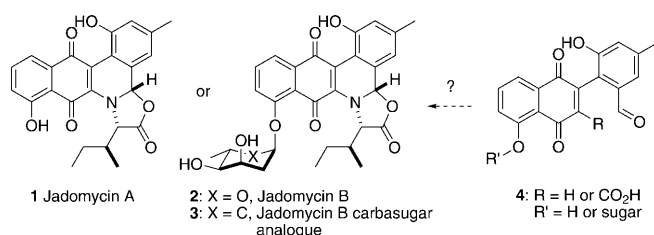


Total Synthesis of Jadomycin A and a Carbasugar Analogue of Jadomycin B**

Mingde Shan, Ehesan U. Sharif, and George A. O'Doherty*

Jadomyins A and B are unique members of the angucycline family of natural products with the unusual 8*H*-benzo[*b*]oxazolo[3,3-*f*]-phenanthridine ring system (Scheme 1). Both jadomyins A and B are produced by the Gram-positive soil bacteria *Streptomyces venezuelae* ISP5230

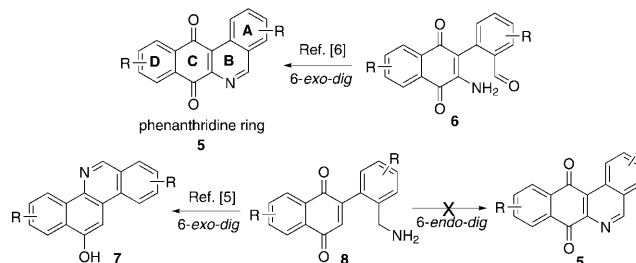


Scheme 1. Jadomycin A, B and its analogue.

under specific nutrient and environmental stress.^[1] They display important bioactivities, including antitumor, antimicrobial, anti-viral, as well as, aurora-B kinase inhibition and DNA-cleaving capacity.^[2–4]

There has been no total synthesis of jadomycin A or B, although libraries of jadomycin analogues have been produced through fermentation approaches.^[5] Several approaches to the benzo[*b*]-phenanthridine ring system have been developed (Scheme 2).^[6,7] From these studies one can conclude that for stereoelectronic reasons the B-ring is best constructed by a cyclization of a 2-amino-3-phenyl-naphthoquinone (e.g., **6** to **5**), rather than an amine cyclization onto the 2-position of an anthroquinone (e.g., **8** to **5**). However, these results stand in contrast to the proposed biosynthetic route (e.g., **1/2** through **4**, Scheme 1).

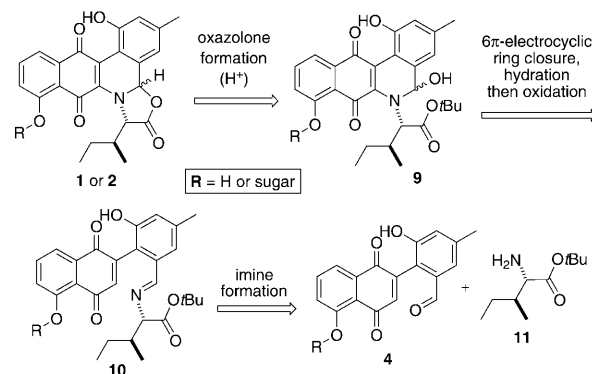
Our ongoing interest in the synthesis and study of biologically active carbohydrate containing natural products,^[8] along with these biosynthetic questions, made us interested in this class of natural products. Specifically, we were interested in testing our hypothesis that by employing a 6*π*-electrocyclic ring closure for the B-ring construction, one



Scheme 2. Retrosynthetic comparison for phenanthridine B-ring formation.

could skirt the stereoelectronic disadvantages of the 6-endo-dig ring closures (**10** to **9**).^[9] To the best of our knowledge this is the first time that switching to an electrocyclic ring closure was used to change the regioselectivity for ring closure. This may also be the biosynthetic mode of ring closure and thus implying that the B-ring cyclization step does not need enzyme mediation to control the regioselectivity of cyclization.

Retrosynthetically, we envisioned **1** and **2** arising from the condensation of a protected amino acid **11** and aldehyde **4** to form imine **10**. A 6*π*-electron electrocyclic ring closure would form the desired benzo[*b*]-phenanthridine **9** after hydration and oxidation. Finally an acid-catalyzed deprotection/dehydration would install the final oxazolone ring in jadomycin (Scheme 3). Following precedent, a Stille coupling could be employed to prepare **4**.^[6] In accordance with the proposed biosynthetic pathway,^[5] the L-digitoxose could be incorporated earlier (e.g., **4**) or on jadomycin A—this would depend on the stability of the glycosidic bond to the required acid-catalyzed transformations.



