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Total Synthesis of Tiacumicin B: Study of the Challenging β -Selective Glycosylations

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ABSTRACT: We report here a full account of the total synthesis of tiacumicin B (Tcn-B), a natural glycosylated macrolide with remarkable antibiotic properties. Our strategy is based on our experience with the synthesis of the tiacumicin B aglycone and on unique 1,2-*cis*-glycosylation steps. It features the conclusive use of sulfoxide anomeric leaving-groups in combination with a remote 3-*O*-picoloyl group on the donors allowing highly β -selective rhamnosylation and noviosylation that rely on H-bond-mediated Aglycone Delivery (HAD). The rhamnosylated C1-C3 fragment was anchored to the C4-C19 aglycone fragment by a Suzuki-Miyaura cross-coupling. Ring-size selective Shiina macrolactonization provided a semi-glycosylated aglycone that was engaged directly in the noviolysation step with a virtually total β -selectivity. Finally, a novel deprotection method was devised for the removal of a 2-naphthylmethylidene (Nap) ether on a phenol and efficient removal of all the protecting groups provided synthetic tiacumicin B.

Nowadays antibiotic resistance is one of the most serious threats to global health, food security and development, degrading the quality of life and heavily impacting the economy with longer hospital stays, higher medical expenses and higher mortality. This is a natural phenomenon whose process is accelerated by the excessive use or misuse of these drugs in humans and animals. An increasing number of infections, such as pneumonia, tuberculosis or salmonellosis, are becoming more and more difficult to treat because of the lack of effectiveness of the antibiotics used. To circumvent this resistance, one of the tracks consists in developing new antibiotics with new biological targets. Tiacumicin B (Tcn-B) meets these criteria and, received marketing authorization in 2011 in the United States for the treatment of *Clostridium difficile* intestinal infections, which are often of nosocomial origin and had previously been fatal in 25% of the cases.¹ Tcn-B interacts with bacterial RNA polymerase (RNAP)² blocking RNA synthesis, a strategy already used in broad-spectrum antibacterial therapy. Since Tcn-B inhibits bacterial RNAP by binding a site that does not overlap with other antibiotic binding sites, there is no known cross-resistance with the other antibiotics in use. There is no cross-resistance with rifamycin,³ so Tcn-B is active on resistant strains of *mycobacterium tuberculosis*, which opens up new therapeutic possibilities.⁴ In this context, designing reliable total syntheses of Tcn-B that could also lead to analogues proves particularly relevant.

Figure 1. Tiacumicin B (Tcn-B)

Tcn-B is representative of a new class of antibiotic macrolides and is also known as clostomicin B1, fidaxomicin or lipiarmycin A3 (Figure 1).⁵ This natural product was first isolated in the 70's from an actinobacterium, *Actinoplanes deccanensis*, found in the soil in India.⁶ Its

structure was partially elucidated between 1983 and 1987 by Martinelli,⁷ Arnone and Nasini⁸ and recently its biosynthesis could be resolved by the team of Zhang.⁹ Tcn-B is among the most complex and heavier macrolide and its structure can be divided into three main parts: 1) a central core composed of an eighteen-membered macrolactone displaying two (*E*),(*E*)-conjugated dienes, a trisubstituted (*E*)-alkene and featuring five stereogenic centers; 2) an eastern part with 2-*O*-methyl-D-rhamnose, linked as the β -anomer and esterified at the 4-position by an homodichloro-orsellinic acid; and 3) a western part consisting in a rare sugar, D-noviose, also linked as the β -anomer and esterified at the 4-position by isobutyric acid.

In 2015, aglycone's syntheses of Tcn-B were achieved by the Gadmann¹⁰ and Altmann¹¹ groups while the Zhu¹² group synthesized the macrolactonic core of a diastereomer of Tcn-B (OH on the C-18). In 2017, we also reported our own syntheses of the aglycone by designing two closely related pathways.¹³ The development of our strategy led us to discover a Kumada-Corriu reaction of vinyl sulfides catalyzed by Pd-nanoparticles.¹⁴ We also reexplored the Grigg's allene/alkyne cross-coupling and proposed an unprecedented mechanism.¹⁵ Until 2020, only the Gadmann's team managed to complete the total synthesis of Tcn-B, providing solutions to the challenging problem of the 1,2-*cis*-glycosylations but there was still room for innovation and improvement of selectivity.¹⁶ For the noviosylation step, the cyclic aglycone was found to show low reactivity toward a variety of different glycosyl donors or led exclusively to the α -glycosylated adduct. The mercury(II) Helferich's protocol¹⁷ was then used for the glycosylation of an acyclic triene fragment with a noviosyl bromide giving the corresponding product in 63% yield (α/β : 1/3). After assembly and cyclization of the aglycone, the rhamnosylation was carried out on the macrolide with an acetimidate glycosyl donor delivering the fully protected Tcn-B (α/β : 1/4, 62%). Note that recently De Brabander used similar glycosylation strategies for the synthesis of the northern and southern glycosylated fragments of Tcn-B.¹⁸ The total syntheses of tiacumicin A¹⁹, mangrolide A²⁰ and D,²¹ three congeners of tiacumicin B with a slightly simplified aglycone structure were also achieved.

