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Enantioselective Synthesis of the Lomaiviticin Aglycon Full Carbon Skeleton Reveals Remarkable Remote Substituent Effects During the Dimerization Event

Hong Geun Lee^[a], Jae Young Ahn^[a], Amy S. Lee^[a], and Matthew D. Shair^{*[a]}

Dedication ((optional))

Lomaiviticin A (**1**) and lomaiviticin B (**2**) are novel C_2 -symmetric diazobenzofluorene glycoside marine natural products (Figure 1).^[1] Lomaiviticin A (**1**) potently inhibits the growth of cultured cancer cell lines (GI_{50} values ranging from 0.007 nM to 72.0 nM), and both **1** and **2** exhibit impressive growth inhibition activity against Gram-positive bacteria. Compound **1** causes damage to DNA in vitro. It has been speculated that a reactive species derived from the diazobenzofluorenones of **1** and **2** causes damage to nucleic acids, leading to the cytotoxic activities of these molecules. Experimental studies have provided support for the formation of reactive species from diazofluorenone structures.^[2] However, it remains to be determined how the full structures of **1** and **2** react under physiologically relevant conditions and what cellular components are perturbed by **1** and **2**.

The structures of **1** and **2** are unprecedented and striking. The unusual structures of **1** and **2** pose interesting questions about how they are biosynthesized, especially how the C2-C2' bond is made, and questions about how to achieve syntheses of these molecules. Although a synthesis of **1** or **2** has not yet been achieved, many approaches to these molecules have been reported,^[3] including

our own enantioselective synthesis of the central (C-D-D'-C') ring system of **1**.^[3b] A synthesis of the full carbon skeleton of **1** and **2** has not yet been accomplished. To achieve syntheses of **1** and **2**, there are significant challenges to overcome; the most difficult of which is likely formation of the C2-C2' σ -bond (Figure 1, see red bond). The C2-C2' bond links two highly functionalized 'halves' of **1** and **2**, which closely resemble the related natural product kinamycins.^[4] The congested environment surrounding the C2-C2' bond, stereochemical control during bond formation, and the potential lability of the D and D' rings render the formation of this bond extremely challenging. Late-stage dimerization and formation of the C2-C2' bond, in which double-processing is kept at a minimum, would be the most efficient means of synthesizing **1** and **2**. Since the C2-C2' bond in **1** and **2** is part of a 1,4-diketone (C1-C2-C2'-C1'), a late-stage oxidative enolate coupling would be the ideal reaction to construct this bond. However, the high potential for β -elimination of the C3 tertiary carbinol from a C1-C2 enolate, coupled with the likely poor stereoselectivity of such an oxidative enolate coupling reaction, renders this approach unattractive. To circumvent these obstacles, we developed a strategy utilizing the oxidative enolate coupling of 7-oxanorbornanones to achieve the first enantioselective synthesis of the central ring system of the aglycon of **1** (Scheme 1, see black structures in brackets). The 7-oxanorbornanone structure prevents β -elimination at C3 due to stereoelectronic effects, and it provides perfect diastereoselectivity during the formation of the C2-C2' bond due to double diastereodifferentiation in the dimerization event. Herein we report stereoselective coupling of the monomeric units of **1** and **2** leading to the first synthesis of the full carbon skeleton of the aglycons of **1** and **2**. During the course of these studies, remarkable remote substituent effects during the key dimerization reaction have been observed, which may further complicate future syntheses of **1** and **2**.

As stated, our synthesis plan is to perform an oxidative dimerization of the enolate of **4** to afford **3** (Scheme 1). An annulation reaction^[5] between enone **5** and a known cyanophthalide reagent^[6] would provide access to **4** (Scheme 3, **12**). Enone **5** would be accessible from intermediate **6**, a compound for which we had previously developed an enantioselective synthesis. Initially, an anionic annulation reaction on an enone similar to **5**, but with C1 at the ketone oxidation state,