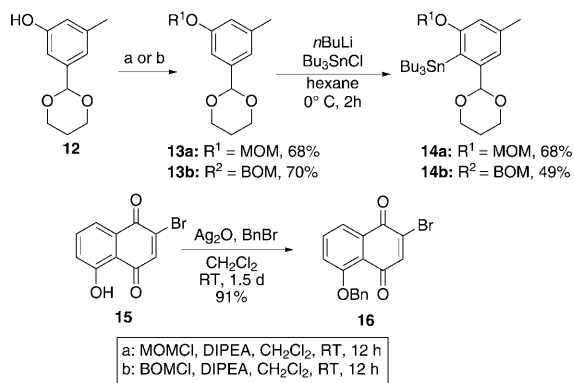
Scheme 3. Our biomimetic approach toward the jadomyins.

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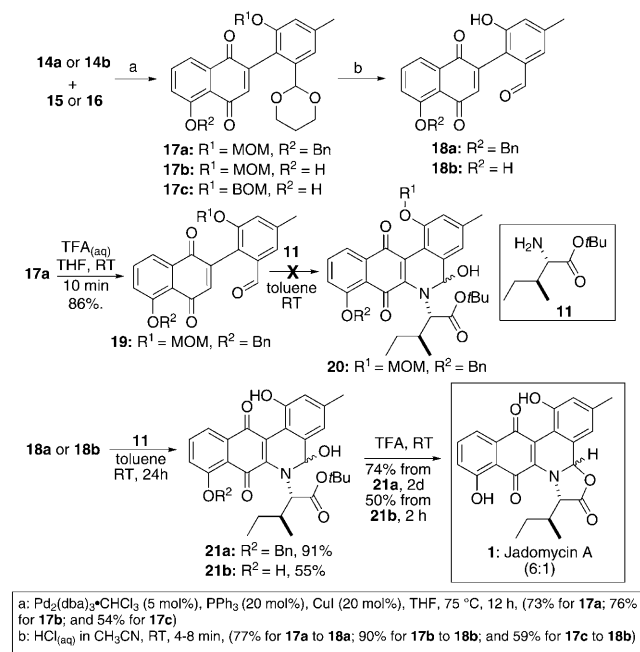
Supporting information for this article is available on the WWW under <http://dx.doi.org/10.1002/ange.201005329>.

Our synthesis began with commercially available phenol **12**,^[6] which was then protected as MOM and BOM ethers **13a** (68%) and **13b** (70%). *ortho*-Metalation of **13a/b** gave stannanes **14a** (68%) and **14b** (49%) in modest yields along with some recovered starting material (ca. 30%). The other coupling partner **16** was prepared by benzylation of commercially available bromojuglone **15** (Scheme 4).^[10]



Scheme 4. Synthesis of the desired coupling partners: MOM = methoxymethyl, BOM = benzyloxymethyl, DIPEA = *N,N*-diisopropylethylamine.

Coupling between stannane **14a** and juglone **16** gave **17a** in 73% yield (5 mol% [Pd₂(dba)₃]-CHCl₃, 20 mol% PPh₃ and 20 mol% CuI co-catalyst in THF, 75 °C for 12 h) (Scheme 5). Under identical conditions, unprotected juglone **15** was coupled with **14a** and **14b** to similar yields of **17b** (76%) and **17c** (54%). Selective acetal cleavage on **17a** (TFA_(aq)/THF, 10 min, 86%) gave the desired cyclization precursor

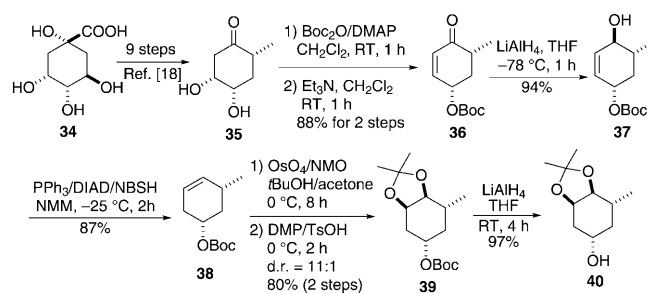


Scheme 7. Synthesis of glycosyl donors: NIS = *N*-iodosuccinimide, AIBN = azobisisobutyronitrile, DBU = 1,8-diazabicyclo[5.4.0]undec-7-ene.

was used in situ. Simply switching the cyclic carbonate protecting group of **30** to an acetonide and debenzoylation gave another potential glycosyl donor **33** (Scheme 7).^[16]

