

Communication

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J. Am. Chem. Soc., **Just Accepted Manuscript** • DOI: 10.1021/jacs.9b05370 • Publication Date (Web): 17 Jun 2019

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Synthesis of (\pm)-Idarubicinone via Global Functionalization of Tetracene

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Supporting Information Placeholder

ABSTRACT: Anthracyclines are archetypal representatives of the tetracyclic type II polyketide natural products that are widely used in cancer chemotherapy. Although the synthesis of this class of compounds has been a subject to several investigations, all known approaches are based on annulations, relying on the union of properly pre-functionalized building blocks. Herein, we describe a conceptually different approach using a polynuclear arene as a starting template, ideally requiring only functional decorations to reach the desired target molecule. Specifically, tetracene was converted to (\pm)-idarubicinone, the aglycone of the FDA approved anthracycline idarubicin, through the judicious orchestration of Co- and Ru-catalyzed arene oxidation and arenophile-mediated dearomative hydroboration. Such a global functionalization strategy, the combination of site-selective arene and dearomative functionalization, provided the key anthracycline framework in five operations and enabled rapid and controlled access to (\pm)-idarubicinone.

The *Streptomyces*-produced type II polyketides doxorubicin (1)¹ and daunorubicin (2)² are among the most effective and most often used chemotherapeutics owing to their broad-spectrum of anticancer activity (Figure 1).³ For example, doxorubicin (1) is used for the treatment of breast and bladder cancers, childhood solid tumors, soft tissue sarcomas, and aggressive lymphomas.⁴ Similarly, daunorubicin (2) is primarily used as an antileukemic drug for multiple myeloma, acute myeloid leukemia, acute lymphocytic leukemia, and Kaposi's sarcoma.⁵ Although extremely effective, anthracyclines threaten patients with cumulative dose-dependent cardiotoxicity, severely limiting their long-term application as well as their use in patients with pre-existing cardiovascular risk.⁶ Therefore, significant research efforts have been devoted to the identification of derivatives with improved pharmacological properties.⁷ The successful result of one such medicinal chemistry campaign is idarubicin (3),⁸ an FDA approved anticancer agent with superior therapeutic efficacy and reduced cardiotoxicity relative to daunorubicin (2).⁹

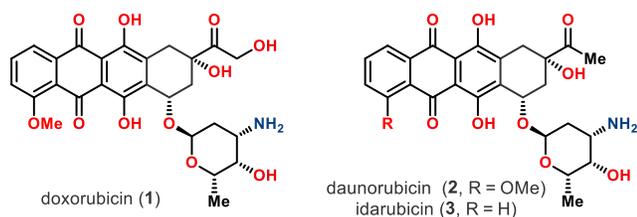


Figure 1. Structures of doxorubicin (1), idarubicin (2), and daunorubicin (3).

The need for tailored analogs has made anthracyclines the subject of rigorous investigation within the synthetic community.¹⁰ Thus, many innovative pathways to the aglycon anthracyclines (anthracyclinones) have been established, all of which rely upon annulation to forge one of the rings (see Figure 2a). The most commonly employed unifying disconnection is C-ring annulation, achieved through cycloadditions, cationic cyclizations, or anionic processes (Figure 2a, left inset).¹¹ Moreover, cycloadditions were also explored to forge other rings of the tetracyclic core of these molecules (Figure 2a, right