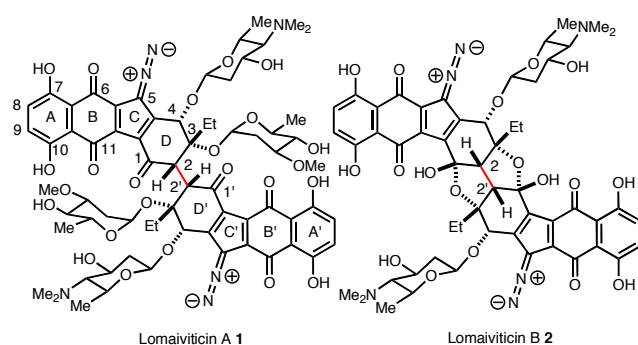
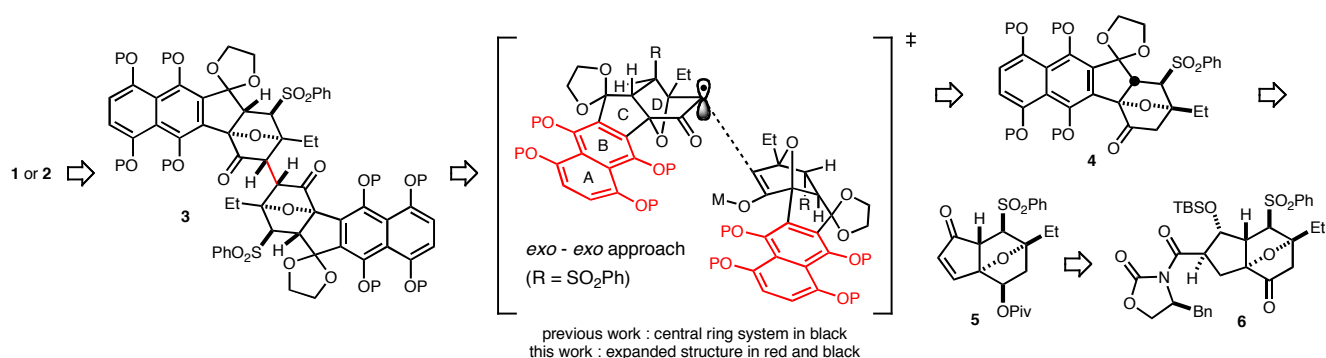


Figure 1. Structures of lomaiviticin A and B.

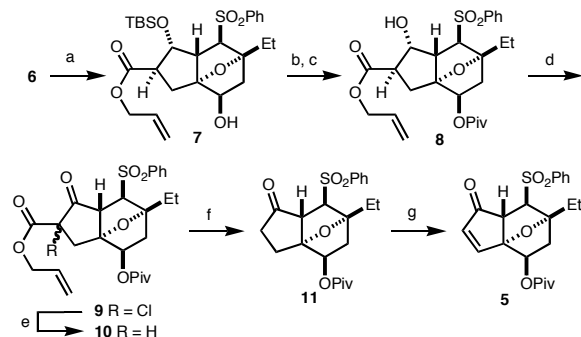
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Scheme 1. Synthesis plan.

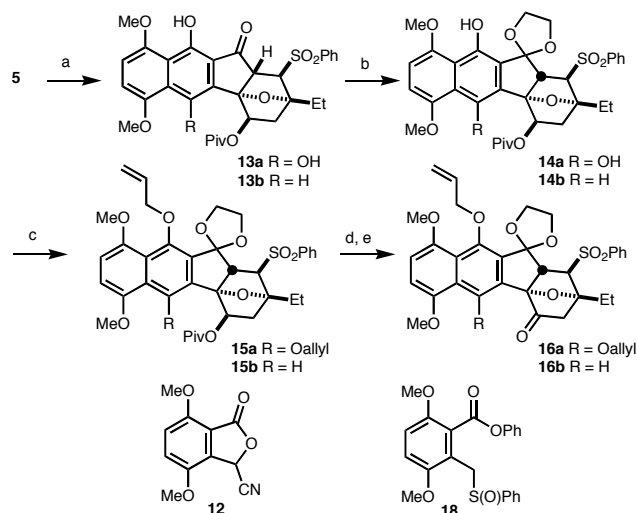
was attempted. Unfortunately, the basic conditions of the annulation were not compatible with the C1 ketone. Therefore, we were forced to protect the C1 ketone of **6** in its reduced form by treatment with NaBH₄ (Scheme 2). Surprisingly, **7** was obtained as the major product when the reduction was performed in allyl alcohol solvent (Scheme 2). We speculate that transesterification of the oxazolidinone chiral auxiliary occurs from the boronate of allyl alcohol. Quenching the remaining borohydride with acetone allowed us to isolate **7** in 89% yield. Protecting group manipulation of **7** afforded **8**. The C5 carbinol of compound **8** was then oxidized under standard Swern conditions. Surprisingly, the product was obtained as the chlorinated β -ketoester **9**.^[7] The reductive dechlorination of **9** with Zn afforded β -ketoester **10**, which underwent decarboxylative deallylation^[8] upon exposure to PdCl₂(PPh₃)₂ and *n*Bu₃SnH, furnishing ketone **11** in 85% yield. Finally, α -phenylselenation of the enolate of **11**, followed by oxidation, delivered enone **5**.^[9]