In this article we wish to report with more details on our total synthesis of Tcn-B,²² highlighting the original glycochemistry that we had to develop to achieve this goal. Displaying axial C-O bonds at C2, D-rhamnose and D-noviose can both be related to D-mannose derivatives, in which the methylhydroxyl group in the C5-position has been replaced by a methyl or a gem-dimethyl group, respectively. For this series, the glycosylation reactions leading to 1,2-*cis* derivatives are particularly challenging since the α -compound is favored for steric and thermodynamic (anomeric effect) reasons and its C-2 configuration precludes the application of conventional neighboring group participation effects.²³

Our total synthesis of Tcn-B relies on our syntheses of the Tcn-B aglycone,¹³ that are based on original and selective assembly of the three main regions of the molecule. We had originally imagined to sequentially glycosylate the aglycone, but finally opted for two more convergent retrosynthetic plans. In these scenarios, Tcn-B was disconnected into fragments **A1**, **B1**, and **C** or into fragments **A2**, **B2**, and **C** (Figure 2).¹³ Connection of fragment **A1** together with fragment **B1** would first be achieved using ruthenium-catalyzed cross metathesis while a Suzuki coupling would allow the assembly of vinylbromide **B2** with the boronic ester **A2**. In both cases, following these steps, a ring-size selective macrolactonization would deliver a monoglycosylated aglycone having an unprotected OH at C-11 ready for the noviosylation step with fragment **C**. Knowing that the macrolactone has been described as a reluctant glycosyl acceptor, this challenging late β -glycosylation carried a high risk of failure.¹¹

Figure 2. Pathways to tiacumicin B (Tcn-B).

Preparation of the eastern part with 2-*O*-methyl-D-rhamnose.

For the preparation of the rhamnose fragments **B1** or **B2**, we initially considered using the mannose β -glycosylation method described by Kahne²⁴ and Crich.²⁵ The activation of a mannosyl donor having a 4,6-benzylidene and an anomeric sulfoxide or a thioaryl leaving group

would allow the selective formation of the β -mannoside compound through S_N2 displacement of the corresponding α -anomeric triflate. The obtained glycoside would therefore be later functionalized into a rhamnoside derivative.

Scheme 1. Synthesis of β -mannoside **4**. BSP: 1-benzenesulfinyl piperidine, TTBP: 2,4,6-tri-*tert*-butylpyrimidine.

To this aim, thioglycosyl donor **1** was prepared²⁶ and submitted to Crich glycosylation conditions using pre-activation at -60 °C by 1-benzenesulfinyl piperidine (BSP) in the presence of 2,4,6-tri-*tert*-butylpyrimidine (TTBP) and Tf_2O (Scheme 1).²⁷ The intermediate triflate was then allowed to react with the acceptor **3**^{3b} leading to the desired glycosylation product in a good 94% yield and a good selectivity of 1/6 in favor of the β -adduct. The stereochemistry was determined by measuring the C-1/H-1 coupling constant ($^1J_{C-1, H-1} \approx 160$ Hz if H-1 is axial and $^1J_{C-1, H-1} \approx 170$ Hz if H-1 is equatorial) derived from its 1H -coupled heteronuclear single quantum correlation spectrum. Unfortunately, we did not succeed in further functionalizing this compound into the desired rhamnoside. We then turned to a glycosylation using the phenylthiorhamnosyl donor **12** comprising the homodichloroorsellinate ester at the 4-position and a picoloyl group (Pico) at *O*-3 (Scheme 3, Table 1)). This glycosylation strategy, developed by Demchenko, involves intermolecular H-bonding between the nitrogen of the picoloyl group and the acceptor, directing the selective facial attack on the glycosyl donor on the same side as the Pico group.²⁸ We started with the synthesis of orsellinate derivative **7** needed for the esterification at the 4-position of *S*-Phenyl-2-*O*-methyl-1-thio- α -D-rhamnopyranoside **10**.²² This compound can be synthesized in a straightforward manner from commercially available carboxylic acid **5a** (Scheme 2). After dichlorination in acetic acid²⁹ and formation of the cyclic ester using acetone and trifluoroacetic anhydride in trifluoroacetic acid,³⁰ the resulting phenol was protected as a 2-naphthylmethylidene ether (Nap) **6**. This protecting group was chosen as it could be removed using the same conditions (DDQ) as for 4-methoxybenzyl (MPM) ethers, already present on the aglycone. Lithiation of the benzylic methyl group of **6** with LDA at -78 °C followed by trapping the corresponding anion with methyl iodide allowed the formation of the targeted **7**. Unfortunately, this reaction proved unreproducible giving yields ranging from 53 to 10% on a larger scale. For this reason, we chose a longer but reliable sequence starting from commercially available ethyl orsellinate **5b**. The latter was chlorinated and both phenols protected as *tert*-butyldimethylsilyl ethers provided **8**.²² With this compound, the sequence of the deprotonation of the benzylic methyl group/methylation proceeded in an almost quantitative yield to furnish reproducibly the homologated compound. After acidic hydrolysis with concentrated sulfuric acid to the deprotected carboxylic acid **9**, treatment with acetone and triflic anhydride in trifluoroacetic acid furnished the cyclic ester. Note that no reaction took place using trifluoroacetic anhydride instead of Tf_2O . The remaining free phenol was finally protected, supplying Nap ether **7**.

Scheme 2. Synthesis of orsellinate derivative **7**. TFA: trifluoroacetic acid, TFAA: trifluoroacetic anhydride, NapBr: 2-naphthalene-methyl bromide, LDA: Lithium di-*iso*-propylamide

The synthesis of donor **12** was performed from diol **10** obtained in a few steps from phenyl-2,3,4,6-tetra-*O*-acetyl-1-thio- β -D-mannopyranoside (Scheme 3).²² Esterification of this diol using homodichloroorsellinate ester **7** and NaH in THF led to compound **11**. As observed by Gademann *et al.*,¹⁹ the *O*-3 position is first esterified and the desired adduct is formed after ester migration on a prolonged reaction time. The liberated phenol was then protected as 2-Nap ether and the *O*-3 position of the rhamnoside was esterified with picolinic acid to give the desired donor **12**.

Scheme 3. Synthesis of rhamnopyranosyl donors **12-14**. DCC: dicyclohexylcarbodiimide, DMAP: 4-dimethylaminopyridine, *m*-CPBA: 3-Chloroperoxybenzoic acid.