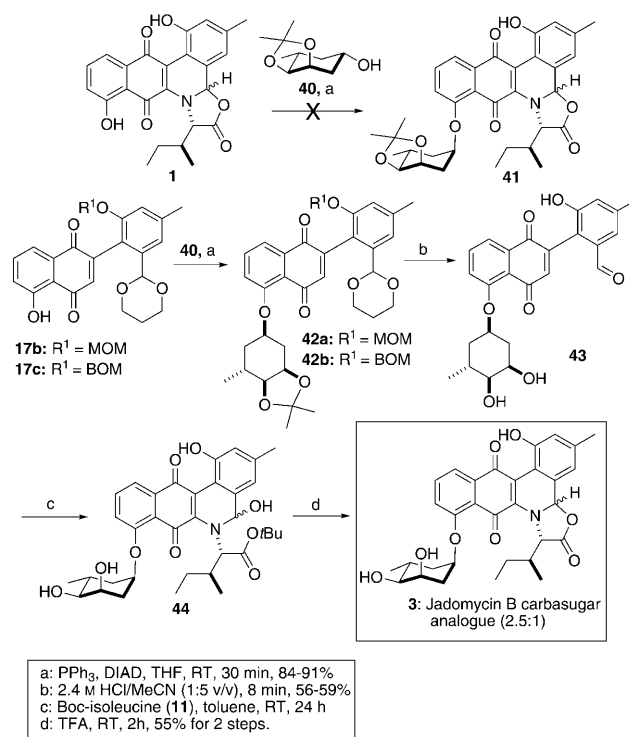
We primarily focused on the installation of the L-digitoxose onto the aglycon employing mild Mitsunobu conditions to construct jadomycin B (e.g., **31** or **33** and **17b/c** or **1** with DIAD/PPH₃, -78°C to RT).^[17] While these reaction conditions had TLC (thin-layer chromatography) evidence consistent with the formation of the desired glycosidic bond (a distinctive wine-red spot),^[18] unfortunately, none of the desired products could be isolated from these reaction mixtures. Changing of solvent and/or order of addition did not seem to have any significant effect, whereas, raising the temperature above -20°C caused the in situ formed wine-red product to disappear. Schmidt glycosylation using **32** failed to produce the desired in situ product as determined by TLC.^[15] The acid sensitivity of the glycosylated hydroxyquinones was evident by the fact that even our non-Lewis acidic Pd-catalyzed glycosylation of **1** and **17b/c** with pyranone **27** also failed. This instability of jadomycin B precluded a practical SAR type study and thus, we decided to investigate the synthesis of a less acid sensitive cyclitol analogue **3**.

The synthesis of carbasugar glycosyl donor equivalent of L-digitoxose starts from quinic acid (Scheme 8). Following a known procedure, D-(–)-quinic acid was converted into dihydroxyketone **35** in nine steps.^[19] Boc protection followed by base mediated elimination resulted in enone **36** (88 %, two steps). A diastereoselective substrate-controlled reduction of ketone **36** afforded **37** (94 %). A Myers reductive rearrangement (NBSH/DIAD/PPh₃) cleanly gave alkene **38** (87 %).^[20] Stereoselective dihydroxylation using OsO₄ and subsequent diol protection gave acetonide **39** (80 %, two steps). Finally, our proposed Mitsunobu-type cyclitolization precursor **40** was prepared by a low-temperature LiAlH₄ removal of the Boc group (97 %).



Scheme 8. Asymmetric synthesis of a cyclitol donor: Boc = *tert*-butyl carboxylate, DIAD = diisopropyl azodicarboxylate, NBSH = *o*-nitrobenzenesulfonylhydrazide, NMO = *N*-methylmorpholine *N*-oxide, DMP = 2,2-dimethoxypropane, TsOH = *p*-toluenesulfonic acid.

With the cyclitol donor **40** in hand, we employed the proposed Mitsunobu cyclitolization with jodomycin A (**1**), but unfortunately no cyclitol product **41** was observed (Scheme 9). Success occurred when we performed the Mitsunobu cyclitolization with jodomycin A (**1**) and cyclitol donor **40** in the presence of 18-crown-6 in CH₂Cl₂ at 0 °C. The cyclitol product **41** was obtained in 10% yield (Scheme 9).



Scheme 9. Synthesis of jadomycin B carbasugar analogue **3**.

nobu reaction on oxazolone precursors (e.g., **17b** and **17c**). As before these reactions' TLCs showed the characteristic wine-red spots, which this time did not disappear upon warming to room temperature. More convincingly, stable products **42a** and **42b** were isolated in excellent yields (84 % and 91 % respectively) when the reactions were run at room temperature. These acid-stable products **42a** and **42b** could be globally deprotected upon exposure to aqueous acid (2.4 M, HCl/MeCN) giving moderate yield of **43** (59 % from **42a**, 56 % from **42b**).

As before with the aglycone, the cyclitol analogues **43** could be annulated with isoleucine *tert*-butyl ester **11**. Thus, condensation of aldehyde **43** with aminoester **11** followed by a 6π -electrocyclic ring closure resulted in the formation of the desired ring system **44**. Finally acid-catalyzed transesterification leads to the formation of carbasugar analogue of jadomycin B as an inseparable mixture of 2.5:1 diastereomers (Scheme 9).

In conclusion, the first total synthesis of jadomycin A (**1**) was accomplished in six longest linear steps in 17 % overall yield. A cyclitol analogue of jadomycin B (**2**) has been successfully achieved in 20 longest linear steps in a 5 % overall yield.^[21] A 6π -electrocyclic ring closure was used to obtain the jadomycin ring system, which should be compatible with the incorporation of various other amino acids for rapid synthesis of analogues. This synthesis supports the conclusions by Rix et al. that the isoleucine incorporation and consequent formation of the oxazolophenanthridine ring system occur non-enzymatically.^[5b] The application of this methodology for further analogue synthesis and biological testing is ongoing.

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