Scheme 2. Synthesis of enone **5**. Reagents and conditions: a) NaBH₄, allyl alcohol, -78 °C, 5 min; then acetone, 23 °C, 15 min, 89%; b) PivCl, pyridine, CH₂Cl₂, 40 °C, 24 h; c) TBAF, THF, 23 °C, 16 h, 91% over two steps; d) DMSO, (COCl)₂, -78 °C, 30 min; then Et₃N, 23 °C, 30 min; e) Zn dust, AcOH, 23 °C, 2 h, 96% over two steps; f) PdCl₂(PPh₃)₂, *n*Bu₃SnH, AcOH, toluene, 0 °C, 30 min; then 110 °C, 30 min, 85%; g) LHMDS, THF, -78 °C, 30 min; then PhSeBr, -78 °C, 5 min; then H₂O₂, CH₂Cl₂/THF (1:1), 0 °C, 4 h, 73%. Piv = trimethylacetyl, TBAF = tetrabutylammonium fluoride, THF = tetrahydrofuran, DMSO = dimethyl sulfoxide, LHMDS = lithium bis(trimethylsilyl)amide.

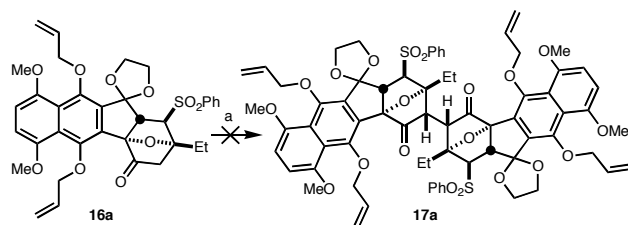
For the key annulation reaction, we used the cyanophthalide method developed by Kraus^[10] (Scheme 3). Addition of the anion of **12** to enone **5** afforded hydroquinone **13a** in 85% yield. The C5 ketone of **13a** was protected as a dioxolane.^[11] Protection of the two phenols of **14a** as their corresponding allyl ethers (compound **15a**), followed by reductive cleavage of the pivaloyl group, and subsequent oxidation at C1 afforded ketone **16a**, setting the stage for the key oxidative enolate coupling.

For synthesis of the C-D-D'-C' central ring system of **1** and **2**, we had developed conditions for the oxidative enolate coupling reaction involving deprotonation of the ketone with LHMDS at



Scheme 3. Synthesis of dimerization precursors **16a** and **16b**. Reagents and conditions: a) For **13a**: **12**, LHMDS, HMPA, THF, -78 °C, 30 min; then **5**, 50 °C, 3 h, 85%; for **13b**: **18**, LHMDS, THF, -60 °C, 6 h; then **5**, -78 °C, 10 min, 79%; b) 1,2-bis(trimethylsilyloxy) ethane Scheme 3. e, TMSOTf, CH₂Cl₂, 23 °C, 48 h, 76% for **14a**, 72% for **14b**; c) Cs₂CO₃, allyl bromide, DMF, 23 °C, 2 h, 87% for **15a**, 76% for **15b**; d) NaBHET₃, THF, 0 °C, 4 h; e) TPAP, NMO, 4 Å M.S., CH₂Cl₂, 23 °C, 2 h, 90% for two steps for **16a**, 88% for two steps for **16b**. LHMDS = lithium bis(trimethylsilyl)amide, HMPA = hexamethylphosphoramide, DMF = dimethyl formamide, TMSOTf = trimethylsilyl trifluoromethanesulfonate, TPAP = tetrapropylammonium perruthenate, NMO = 4-methylmorpholine-*N*-oxide.