Rhamnosyl donor **12** was first engaged in a glycosylation reaction with the monoprotected diol **16a** as the acceptor (Table 1, entry 1). This compound was prepared from 2,2-dimethyl-1,3-dioxan-5-one³¹ after a Wittig reaction with bromotrimethylphenylphosphonium bromide in the presence of sodium bis(trimethylsilyl)amide (NaHMDS) in THF (Scheme 4). This was followed by deprotection of the acetonide and monoprotection of the resulting diol as TBS-ethers **15a** and **15b**.³² A Suzuki cross coupling of a mixture of **15a,b** with phenylvinylboronic acid using catalytic Pd(OAc)₂ combined with 2-dicyclohexylphosphino-2',6'-dimethoxybiphenyl (Sphos) as ligand and potassium phosphate in THF/water at room temperature was then carried out giving **16a** and **16b** that could be separated using preparative HPLC.

Scheme 4. Synthesis of acceptors **15a** and **16a**. NaHMDS: sodium bis(trimethylsilyl)amide, PTSA: *para*-toluene sulfonic acid, SPhos: 2-dicyclohexylphosphino-2',6'-dimethoxybiphenyl.

The glycosylation reaction of **16a** was carried out in 1,2-dichloroethane (DCE) with dimethyl(methylthio)sulfonium triflate (DMTST), a classical promoter used in H-bond-mediated Aglycone Delivery (HAD). However, no glycosylation adduct was obtained with donor **12**, **16a** proving very unstable under these conditions. Acceptor **15a** was then used and with DMTST (2 eq.) in DCE at room temperature, we were pleased to obtain the glycosylation product **18** in 85% yield (Table 1, entry 2). A good α/β selectivity of 1/6.5 was also achieved suggesting that HAD might have worked to mediate a β -selective glycosylation through an intermediate such as **21**. Recrystallization allowed us to separate both anomers and to unambiguously determine the configuration of the major one through X-ray crystal diffraction analysis (Figure 3).³³ The reaction was also carried out in dichloromethane using *N*-iodosuccinimide (NIS, 1.2 eq.) and a catalytic amount of triflic acid (TfOH, 0.24 eq.), which gave a good yield (86%) but with a slightly lower stereoselectivity ($\alpha/\beta = 1:4$) (Table 1, entry 3).

Figure 3. X-ray crystal structure of **18**.

Table 1. Rhamnosylation conditions of different acceptors with donors **12-14**.

Entry	D (eq.)	A	Conditions ^a	P, yield (%) ^b	α/β ^c
1	12 (1.3)	16a	DMTST ^d	-	-
2	12 (1.3)	15a	DMTST ^d	18 , 85	1/6.5
3	12 (1.3)	15a	NIS, cat. TfOH ^e	18 , 86	1/4
4	12 (1.3)	17	DMTST ^d	-	-
5	12 (1.3)	17	NIS, TfOH cat. ^e	-	-
6	12 (1.3)	17	BSP, TTBP, Tf ₂ O, -60 °C ^f	-	-
7	12 (1.3)	17	NIS, TfOH ^g	19 , 75-86	1/3-1/4
8	13 (1.3)	17	Tf ₂ O, -50 °C ^h	19 , 52	1/12
9	13 (1.3)	17	Tf ₂ O, -70 °C ^h	19 , 48	1/16
10	13 (1.7)	17	Tf ₂ O, -70 °C ^h	19 , 64	1/20
11	13 (2.1)	17	Tf ₂ O, -70 °C ^h	19 , 70	1/20
12	13 (1.3)	17	Tf ₂ O, -70 °C ⁱ	19 , 20	1/2
13	14 (1.7)	17	Tf ₂ O, -70 °C ^h	20 , 95	20/1

^a Reaction performed in CH₂Cl₂ (c 0.01M) with 4Å MS unless otherwise stated. ^b After chromatography on silica gel. ^c Ratio and stereochemistry determined by ¹H NMR analysis of the crude mixture and by measuring ¹J_{C1,H1} coupling respectively. ^d With 2.6 eq. of the promotor in DCE as solvent from 0 °C to r.t. ^e With 1.2 eq. of NIS and 0.24 eq. of TfOH from -40 °C to r.t. ^f With 1.3 eq. of BSP, 2.5 eq. of TTBP and 1.4 eq. of Tf₂O. ^g With 1.2 eq. of NIS and 1 eq. of TfOH from -40 °C to r.t. ^h With 1.3-1.9 eq. of Tf₂O, 3.5-4.6 eq. of DTBMP and 4.2-6 eq. of ADMB. ⁱ Preactivation protocol.

Scheme 5. Functionalization of **18**. CSA: camphor sulfonic acid.

Before optimizing this glycosylation step, we chose to test the synthetic sequence leading to the key building block **B1** necessitating a terminal carboxylic acid group. Thus, the Pico group in **18** was smoothly and selectively removed with Cu(OAc)₂ in dichloromethane/methanol (Scheme 5). This was followed by a Pd-catalyzed Suzuki reaction with phenylvinylboronic acid using catalytic Pd(OAc)₂/Sphos and K₃PO₄ in THF/water at room temperature for 18 hours. Following these mild conditions, we noticed the presence of

about 20% of products displaying deprotected phenols on the orsellinate moiety. Direct treatment of the crude reaction mixture with NapBr in the presence of K_2CO_3 in DMF allowed us to reinstall the protective groups and obtain the targeted diene compound **23** in 73% yield. The 3-OH on the rhamnoside moiety was then protected as a TBS-ether and the primary alcohol was deprotected using camphor sulfonic acid in methanol/ CH_2Cl_2 , providing **24** in a 75% yield. Unfortunately, we did not succeed in oxidizing this compound into the corresponding carboxylic acid prior to the next Ru-catalyzed cross metathesis with fragment **A1**.

