Kentucky are acknowledged for the access to their instruments. Dr Guy Carter, Wyeth, is acknowledged for providing the ravidomycin producer *Streptomyces ravidus*.

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