-78 °C, addition of Cp₂FePF₆, and warming to -60 °C.^[12] We then applied these conditions to the oxidative dimerization of ketone **16a** (Scheme 4). Unfortunately, none of the desired dimer (**17a**) was formed under these conditions, and only starting material **16a** or decomposed material was isolated (Scheme 4).



Scheme 4. Attempted oxidative dimerization of **16a**. Reagents and conditions: a) LHMDS, HMPA, THF, -78 °C, 2 h; then Cp₂FePF₆, -60 °C, 72 h, 0%. LHMDS = lithium bis(trimethylsilyl)amide, HMPA = hexamethylphosphoramide, Cp₂FePF₆ = ferrocenium hexafluorophosphate.

In light of these disappointing results, we tried to rationalize why our C-D ring system published earlier underwent successful dimerization, but the A-D ring system here failed to dimerize. We speculated that non-bonded interactions may be developing in the

transition state, which could be inhibiting dimerization. Since little is known about the mechanism of this reaction and the trajectory by which substrates approach each other, we attempted to understand possible developing non-bonded interactions in the transition state by studying the ground state conformations of the desired product (compound **17a**).^[13] We examined compound **17a** (Figure 2) and each of the three staggered conformations about the C2-C2' bond, specifically H2-H2' dihedral angles of 60°, 180°, and -60° (see Figure 2). The conformation with a H2-H2' dihedral angle of 180° adopted an 'n' conformation, the name of which is given because the shape of the conformer resembles that of the letter 'n'. This conformer appeared to be very high in energy and unlikely to be a relevant conformer since the C3 and C3' ethyl groups were suffering from severe non-bonded interactions. The 60° H2-H2' dihedral angle conformation also exists in an 'n' conformation, but it does not suffer from the destabilizing C3-C3' ethyl interactions. However, we did notice potential non-bonded interactions between the C11 and C11' allyloxy groups (see red arrow). Finally, the -60° H2-H2' dihedral angle conformation exists in a 'z' conformation, which was also given due to the resemblance to the letter 'z' in shape, and in this conformation non-bonded interactions may develop between C11 (and C11') allyloxy groups and the C3' (and C3) ethyl groups. Since in both the 'n' and 'z' conformations, the C11 (and C11') allyloxy groups may be preventing dimerization, we decided to synthesize the tetracyclic dimerization precursor where the C11 allyloxy group is replaced with a hydrogen atom (**16b**, R= H). This forced us to devise a new annulation strategy to access this structure from enone **5**.

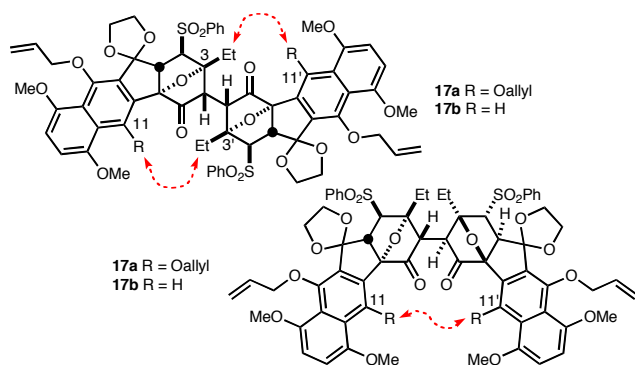
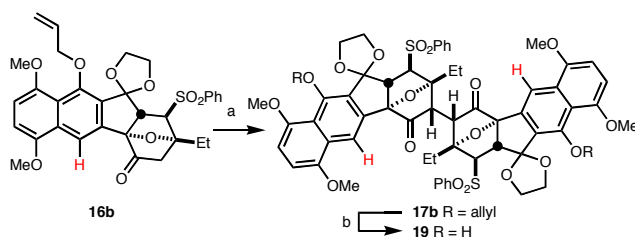


Figure 2. Postulated steric interactions in the dimeric products.