We then turned our attention toward the glycosylation of the alternative acceptor **17** already bearing the carboxylic function. Contrary to previous results with **15a**, activation of **12** with DMTST or NIS/cat. TfOH were disappointingly unsuccessful since no glycosylation adduct was observed (Table 1, entries 4 and 5). Activation with the BSP/ Tf_2O method at $-60\text{ }^\circ\text{C}$ in the presence of TTBP was also tried without success leading to the degradation of the acceptor (Table 1, entry 6). However, by increasing the amount of TfOH (0.92 eq./donor)³⁴ used in combination with *N*-iodosuccinimide (1 eq./donor) in CH_2Cl_2 at $-40\text{ }^\circ\text{C}$ to r.t. produced **19** in a 76% yield as a mixture of anomers (α/β : 1/4) (Table 1, entry 7), that could be separated by preparative HPLC. Note that in this particular case, the excess of TfOH could disrupt the HAD pathway and the glycosylation would follow a different mechanism, probably involving the protonation of the nitrogen on the picoloyl and formation of a glycosyl triflate.³⁵ By changing parameters such as temperature, promoter or donor amount, dilution, etc., we were not able to obtain a better selectivity. Seeking for higher β -selectivity, we decided to explore the reaction with sulfoxide **13**, a type of anomeric leaving group never before used in combination with a directing picoloyl group. Following *m*-CPBA oxidation of sulfide **12**, the activation of donor **13** (1.3 eq.) was first examined in CH_2Cl_2 at $-50\text{ }^\circ\text{C}$ using Tf_2O in the presence of the acceptor **17**, 2,6-di-*tert*-butyl-4-methylpyridine (DTBMP) as acid scavenger, and 4-allyl-1,2-dimethoxybenzene (ADMB) (Table 1, entry 8).^{36,37} We were pleased to find that under these conditions the desired glycosylated compound **19** with an upgraded facial selectivity (α/β : 1/12) and a yield of 52%. Higher yield (70%) and selectivity (α/β : 1/20) could be further attained by increasing the donor amount (2.1 eq.) and by lowering the temperature of the activation at $-70\text{ }^\circ\text{C}$ (see Table 1, entries 8 to 11 for comparison). To verify the influence of the remote picoloyl group in stereodirecting the nucleophilic attack on the β -face by H-bonding, we carried out two control experiments. The first consisted in performing the reaction by pre-activating the donor using Tf_2O at $-70\text{ }^\circ\text{C}$ for 15 minutes followed by the addition of the acceptor (Table 1, entry 12). These conditions led to the formation of the expected glycoside **19** in only 20% yield and poor selectivity (α/β : 1/2). The second control experiment was achieved with donor **14** having a benzoyl group in place of the picoloyl (Table 1, entry 13). This reaction led to pure α -glycosylation product **20**, indicating that the HAD mechanism may effectively take place only with the picoloyl group. The picoloyl of **19** was then removed with $Cu(OAc)_2$ in CH_2Cl_2 /MeOH, and replaced by a TBS group to lead to fragment **B2**.

Assemblage of A2 and B2 and macrolactonization

We could easily scale up our synthesis of the aglycone fragment **A2**. The convergent step of Pd-catalyzed cross-coupling of boronic ester **A2** with rhamnoside **B2** proceeded cleanly with catalytic $Pd(OAc)_2$ /Ruphos and potassium phosphate in a mixture of THF/water at r.t., furnishing ester **25** in 79% yield (Scheme 6). As reported by Nicolaou *et al.*,³⁸ the use of Me_3SnOH allowed us to hydrolyze selectively the methyl ester in the presence of the orsellinate on the rhamnoside moiety. Performed in toluene at $120\text{ }^\circ\text{C}$, the reaction led cleanly to seco-acid in 70 to 93% yields. With the latter in hand, we then focused on a ring-size selective macrolactonization using Yamaguchi conditions as described for the synthesis of our aglycone.¹³ However, the transposition of these conditions on this substrate led to the desired hemi-glycosylated tiacumicin B **26** in a low 23% yield. With the Boden-Keck's protocol,³⁹ the yield reached 58% yield, but **26** proved to be a mixture of two products resulting from the isomerization of the C4-C5 alkene. Finally, a far cleaner and reproducible macrolactonization was achieved using the Shiina's conditions⁴⁰ with 2-methyl-6-nitrobenzoic anhydride furnishing **26** in 72% yield, and an isomerization minimized at 15%. This selective strategy allowed us to keep the free OH at C-11 so that **26** could be directly engaged in the next glycosylation step.

Scheme 6. Assemblage of **A2** and **B2** and macrolactonization to monoglycosylated **26**. Ruphos : 2-dicyclohexylphosphino-2',6'-diisopropoxybiphenyl.

Preparation of the western part with D-noviose and noviosylation step.

