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Introduction

Fredericamycin A (FDM-A) was isolated from Streptomyces griseus (FCRC-48) by Pandey and co-workers in 1981 (Fig. 1a).^{1a} It is a hexacyclic guinone-based natural product with a novel spiro [4.4] nonane ring, not previously observed in any antibiotic connected with naphthoquinone or isoquinoline aromatic moieties.1b The all-carbon quaternary chiral center in FDM-A is a unique symmetrical structure, whose chirality is enabled by a methoxy substituent in a remote position (6 atoms away from the chiral carbon center). The biosynthesis of natural FDM-A probably relies on an enzymatic asymmetric epoxidation of the highly symmetrical benastatin core and the following stereospecific transformations.² In comparison, the asymmetric chemical synthesis of this symmetrical chiral structure is much more challenging (Fig. 1b). Although several different strategies,^{3a} including the aldol reaction,^{3b,c} radical cyclization,^{3d-g} Diels-Alder reaction,^{3h,i} Tamura annulation^{3j,k} and Hauser-

^aKey Laboratory of Bioinorganic and Synthetic Chemistry of Ministry of Education, Guangdong Provincial Key Laboratory of Chiral Molecule and Drug Discovery, School of Chemistry, IGCME, Sun Yat-Sen University, Guangzhou 510275, China. E-mail: zhutshun@mail.sysu.edu.cn

Carbene-catalyzed enantioselective construction of a quasi-symmetrical spirocyclic hydroquinone with a minor chiral distinction[†]

Panlong Ren, 🔎 ^a Qing Zhao, ^a Yonggui Robin Chi ^{(b) *bc} and Tingshun Zhu ^{(b) *a}

Constructing a nearly symmetrical chiral center with tiny chiral differences is a challenging task in asymmetric synthesis. The natural antibiotic fredericamycin A (FDM-A), a representative example, has a unique structure with a quasi-symmetrical spirocyclic hydroquinone and remains difficult to chemically synthesize. Herein we developed an *N*-heterocyclic carbene–catalyzed enantioselective hydroquinone formation reaction with desymmetrization of spirocyclic cyclopentene-1,3-diones to construct these challenging structures. Using our method, the asymmetric synthesis of FDM-A (previously requiring a 26-step or 32-step synthesis) was shortened to 11 steps. Several analogs of FDM-A were also readily made. Moreover, a more challenging all-carbon quaternary chiral center with minimal differences (H *vs.* D) in a remote position (6 atoms away from the chiral center) was also constructed to investigate the performance of the extremely weakly chiral small molecule.





c) This work: carbene-catalyzed asymmetric hydroquinone formation with desymmetrization of spirocyclic



Fig. 1 Enantioselective construction of quasi-symmetrical spirocyclic hydroquinone with a minor chiral distinction.



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^bState Key Laboratory of Green Pesticide, Key Laboratory of Green Pesticide and Agricultural Bioengineering, Ministry of Education, Guizhou University, Guiyang 550025, China. E-mail: robinchi@ntu.edu.sg

^cSchool of Chemistry, Chemical Engineering, and Biotechnology, Nanyang Technological University, Singapore 637371, Singapore

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(N-methylynetoluenesulfonamide)⁶ as the coupling reagent,

triazolium salt C1 (ref. 4b) as the precatalyst, DABCO as the base

and CH₃CN as the solvent, the reaction proceeded smoothly at

room temperature and furnished the product 3a in 65% yield

with a 96:4 enantiomeric ratio (er) (Table 1, entry 1). Control

tests showed that the absence of NHC catalyst or coupling

reagent each totally deactivated the reaction (entry 2). Note the

considerable difference between catalyst chosen in our reaction

and the traditional examples for asymmetric NHC-organo-

catalysis.7 N-Aryl-substituted triazolium salts such as C2 (ref. 4a

and 8) or C3,⁹ which have been widely applied in asymmetric

NHC-organocatalysis, were totally inactive in our reaction (entry

3). *N*-Phenyl triazolium catalyst C4 (ref. 10) showed acceptable enantioselectivity but with only 8% yield (entry 4). *N*-Benzyl

triazolium salts, while easily synthesized from S_N2 alkylation of

triazole, were normally considered as unsuitable catalysts for asymmetric reactions¹¹ due to the lack of sufficient steric

hindrance. Possibly due to the relatively crowded transition

state with the phthalide-type Breslow intermediate, the reaction

showed remarkable enantioselectivity when we used the N-

benzyl triazolium catalyst. Another N-benzyl catalyst, namely

C5, also gave the desired product in good yield but with lower er

Kraus annulation^{31,m} have been applied in the total synthesis of FDM-A since the 1990s, few of them can achieve the desired enantioselective control. In 1995, Boger and coworkers obtained both enantiomers by conducting a chiral separation of racemic FDM-A, but with a low separation factor ($\alpha = 1.14$).^{3c} The only two examples of asymmetric synthesis of FDM-A were reported by the Kita group, involving a 26-step (longest linear sequence, LLS) synthesis with a chiral sulfoxide auxiliary in $1999^{3j,k}$ and a 32-step (LLS) synthesis with enzymatic esterification and enantiospecific transformations in 2005.³¹ In comparison to the racemic synthesis, tedious steps (>15 steps) were required to achieve the enantioselective control. Our research group has continuous interest in building sophisticated aromatic cycles with asymmetric benzannulation.4 Herein we developed a carbene-catalyzed asymmetric hydroguinone formation reaction to synthesize this symmetrical structure with small differences in a remote position by desymmetrization of spirocyclic dienophiles. An 11-step (LLS) synthesis of FDM-A as well as facile synthesis of FDM-A analogs were achieved. Moreover, we obtained a more challenging structure with only H/D differences in the remote position, which can hardly be identified even in the enzyme catalysis or chiral separation.