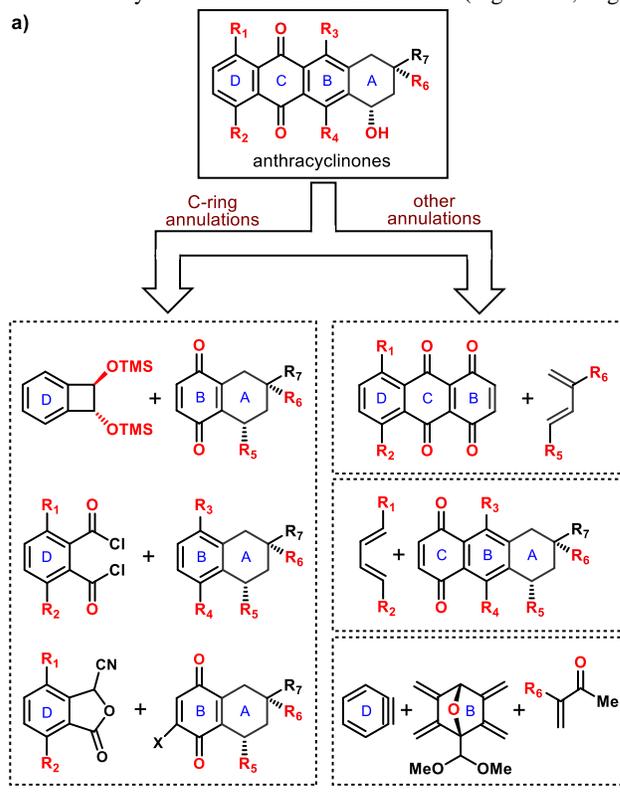


Figure 2. (a) Selected annulation-based strategies to anthracyclinones. (b) This work: synthesis of (\pm)-idarubicinone (4) from tetracene (5) using a non-annulative approach.

insets).¹² Herein, we report a conceptually different, non-annulative approach to anthracyclinones, starting from a simple aromatic hydrocarbon *via* a global functionalization strategy (Figure 2b). Specifically, (\pm)-idarubicinone (**4**) was synthesized from tetracene (**5**), an ideal aromatic precursor containing the essential tetracyclic framework, through a manifold of arene functionalizations and a site-selective dearomatative elaboration.

Following this global functionalization strategy, we commenced our studies by exploring functionalization reactions of tetracene (**5**), which would establish the proper oxidation states of the internal rings B and C within idarubicinone (**4**) (Figure 3). Thus, inspired by a similar transformation reported on anthracene, we achieved the first oxidation of **5** with catalytic amounts of cobalt(II) tetraphenylporphyrin (CoTPP, 5 mol %) and phenyliodine(III) sulfate as an oxidant,¹³ delivering 5,12-tetracenequinone (**6**) in 77% yield. Although this transformation proceeded readily, the second oxidation to the corresponding 6,11-dihydroxy-5,12-tetracenequinone derivative **7** proved more challenging. Several oxidants known for direct arene oxidation, such as CAN, Frémy's salt, hypervalent iodine reagents, or oxidizing metal complexes¹⁴ were found to be unsuitable for this transformation. This setback was not surprising, as this type of *peri*-oxidation remains a largely unsolved synthetic challenge owing to the high oxidation potential of quinones. Therefore, we decided to evaluate C–H activation, anticipating that the quinone carbonyl groups, would serve as weakly coordinating directing groups for the *peri*-(C-6) and (C-11) positions.¹⁵ After examining several carbonyl-directed hydroxylation protocols, we developed a one-pot procedure involving a modification of Ru-catalyzed *sp*² C–H oxygenation pioneered by Ackermann ([Ru(cymene)Cl₂]₂ and PIFA),¹⁶ followed by sequential one-pot hydrolysis and methylation to give desired product **7**. Control experiments revealed that this functionalization likely proceeds through the *peri*-selective formation of ruthenacycle intermediate **I-1**, delivering phenol derivative, which underwent further oxidation to the hydroquinone stage in the presence of excess PIFA.¹⁷

With arene oxidation completed, which set the required oxidation state of the B and C rings, we turned our attention to the dearomatative functionalization of the terminal ring A. We have recently reported a series of dearomatization strategies that employ visible-light-promoted *para*-cycloaddition between arenes and the arenophile *N*-methyl-1,2,4-triazoline-3,5-dione (MTAD, **8**) and subsequent *in situ* manipulation of the resulting cycloadducts.¹⁸ With polynuclear arenes, we consistently observed highly site-selective cycloadditions onto the terminal rings. Because tetracenequinone derivative **7** contains two such regions, rings A and D, amenable to cycloaddition with MTAD, another level of complexity to this process is introduced. However, based on previous studies, we know that the relevant mechanistic feature of this process is a photoinduced charge- and electron-transfer from the arene to the arenophile;¹⁹ therefore, the HOMO of the arene should dictate the site-selectivity in polynuclear aromatic settings. Accordingly, computational studies (at the B3LYP/def2-TZVPPD level of theory) of **7** predicted a strong bias for the A ring, which has profoundly larger HOMO orbital coefficients, (see Figure 3, bottom inset for the corresponding HOMO surface). Indeed, this prediction correlated well with experiment, as we observed exclusive cycloaddition onto the A-ring to provide intermediate **9**. With this site-selective dearomatization, we explored several strategies to introduce the remaining two carbon atoms needed to complete the idarubicinone framework. We found that the arenophile-based cycloaddition in combination with *in situ* Rh-catalyzed alkene hydroboration (**7**→**9**)→**10**) installed the boron moiety as a suitable handle for the introduction of the requisite acetyl group. Several hydroboration procedures were evaluated, but ultimately

the cationic rhodium complex [Rh(cod)₂BF₄] with 1,4-bis(diphenylphosphino)butane (dppb) and catecholborane provided the best outcome (for optimization details see Table S1 in Supplementary Information).²⁰ Although catecholborane was essential for the hydroboration step, the inherent instability of the resulting alkyl catechol boronic ester required immediate transesterification of catechol to pinacol to enable product isolation in higher yields.²¹ Importantly, following this protocol, we were able to prepare multigram quantities of boronic ester **10** in a single pass in 55% yield and an endo/exo 3:1 dr (see Figure 3, bottom inset for an X-ray diffraction structure of **10**).