For the noviosylation, we had first programmed Intramolecular Aglycone Delivery (IAD) using a silicon tether anchoring **26** to the 2-*O* position of the noviosyl moiety.⁴¹ The axial configuration of the 2-hydroxyl group of the donor being favorable, this approach would *a priori* provide a 1,2-*cis* glycosidic linkage with complete β -stereocontrol. For the preparation of the required noviosyl donor, we started from D-arabinose (Scheme 7). Selective acetonide protection of the *cis*-diol, oxidation of the hemiacetal with trichloroisocyanuric acid (TCCA) and catalytic TEMPO followed by silylation of the remaining 2-OH group provided lactone **27**. Introduction of the gem-dimethyl groups was achieved using the reaction of **27** with the methyl Grignard reagent and the corresponding diol was re-oxidized to lactone **28**. Dibal-H reduction led to the lactol and a mild acidic treatment with formic acid allowed the selective removal of the acetonide protection, prior to the introduction of a thiophenyl group at the anomeric position. The resulting diol **29** was protected as dichloroacetyl esters, which upon treatment by $(i\text{PrCO})_2\text{O}$ in MeCN with a catalytic amount of $\text{Sc}(\text{OTf})_3$ allowed the direct replacement of the TBS group by an isobutyrate.⁴² The two dichloroacetates in the obtained compound were then selectively removed with *sym*-collidine in MeOH giving diol **30**. The latter was then selectively mono-alkylated (NapBr, CsF) after the prior formation of the stannylene **31** giving predominantly the compound **32** with 2-naphthylmethylidene ether at *O*-2 position (2-*O*-Nap/3-*O*-Nap = 95/5) in 80% yield.

Scheme 7. Preparation of the noviosyl thioglycoside **32** from D-arabinose, conformational preference of diol **30** and likely structure of stannylene **31** with an equatorial C2-O2 bond. TEMPO: 2,2,6,6-tetramethyl-1-piperidinyloxy, TCCA: trichloroisocyanuric acid.

Scheme 8. Preparation of the silaketals **35-37** and glycosylation. NIS : N-iodosuccinimide; DTBMP: 2,6-Di-tert-butyl-4-methylpyridine.

This unexpected selectivity can be rationalized by the preferred conformation of the starting diol **30**, which is not the conventional 4C_1 chair but the 1C_4 chair, certainly imposed by the presence of the gem-dimethyl group. This was confirmed by the ${}^3J_{1,2}$ coupling constant of 9.5 Hz in the 1H NMR spectrum of diol **30** indicating a H-1/H-2 *trans* diaxial orientation. The equatorial OH bond being more accessible, this accounts for the selectivity of this protection.

We chose to investigate IAD approach with **32**, although to our knowledge, no other example has been described with a glycosyl acceptor linked to the 3-*O* position of the donor (Scheme 8). Starting our study with (–)-menthol **33** as a model, the corresponding silaketal **35** could be easily synthesized with the recently reported method of Montgomery.^{41c} After treatment of noviosyl adduct **32** with dimethylchlorosilane, the corresponding alkoxyisilane reacted with (–)-menthol **33** in the presence of $B(C_6F_5)_3$ as catalyst. As indicated by the 1H NMR spectrum, the silylated compound **35** also adopts exclusively a 1C_4 chair conformation. Considering that intermediate oxonium **38**, resulting from **35**, must display a half-chair conformation (Scheme 8), the alcohol to be transferred must then occupy a position favorable for the formation of the desired β -derivative. Optimized glycosylation conditions on **35** were obtained when the reaction was carried out with NIS (1.3 eq.) and TMSOTf (1.8 eq.) in CH_2Cl_2 in the presence of DTBMP (3 eq.) at $-40\text{ }^\circ C$ for 1 h. Treatment of the reaction mixture with a solution of TBAF in THF led to the desired glycosylation product **39a** (X = H) in 55% yield and as a single β -anomer. In this case, the obtained compound **39a** adopted a 4C_1 conformation with a measured ${}^1J_{C-1,H-1}$ of 159 Hz confirming the β -configuration. The moderate yield obtained here is likely due to the presence of the TBAF reagent that can cleave the *i*-butyrate ester. However, without TBAF treatment, we isolated, after silica gel chromatography, two stable glycosylation adducts bearing various silyl groups (**39b**, X = Me_2SiOH ,⁴³ and **39c**, X = $Me_2SiOTMS$ ⁴⁴) at the 3-*O* position of the novioside. To avoid their formation, an anhydrous HCl solution in methanol was added after completion of the glycosylation which delivered targeted product **39a** in a 81% yield with complete β -stereocontrol. The reaction was also carried out with the more complex alcohol **34**, a synthetic precursor of the aglycone fragment **A2**. The formation of the silaketal was performed as with (–)-menthol which produced **36** exclusively in a 1C_4 chair conformation and a good 76% yield. Using the glycosylation conditions described previously and acid treatment, the corresponding adduct **40** was obtained in 41% yield also as a single β -anomer. As for **39**, the obtained glycosylated compound **40** adopted a 4C_1 conformation with a measured ${}^1J_{C-1,H-1}$ of 159 Hz. Despite this unsatisfactory result, we decided to attempt the reaction with the semi-glycosylated tiacumicin B **26**. In this case we failed at preparing the silaketal using Montgomery's conditions but good results were obtained with the preliminary formation of the chloroalkoxyisilane^{41a} that reacted with **26** providing the corresponding dialkoxyisilane **37** in 73% yield. Unfortunately, glycosylation using NIS and TMSOTf in DCM at $-40\text{ }^\circ C$ followed by an HCl treatment led to degradation of the compound without evidence of the formation of the targeted glycosylated adduct.

Scheme 9. Preparation of the noviosyl donors **D** (**41-42**) and glycosylation (structure of R^1OH in Scheme 8).

We then shifted to another strategy based this time on an H-bond-mediated Aglycone Delivery (HAD) approach involving the use of noviosyl donor **41** bearing a Pico group at the 3-position (Scheme 9). Following esterification of **32** with picolinic acid, the corresponding sulfide **41** was engaged in a glycosylation reaction with (–)-menthol as the acceptor (Table 2, entry 1). The reaction was carried out with DMTST as the promoter and led predominantly to the β -compound **43** (α/β : 1/5) in 83 % yield. With elaborated unsaturated alcohol **34**, this approach led to poor yields (11 to 21%) either with DMTST (Table 2, entry 2) or NIS/TfOH (Table 2, entry 3) and unfortunately turned out unsuccessful with macrolactonic acceptor **26** since no glycosylated adduct was detected.