To construct **16b**, where the C11 substituent is a hydrogen atom, we synthesized Hauser-type sulfoxide annulation partner **18** (Scheme 3).^[14] Deprotonation of **18** with LHMDS at -60 °C, followed by exposure to enone **5**, delivered phenol **13b** in 79% yield. It is worth noting that the use of the phenyl ester of **18**,



Scheme 5. Oxidative dimerization of **16b**. Reagents and conditions: a) LHMDS, HMPA, THF, -78 °C, 2 h; then Cp₂FePF₆, -60 °C, 72 h, 80%; b) PdCl₂(PPh₃)₂, *n*Bu₃SnH, AcOH, CH₂Cl₂, 0 °C, 30 min, 90%. LHMDS = lithium bis(trimethylsilyl)amide, HMPA = hexamethylphosphoramide, Cp₂FePF₆ = ferrocenium hexafluorophosphate.

compared to the more commonly used methyl ester, dramatically enhanced the rate and yield of the annulation reaction—a finding that may be useful in other applications of this reaction.^[15] Following the process we developed previously, **13b** was converted to dioxolane **14b**. Allyl protection of the phenol to afford **15b**, followed by cleavage of the C1 pivaloyl group, and subsequent oxidation, delivered ketone **16b**. With ketone **16b** in hand, we again attempted to perform oxidative enolate coupling under identical conditions (LHMDS, HMPA, Cp₂FePF₆, -60 °C, THF). We were gratified to find that oxidative dimerization of **16b** afforded **17b** in 80% yield as a single diastereomer (Scheme 5).

To confirm the relative stereochemistry at C2 and C2', and to determine the conformation of dimeric structure **17b**, we set out to obtain an X-ray crystal structure. Fortunately, we discovered that removal of the allyl groups of **17b** afforded **19** as a crystalline solid (Scheme 5), which was subjected to X-ray crystallographic analysis to provide the structure shown in Figure 3.^[16] The crystal structure confirmed that the correct C2-C2' stereochemistry had been established and also that the compound can exist in an 'n'

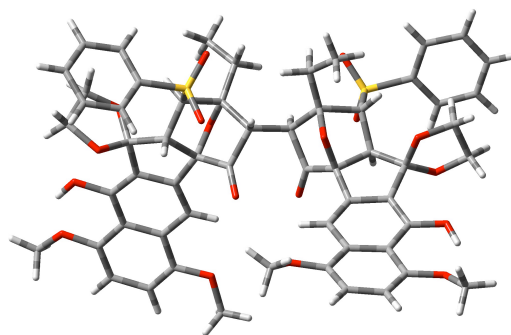


Figure 3. X-ray crystal structure of **19**.

conformation.

Closer inspection of the X-ray crystal structure revealed that the C11 hydrogen atom is in close proximity to the C1' ketone oxygen atom (and conversely the C11' hydrogen and C1 ketone oxygen; see Figure 4). In fact, the interatomic distance between H11 and O1' (and H11' and O1) is only 0.29 Å greater than the sum of their van der Waals radii. This close contact between the C11 substituent and O1' (as well as C11' and O1) may explain why the first attempted oxidative dimerization with a C11 allyloxy group failed. If, as in our first attempted dimerization, C11 bears an oxygen atom, its greater van der Waals radii of 1.52 Å would have suffered a non-bonded interaction with O1'. Interestingly, although our analysis of ground state conformations

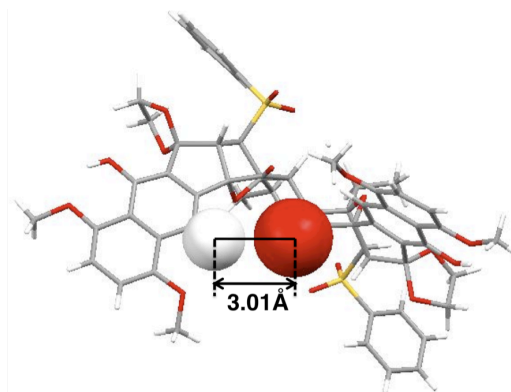


Figure 4. Interatomic distance of C11 hydrogen and C1' oxygen.

led us to convert C11 from an oxygen substituent to hydrogen, we were surprised to find that it may be a steric clash between C11 substituent and O1' that may be preventing dimerization.

Further examination of the X-ray crystal structure of **19** revealed unusual torsional strain in the C2-C2' bond, with a C1-C2-C2'-C1' dihedral angle of 29° (compared to a desired dihedral angle of 60°). The presence of C2-C2' torsional strain in **19** and the potential for severe non-bonded interactions in the crystal structure suggested to us that there may be a measurable barrier to rotation about C2-C2', resulting in atropisomerism. Unfortunately, heating to 140 °C or cooling to -80 °C did not result in a change in the 1H NMR spectrum of **19**, preventing us from determining whether **19** exhibits atropisomerism.