Elaboration of organoborate **10** to the full skeleton of idarubicinone required installation of a two-carbon fragment through a seemingly straightforward *B*-alkyl Suzuki coupling reaction. However, since several standard Pd- and Ni-catalyzed reaction conditions failed,²² we decided to explore the C–C bond forming strategies involving the rich chemistry of boron 1,2-metalate rearrangements. Particularly, we were keen to explore Zweifel olefination with lithiated ethoxyvinyl ether,²³ which would provide rapid access to the C-9 acetyl group. Nevertheless, a major pitfall of this design was the presence of the quinone and its general incompatibility with organolithium reagents. Indeed, prospecting experiments involving boronic ester **10** and 1-ethoxyvinyl lithium resulted in the addition of organolithium species to quinone, delivering a mixture of products without any traces of the desired olefinated product. To address this chemoselectivity issue, we developed a one-pot process that involved *in situ* masking of the quinone. Thus, a THF solution of boronic ester **10** was sonicated with Zn powder in the presence of TMSCl, resulting in the formation of a fully protected *bis*-hydroquinone.²⁴ This intermediate was exposed to a freshly-prepared 1-lithioethyl vinyl ether to form the boronate complex **I-2**, which was immediately subjected to Zweifel olefination by addition of iodine and base.²⁵ Concurrently with olefination, the excess iodine also oxidized the labile silylated hydroquinone back to the quinone, and workup of the reaction mixture with an aqueous HCl solution hydrolyzed the newly introduced vinyl ether to the corresponding methyl ketone **11**. Remarkably, this one-pot operation involved several distinct transformations and was performed on a multigram scale in 72% yield.²⁶

Although the arenophile-mediated dearomatative hydroboration and subsequent Zweifel olefination introduced the desired methyl ketone, this sequence also installed a bridging urazole moiety, which had to be strategically transmuted to reveal the fully decorated A ring of idarubicinone (**4**). This task was partially accomplished by treatment of ketone **11** with base followed by Me₂SO₄, initiating β -elimination of urazole at position C-10 with subsequent methylation of the urazole hydrazyl nitrogen, furnishing α,β -unsaturated ketone **12** in 79% yield. The *N*-alkylation of the urazole motif proved necessary to prevent undesired side reactions during subsequent manipulations (for details see Table S2 in Supplementary Information). Finally, subjecting olefin **12** to Mukaiyama hydration conditions²⁷ selectively introduced the tertiary alcohol at position C-9, as α -ketol product **13** was obtained in 70% yield as a single diastereoisomer. Notably, the use of recently reported silane, PhSiH₂(*Oi*-Pr),²⁸ was beneficial for high conversions of this hydrogen-atom transfer process.

This hydration achieved the proper oxidation state of the A-ring, and the only difference between intermediate **13** and idarubicinone (**4**) at this stage resided in two hydroquinone protecting groups and the urazole moiety instead of a hydroxy group at C-7 position. Although deprotection of methyl ethers to hydroquinone proceeded without any difficulties using BCl₃ (**13**→**14**, 98% yield), the removal of the urazole proved to be an arduous task. Eventually, the inspiration for the direct urazole-to-