The success of the above-mentioned rhamnosylation led us to consider that sulfoxide **42** derived from sulfide **41** could be a far more reactive donor. A first trial with (–)-menthol as acceptor under Tf_2O activation at $-70\text{ }^\circ C$ revealed the potency of this method, as the β -anomer **43** was obtained as the only adduct in 66% yield (Table 2, entry 4). With alcohol **34**, the use of the sulfoxide approach proved to be more efficient as well providing β -glycosylation product **44** in a 62% yield (Table 2, entry 5). Moreover, these reaction conditions applied to the hemi-glycosylated tiacumicin B **26** delightfully furnished the desired noviosylated product **45** in 68% yield, with high facial selectivity (Table 2, entry 6, $\alpha/\beta > 1/20$).

Table 2. Glycosylation conditions of donors D (41–42) with different acceptors.

Entry	D (eq.)	R ¹ -OH	Conditions ^a	P, yield (%) ^b	α/β ^c
1	41 (1.3)	33	DMTST ^d	43, 83	1/5
2	41 (1.3)	34	DMTST ^d	44, 11	nd
3	41 (1.3)	34	NIS, TfOH ^e	44, 21	nd
4	42 (1.3)	33	Tf ₂ O, -70 °C ^f	43, 66	>1/20
5	42 (2.1)	34	Tf ₂ O, -70 °C ^g	44, 62	1/20
6	42 (2.1)	26	Tf ₂ O, -70 °C ^g	45, 68	>1/20

^a Reaction performed in CH₂Cl₂ (*c* 0.01M) with 4Å MS unless otherwise stated. ^b After chromatography on silica gel. ^c Ratio and stereochemistry determined by ¹H NMR analysis of the crude mixture and by measuring ¹J_{C1,H1} coupling. ^d With 2.3 eq. of the promotor in DCE (0.02M) from 0 °C to r.t. ^e With 1.4 eq. of NIS and 1.8 eq. of TfOH from -40 °C to r.t. ^f With 1.5 eq. of Tf₂O and 3 eq. of DTBMP. ^g With 1.9 eq. of Tf₂O, 4.7 eq. of DTBMP and 4.2 eq. of ADMB.

Final deprotection stages

The last steps consisting in the removal of all protective groups (2 MPM, 3 Nap, 1 Pico and 1 TBS) from compound **45** proved unpleasantly more difficult than expected (Scheme 10). Using HF·NEt₃ in THF allowed us to remove first the TBS group located on the rhamnosyl moiety giving the corresponding alcohol. The 2 MPM as well as the Nap located on the novioside were then oxidized with DDQ in CH₂Cl₂/H₂O. At this stage, the two Nap groups protecting the phenol functions of the rhamnosyl moiety resisted these conditions at 0 °C, and a longer reaction time at 20 °C led to an intractable mixture of products. The removal of the picoloyl was then cleanly carried out using Cu(OAc)₂ in CH₂Cl₂/MeOH at 0 °C to produce **46**. This deprotection sequence order was important as the Pico group had to be removed after its neighboring Nap group to avoid the DDQ-promoted formation of a 2,3-*O*-naphthylmethylidene on the novioside. These three operations were performed with no intermediate purifications providing an overall yield of 74% of **46**. However, the unexpected problem of cleavage of the two Nap groups located on the two phenol functions remained to be addressed. Lewis acid-mediated treatment of the Nap led only to the degradation of the molecule. Pd-catalyzed hydrogenation (Pd/C, cyclohexene) allowed the phenol deprotection but along with the reduction of the C4-C5 alkene.⁴⁵ As above mentioned, we observed a partial loss of the Nap groups located on the phenol moiety during the Pd-catalyzed Suzuki reaction of **23** with phenyl vinylboronic acid (see Scheme 5 and text). Exploiting this observation, we finally discover a new and selective method of deprotection of Nap ether on phenol. After a few optimizations, we found that using Pd₂(dba)₃ as a catalyst in combination with 4 PPh₃ along with 1,3-dimethylbarbituric acid (NDMBA) as methyl naphthyl scavenger and pyridine in DMF at 80 °C provided selective and smooth deprotection conditions. This ultimate step supplied the target compound tiacumicin B in 73% yield, with chemical data identical to those of the naturally occurring compound.^{6b}

Scheme 10. Final deprotection steps. DDQ: 2,3-dichloro-5,6-dicyano-1,4-benzoquinone. NDMBA: 1,3-dimethylbarbituric acid

In conclusion we have achieved a convergent total synthesis of tiacumicin B, by assembling the three main regions of the molecule. A highly β -selective rhamnosylation of the C1-C3 fragment followed by a Suzuki cross-coupling allowed assembling the rhamnoside **19** with the C4-C19 aglycone fragment **A2**. A ring-size selective macrolactonization using Shiina conditions was carried out followed by a final highly selective β -noviosylation of the cyclic aglycone and removal of all the protecting groups. During our glycosylation studies, we discovered a novel variant of the Demchenko procedure thanks to the conjoint use of a phenylsulfoxide leaving-group and a remote 3-*O*-picoloyl group on the donor. This combination allowed us to solve the problem of 1,2-*cis* glycosylation with a sensitive and complex aglycone and to reach a remarkable facial selectivity relying on an H-bond-directed effect. This new procedure will certainly prove useful in addressing the biological relevance of the tiacumicin B carbohydrate sugars or for the preparation of a set of glycosylated analogues. We also found a new and effective method for the removal of a 2-naphthylmethylidene (Nap) ether protecting group on a phenol through the use of palladium catalysis.