In summary, we report the first synthesis of the full carbon skeleton of the lomaiviticin aglycon. An important aspect of this study was demonstrating that the oxidative enolate coupling strategy we reported earlier can be applied to the full monomeric units of the lomaiviticins to assemble the full carbon skeleton, again with complete stereochemical control in the formation of the C2-C2' bond. During the course of these studies, we have discovered that remote oxygen substituents at C11 (and C11'), surprisingly, have dramatic effects on the success of the oxidative dimerization. It is interesting to speculate whether our observation that dimerization to form the lomaiviticin aglycon skeleton requires hydrogen atoms at C11 and C11' extends to other dimerization strategies or even to the biosynthesis of **1** and **2**. It may be that a biosynthetic dimerization also occurs with hydrogen at C11 and C11' and that late stage oxidation leads to the lomaiviticins. An X-ray crystal structure of the lomaiviticin aglycon skeleton revealed that a non-bonded interaction between substituents at C11 and O1' (and C11'-O1) may be suppressing oxidative dimerization when the C11 substituent is larger than hydrogen. In addition, the lomaiviticin aglycon skeleton exists (or at least crystallizes) in an unusual 'n' conformation with significant torsional strain around the C2-C2' σ -bond. The lomaiviticins are challenging synthesis targets and the advances reported herein are important steps in eventually achieving their syntheses.

Experimental Section

Oxidative dimerization of 16b: A two neck flask charged with a solid addition adaptor with ferrocenium hexafluorophosphate (2.32 g, 7.00 mmol, 5.00 equiv) in it was flame dried and charged with argon. Then the flask was charged with 8 mL of THF, HMDS (0.502 mL, 2.38 mmol, 1.70 equiv), 2.29 M *n*-BuLi solution in hexane (1.04 mL, 2.38 mmol, 1.70 equiv) and HMPA (0.487 mL, 2.80 mmol, 2.00 equiv) at -78°C, successively. 40 minutes later a solution of ketone **15b** (830 mg, 1.40 mmol, 1.00 equiv) in 10 mL of THF (plus 2 x 1 mL rinse) was cannulated to the LiHMDS-HMPA solution to give a pale yellow solution. 2 hours later, ferrocenium hexafluorophosphate was added to the reaction system from the solid addition adaptor. The originally formed deep blue suspension turned yellowish green in 30 minutes. The reaction mixture was stirred at -60 °C for 3 days. The reaction mixture was quenched with a saturated solution of NH₄Cl and warmed to room temperature. The aqueous phase was extracted with dichloromethane (3 x 20 mL) and the combined organic phase was dried with anhydrous Na₂SO₄. The crude product was purified by flash column chromatography (EtOAc/Hexane = 1/1 to 1.5/1) to give dimer **17b** (665 mg, 1.12 mmol, 80%) as a pale yellow solid.

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acknowledge Sanofi-Aventis and Eli Lilly, respectively, for graduate fellowships. Shao-Liang Zheng is acknowledged for X-ray crystallographic analysis.

Keywords: annulation • antitumor agents • natural products • remote steric effect • total synthesis

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- [16] CCDC 777057 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.

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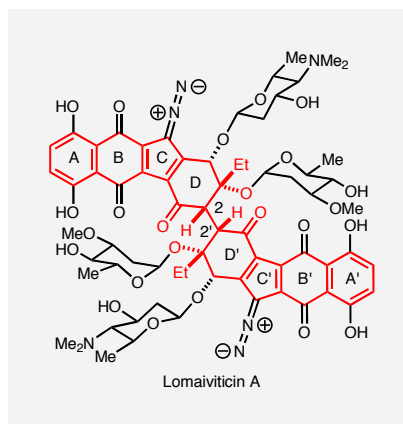
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Enantioselective Synthesis of the Lomaiviticin Aglycon Full Carbon Skeleton Reveals Remarkable Remote Substituent Effects During the Dimerization Event



A full carbon skeleton of natural product lomaiviticin has been synthesized. The approach features anionic annulation reactions to deliver A and B rings of the natural product, and stereoselective oxidative enolate coupling to form the central C2-C2' σ -bond. In the course of the investigation, a remarkable substituent effect at C11 position was observed.