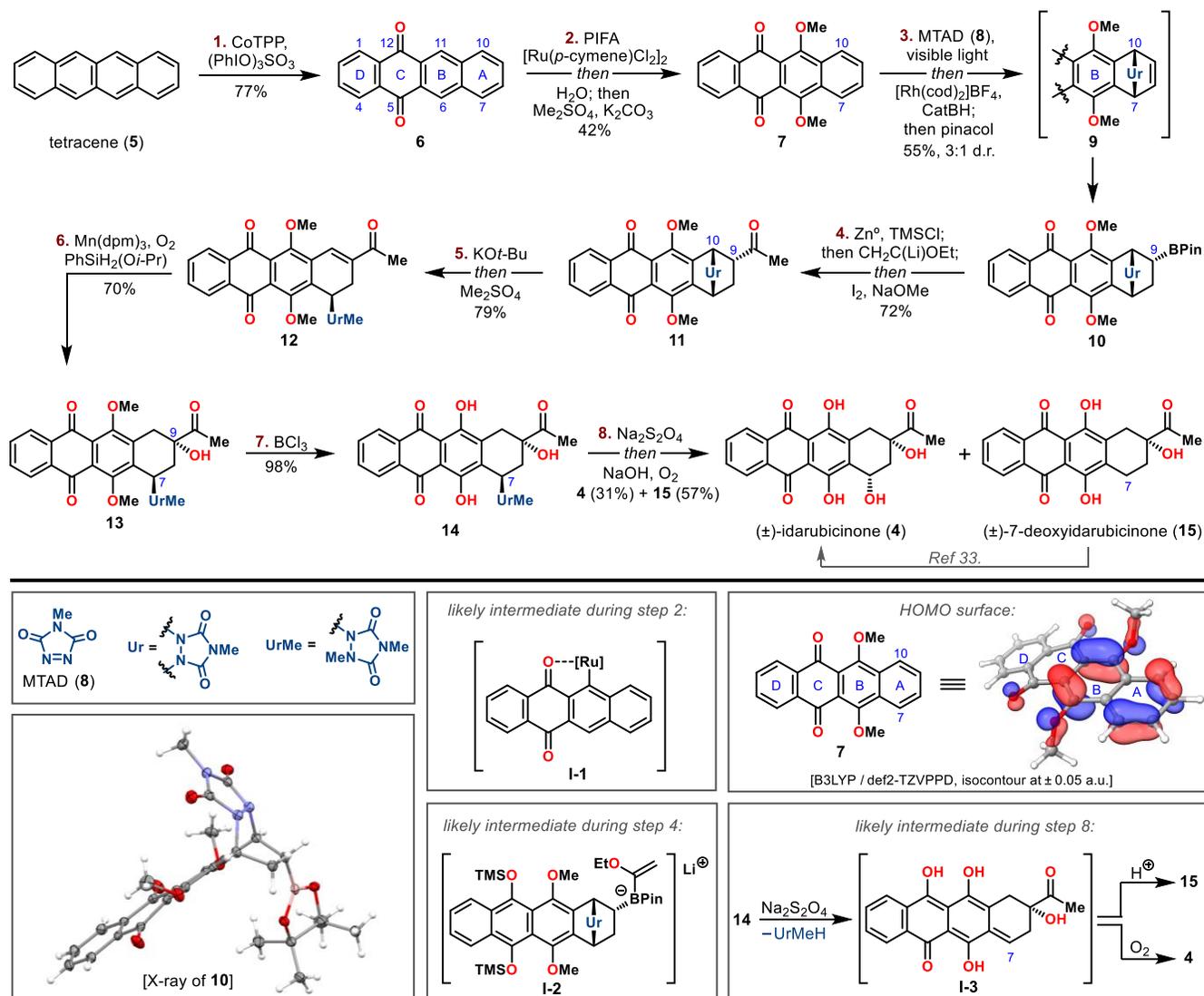


Figure 3. Synthesis of (±)-idarubicinone (**4**) from tetracene (**5**). Reagents and conditions: 1. CoTPP (5 mol%), (PhIO)₃SO₃, CH₂Cl₂, 25 °C, 2 h, 77%; 2. [Ru(*p*-cymene)Cl₂]₂ (2.5 mol%), PIFA, DCE, 100 °C, 12 h; then H₂O, 100 °C, 12 h; then Me₂SO₄, K₂CO₃, (CH₃)₂CO, 74 °C, 24 h, 42%; 3. MTAD (**8**), CH₂Cl₂, -50 °C, 36 h; then [Rh(cod)₂]BF₄ (10 mol%), dppb (10 mol%), HBCat, THF, -30 °C, 12 h; then pinacol, -78 to 25 °C, 12 h, 55% (3:1 dr); 4. Zn⁰, TMSCl, THF, ultrasonication, 40 °C, 30 min; then CH₂C(Li)OEt, -78 to -25 °C, 30 min; then I₂, -78 to -25 °C, 30 min; then NaOMe, -78 to 25 °C, 4 h; then 1 M HCl, 25 °C, 2 h, 72%; 5. KO^t-Bu, THF, -78 °C, 20 min; then Me₂SO₄, -78 to 0 °C, 1.5 h, 79%; 6. Mn(dpm)₃ (10 mol%), PhSiH₂(*Oi*-Pr), O₂, *i*-PrOH/DCE 1:1, 0 to 25 °C, 2 h, 70%; 7. BCl₃, CH₂Cl₂, -78 °C, 1 h, 98%; 8. Na₂S₂O₄, H₂O/THF/MeOH 1:1:1, -20 °C, 20 min; then NaOH, -20 °C, 60 s; then O₂, -20 °C, 5 min, 88%, **4:15** = 1:1.8.