ASSOCIATED CONTENT

Supporting Information

Crystallographic data for structure **18**, preparation of all of the compounds, ^1H and ^{13}C NMR spectra for new compounds **4**, **6**, **15-16**, **18**, **22-24**, **39-40**, **43-44**, ^1H and ^{13}C NMR spectra of the synthetic and natural Tcn-B

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CONFLICT OF INTEREST

The authors declare no conflict of interest.

REFERENCES

- 1 Traynor, K., *Am. J. Health Syst. Pharm.* **2011**, *68* (14), 1276-1276.
- 2 Tupin, A.; Gualtieri, M.; Leonetti, J.-P.; Brodolin, K., *The EMBO J.* **2010**, *29* (15), 2527-2537. b) Gualtieri, M.; Tupin, A.; Brodolin, K.; Leonetti, J.-P., *Int. J. Antimicrobial Agents* **2009**, *34* (6), 605-606.
- 3 Srivastava, A.; Talaue, M.; Liu, S.; Degen, D.; Ebright, R. Y.; Sineva, E.; Chakraborty, A.; Druzhinin, S. Y.; Chatterjee, S.; Mukhopadhyay, J.; Ebright, Y. W.; Zozula, A.; Shen, J.; Sengupta, S.; Niedfeldt, R. R.; Xin, C.; Kaneko, T.; Irschik, H.; Jansen, R.; Donadio, S.; Connell, N.; Ebright, R. H., *Curr. Opin. Microbiol.* **2011**, *14* (5), 532-543.
- 4 Kurabachew, M.; Lu, S. H. J.; Krastel, P.; Schmitt, E. K.; Suresh, B. L.; Goh, A.; Knox, J. E.; Ma, N. L.; Jiricek, J.; Beer, D.; Cynamon, M.; Petersen, F.; Dartois, V.; Keller, T.; Dick, T.; Sambandamurthy, V. K., *J. Antimicrob. Chemother.* **2008**, *62* (4), 713-719.
- 5 For recent reviews on chemistry and biology of tiacumicin B see : a) Roulland, E., *Synthesis* **2018**, *50* (21), 4189-4200. b) Dorst, A.; Gademann, K., *Helv. Chim. Acta* **2020**, *103* (4), e2000038.
- 6 a) Parenti, F.; Pagani, H.; Beretta, G., *J. Antibiot.* **1975**, *28* (4), 247-252. b) Coronelli, C.; White, R. J.; Lancini, G. C.; Parenti, F., *J. Antibiot.* **1975**, *28* (4), 253-259.
- 7 Martinelli, E.; Faniuolo, L.; Tuan, G.; Gallo, G. G.; Cavalleri, B., *J. Antibiot.* **1983**, *36* (10), 1312-1322.
- 8 Arnone, A.; Nasini, G.; Cavalleri, B., *J. Chem. Soc., Perkin Trans. 1* **1987**, 1353-1359.
- 9 a) Xiao, Y.; Li, S.; Niu, S.; Ma, L.; Zhang, G.; Zhang, H.; Zhang, G.; Ju, J.; Zhang, C., *J. Am. Chem. Soc.* **2011**, *133* (4), 1092-1105. b) Yu, Z.; Zhang, H.; Yuan, C.; Zhang, Q.; Khan, I.; Zhu, Y.; Zhang, C., *Org. Lett.* **2019**, *21* (18), 7679-7683.
- 10 Miyatake-Ondozabal, H.; Kaufmann, E.; Gademann, K., *Angew. Chem., Int. Ed.* **2015**, *54* (6), 1933-1936.
- 11 Glaus, F.; Altmann, K. H., *Angew. Chem., Int. Ed.* **2015**, *54* (6), 1937-1940.
- 12 Erb, W.; Grassot, J. M.; Linder, D.; Neuville, L.; Zhu, J., *Angew. Chem., Int. Ed.* **2015**, *54* (6), 1929-1932.
- 13 a) Jeanne-Julien, L.; Masson, G.; Astier, E.; Genta-Jouve, G.; Servajean, V.; Beau, J.-M.; Norsikian, S.; Roulland, E., *Org. Lett.* **2017**, *19* (15), 4006-4009. b) Jeanne-Julien, L.; Masson, G.; Astier, E.; Genta-Jouve, G.; Servajean, V.; Beau, J. M.; Norsikian, S.; Roulland, E., *J. Org. Chem.* **2018**, *83* (2), 921-929.
- 14 Jeanne-Julien, L.; Astier, E.; Lai-Kuen, R.; Genta-Jouve, G.; Roulland, E., *Org. Lett.* **2018**, *20* (5), 1430-1434.
- 15 Jeanne-Julien, L.; Masson, G.; Kouoi, R.; Regazzetti, A.; Genta-Jouve, G.; Gandon, V.; Roulland, E., *Org. Lett.* **2019**, *21* (9), 3136-3141.
- 16 Kaufmann, E.; Hattori, H.; Miyatake-Ondozabal, H.; Gademann, K., *Org Lett* **2015**, *17*, 3514-3517.
- 17 Helferich, B.; Wedemeyer, K. F., *Liebigs Ann. Chem.* **1949**, *563*, 139-145.
- 18 Hollibaugh, R.; Yu, X.; De Brabander, J. K., *Tetrahedron* **2020**. DOI : 10.1016/j.tet.2020.131673
- 19 Hattori, H.; Kaufmann, E.; Miyatake-Ondozabal, H.; Berg, R.; Gademann, K., *J. Org. Chem.* **2018**, *83* (13), 7180-7205.

- ²⁰ Hattori, H.; Roesslein, J.; Caspers, P.; Zerbe, K.; Miyatake-Ondozabal, H.; Ritz, D.; Rueedi, G.; Gademann, K., *Angew. Chem., Int. Ed.* **2018**, *57* (34), 11020-11024.
- ²¹ Hattori, H.; Hoff, L. V.; Gademann, K., *Org. Lett.* **2019**, *21* (9), 3456-3459.
- ²² a) Norsikian, S.; Tresse, C.; François-Eude, M.; Jeanne-Julien, L.; Masson, G.; Servajean, V.; Genta-Jouve, G.; Beau, J. M.; Roulland, E., *Angew. Chem., Int. Ed.* **2020**, *59* (16), 6612-6616. b) Norsikian, S.; Tresse, C.; François-Eude, M.; Jeanne-Julien, L.; Masson, G.; Servajean, V.; Genta-Jouve, G.; Beau, J.-M.; Roulland, E., *Angew. Chem.* **2020**, *132* (16), 6674-6678.
- ²³ For a recent review see: Nigudkar, S. S.; Demchenko, A. V., *Chem. Sci.* **2015**, *6* (5), 2687-2704.
- ²⁴ Kahne, D.; Walker, S.; Cheng, Y.; Van Engen, D., *J. Am. Chem. Soc.* **1989**, *111*, 6881.
- ²⁵ Crich, D.; Sun, S., *J. Org. Chem.* **1996**, *61* (14), 4506-4507.
- ²⁶ Elferink, H.; Mensink, R. A.; Castelijns, W. W. A.; Jansen, O.; Bruekers, J. P. J.; Martens, J.; Oomens, J.; Rijs, A. M.; Boltje, T. J., *Angew. Chem., Int. Ed.* **2019**, *58* (26), 8746-8751.
- ²⁷ Crich, D.; Smith, M., *J. Am. Chem. Soc.* **2001**, *123* (37), 9015-9020.
- ²⁸ a) Yasomane, J. P.; Demchenko, A. V., *J. Am. Chem. Soc.* **2012**, *134* (49), 20097-20102. b) Pistorio, S. G.; Yasomane, J. P.; Demchenko, A. V., *Org. Lett.* **2014**, *16* (3), 716-719.
- ²⁹ Milligan, R. F.; Hope, F. J., *J. Am. Chem. Soc.* **1941**, *63* (2), 544-544.
- ³⁰ Marriott, J. H.; Barber, A. M. M.; Hardcastle, I. R.; Rowlands, M. G.; Grimshaw, R. M.; Neidle, S.; Jarman, M., *J. Chem. Soc., Perkin Trans. 1* **2000**, (24), 4265-4278.
- ³¹ González-García, E. M.; Grognum, J.; Wahler, D.; Raymond, J.-L., *Helv. Chim. Acta* **2003**, *86* (7), 2458-2470.
- ³² As the reaction is completely unselective, both isomers were separated on preparative HPLC. The stereochemistry was determined by NMR spectroscopy using NOESY experiments.
- ³³ See the Supporting Information. CCDC-2027426 contains the supplementary crystallographic data for this paper. These data can be obtained free of charge from The Cambridge Crystallographic Data Centre via www.ccdc.cam.ac.uk/data_request/cif.
- ³⁴ a) Escopy, S.; Geringer, S. A.; De Meo, C., *Org. Lett.* **2017**, *19* (10), 2638-2641; b) Mannino, M. P.; Demchenko, A. V., *Chem. Eur. J.* **2020**, *26* (13), 2938-2946.
- ³⁵ Mannino, M. P.; Demchenko, A. V., *Chem. Eur. J.* **2020**, *26* (13), 2927-2937.
- ³⁶ ADMB scavenge phenylsulfenyl triflate, a highly reactive byproduct, which forms after activation of anomeric sulfoxides with Tf₂O. Gildersleeve, J.; Smith, A.; Sakurai, K.; Raghavan, S.; Kahne, D., *J. Am. Chem. Soc.* **1999**, *121* (26), 6176-6182.
- ³⁷ Zeng, J.; Liu, Y.; Chen, W.; Zhao, X.; Meng, L.; Wan, Q., *Top. Curr. Chem. (Cham)* **2018**, *376* (4), 27.
- ³⁸ Nicolaou, K. C.; Estrada, A. A.; Zak, M.; Lee, S. H.; Safina, B. S., *Angew. Chem., Int. Ed. Engl.* **2005**, *44* (9), 1378-1382.
- ³⁹ Boden, E. P.; Keck, G. E., *J. Org. Chem.* **1985**, *50* (13), 2394-2395.
- ⁴⁰ Shiina, I.; Kubota, M.; Oshiumi, H.; Hashizume, M., *J. Org. Chem.* **2004**, *69* (6), 1822-1830.
- ⁴¹ a) Stork, G.; Kim, G., *J. Am. Chem. Soc.* **1992**, *114*, 1087. Bols, M., *J. Chem. Soc., Chem. Commun.* **1992**, (12), 913-914. c) Walk, J. T.; Buchan, Z. A.; Montgomery, J., *Chem. Sci.* **2015**, *6* (6), 3448-3453.
- ⁴² Norsikian, S.; Holmes, I.; Lagasse, F.; Kagan, H. B., *Tetrahedron Lett.* **2002**, *43* (33), 5715-5717.
- ⁴³ Packard, G. K.; Rychnovsky, S. D., *Org. Lett.* **2001**, *3* (21), 3393-3396.
- ⁴⁴ Beignet, J.; Cox, L. R., *Org. Lett.* **2003**, *5* (22), 4231-4234.
- ⁴⁵ Smith, III, A. B.; Sfougataki, C.; Risatti, C. A.; Sperry, J. B.; Zhu, W.; Doughty, V. A.; Tomioka, T.; Gotchev, D. B.; Bennett, C. S.; Sakamoto, S.; Atasoylu, O.; Shirakami, S.; Bauer, D.; Takeuchi, M.; Koyanagi, J.; Sakamoto, Y., *Tetrahedron* **2009**, *65*, 6489-6509.