hydroxy exchange was found in the Moore hypothesis for the biological mode of reactivity known as bioreductive alkylation.²⁹ Thus, it was proposed that anthracyclines undergo *in vivo* quinone reduction and subsequent C-7 amino sugar elimination, producing a reactive species in the form of a phenylogous quinone methide. Moreover, this concept was demonstrated in solution with several anthracyclines, which formed the corresponding semiquinone intermediates upon subjection to specific reducing agents.³⁰ The direct translation of these findings to our system, for example the addition of sodium dithionite to precursor **14**, did not eliminate the urazole; however, after the addition of base (NaOH) we observed elimination and exclusive formation of 7-deoxyidarubicinone (**15**) under anaerobic conditions. This result was in accordance with the literature, since deoxygenated anthracyclones were commonly observed upon reduction of anthracyclines.³¹ Mechanistically, the reduction of quinone **14** to hydroquinone, followed by base-induced elimination of the

urazole, likely formed the semiquinone methide **I-3**, which after protonation gave deaminated product **15**. However, we noticed that in the presence of oxygen, this reactive intermediate underwent competitive oxidation,³² delivering idarubicinone (**I-3**→**4**). Accordingly, short exposure of **14** to an aqueous solution of sodium dithionite and NaOH, followed by rapid saturation of reaction mixture with oxygen provided (±)-idarubicinone (**4**) and (±)-7-deoxyidarubicinone (**15**) in 88% yield and 1:1.8 ratio. Although extensive optimization of this protocol did not result in a higher ratio of desired anthracyclinone **4** to **15** (for details see Table S3 in Supplementary Information), this deoxygenated side-product could be readily converted to aglycone **4** in one or two steps using known protocols.³³

In summary, we have described a functionalization-based approach to (±)-idarubicinone (**4**) from tetracene (**5**). The salient feature of this strategy is a judicious orchestration of two arene functionalizations and dearomatization, introducing the

functionality of the A, B, and C rings of the anthracyclonone skeleton. Specifically, Co- and Ru-catalyzed arene oxidations, site-selective arenophile-mediated dearomative hydroboration, and subsequent Zweifel olefination provided the fully decorated anthracyclonone framework. Moreover, adjustment of the A ring, including a formally redox neutral urazole-to-hydroxy exchange delivered (\pm)-idarubicinone (**4**) in 8 operations and 2% overall yield from tetracene (**5**).

Importantly, by employing a simple polynuclear hydrocarbon aromatic starting material, the described work also presents a notable departure from previously reported syntheses of anthracyclonones in which annulations were critical to the overall synthetic design. In fact, polynuclear arenes are not commonly considered in synthetic planning for construction of stereochemically complex scaffolds. However, through the development and application of new methods, the present study provides a compelling case in which tetracene serves as an ideal template for imprinting of desired functionality. Thus, the range of available polynuclear arenes, as well as numerous functionalization opportunities, can be combined to render this global functionalization approach an appealing and complementary entry for the preparation of other type II polyketide-like compounds.

ASSOCIATED CONTENT

Supporting Information

This material is available free of charge via the Internet at <http://pubs.acs.org>.

Experimental procedures and spectral data for all new compounds (PDF)

X-ray crystallographic data.

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Notes

The authors declare no competing financial interests.

ACKNOWLEDGMENT

Financial support for this work was provided by the University of Illinois, the NIH/National Institute of General Medical Sciences (R01 GM122891), and the donors of the American Chemical Society Petroleum Research Fund (PRF#57175-DN11). D.S. is an Alfred P. Sloan Fellow. M.O. thanks the Honjo International Scholarship Foundation. Solvias AG is acknowledged for a generous gift of chiral ligands. We thank Prof. S. E. Denmark (University of Illinois) for critical proofreading of this manuscript. We also thank Dr. D. Olson and Dr. L. Zhu for NMR spectroscopic assistance, Dr. D. L. Gray and Dr. T. Woods for X-ray crystallographic analysis assistance, and F. Sun for mass spectrometric assistance.

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