Results and discussion

Our present study commenced with the 6-chloro phthalide **1a** and spirocyclic **1**,3-cyclopentenedione **2a** (ref. 5) (spirocyclic core of FDM-A) used as model substrates (Table 1). With MYTSA



Entry	Variation of conditions	Yield ^{a,b} (%)	e.r ^c
1	None	6504	06.4
1	None	63%	96:4
2	w/o MYTsA or NHC	0	_
3	C2 or C3 instead of C1	Trace	_
4	C4 instead of C1	8%	96:4
5	C5 instead of C1	62%	75:25
6	DBU instead of DABCO	58%	93:7
7	Cs ₂ CO ₃ instead of DABCO	63%	95:5
8	CH ₂ Cl ₂ instead of CH ₃ CN	54%	91:9
9	0 °C instead of 25 °C	64%	97:3
10^d	with 100 mg 4 Å MS	73%	97:3

 a Conditions: (1) **1a** (0.2 mmol), MYTsA (0.24 mmol), CH₂Cl₂ (1 mL), 25 ° C, 0.5 h. (2) **2a** (0.1 mmol), C**1** (20 mol%), DABCO (50 mol%), CH₃CN (2 mL), 25 °C, 24 h. (3) CH₃I (0.24 mmol), K₂CO₃ (0.24 mmol), DMF (1 mL), 25 °C, 2 h. ^{*b*} Yield of isolated product. ^{*c*} Determined using chiral SFC analysis. ^{*d*} Reaction performed at 0 °C.



Scheme 1 Substrates scope ^aAll yields are isolated yields and the er values were determined from the results of chiral SFC analysis. ^bWithout methylation.

(entry 5). Reactions with other bases such as DBU (entry 6) or Cs_2CO_3 (entry 7) showed slightly lower yields and enantioselectivities. Other solvents such as CH_2Cl_2 gave inferior results (entry 8, see the ESI† for details). Performing the reaction at 0 °C instead of 25 °C resulted in a slight enhancement of enantioselectivity (entry 9). Finally, the use of 4 Å molecular sieves to maintain a more anhydrous environment gave the optimized results, affording 3a in a 73% yield with a 97:3 er (entry 10).

With the optimized conditions in hand, we next turned our attention to examine the scope of the reaction (Scheme 1). First, we evaluated the scope of 3-hydroxyl phthalides by using 2a as a model substrate. Halogenation (products 3a–3c), methylation (product 3d), methoxylation (product 3e), and phenylation (product 3f) in the 6-position of phthalides were all well tolerated, affording the desired products 3a–3f in moderate to good yields (55–73%) with excellent enantioselectivities (96:4–97:3

er). The absolute configurations of **3a** and **3c** were unambiguously confirmed from the results of single-crystal X-ray diffraction analysis. 5-Substituted phthalides with the electron-withdrawing carbonyl group (product **3g**) or electron-donating phenyloxy group (product **3h**) were also applicable substrates in our reaction, giving similar good results in both yields and enantioselectivities (51-64% yield, 96:4-97:3 er).

Using **1a** as a model phthalide substrate, the scope of the 1,3-cyclopentenediones was also evaluated. Replacing the methoxy group of **2a** with a bigger bromo group led to a slightly enhanced enantioselectivity (product **3i**, 98:2 er), while replacement with a methyl group gave product **3j** in 66% yield with 97:3 er. Introducing a *para*-methoxy substitution in **2a** gave product **3k** in 82% yield with 96:4 er. Replacing the phenyl ring of **2a** with a naphthyl unit (product **3l**) and dihydrobenzofuranyl unit (product **3m**) also afforded the desired



Scheme 2 Asymmetric synthesis of fredericamycin A and its analogs. ^aUnless otherwise specified, all the reactions were conducted without methylation. ^bSee the ESI† for details.

products with good results (71–74%, 92:8–96:4 er). The rigid spirocyclic structure and substitution maintaining the enantiofacial differences are very important for the enantioselectivity of our reaction. Removing the methoxy group in **2a** led to a sharp decline in enantioselectivity (product **3n**, 56:44 er). The more flexible 2,2-dialkyl 1,3-cyclopentenediones all gave products with reduced enantioselectivities (products **3o–3q**, 86:14–92:8 er).

Encouraged by the general good performance of our method as shown in Scheme 1, we turned our attention to the total synthesis of FDM-A. As shown in Scheme 2, the previously shortest synthesis of FDM-A, reported by Kita, relied on a Tamura annulation of the anhydride 6 and the chiral sulfoxide-auxiliary-attached dienophile 7. The anhydride 6 required a 5-step synthesis from 4a (8.5% yield), and the dienophile 7 required a 20-step synthesis from 5 (2.6% total yield).^{3j,k} In our synthesis, phthalide **1r** can be obtained with a 3-step synthesis from 4b and the yield was found to be nearly 7 times higher than that of the synthesis of 6 (58% total yield). In comparison to the synthesis of dienophile 7, from the same starting material 5, our synthesis of dienophile 2r only required one-fourth of the synthetic steps (5 steps) and gave more than 5 times the yield (14% yield). Overall, the key intermediate 3r was successfully obtained in 7 LLS steps with 4.3% total yield, that is fewer than one-third of the synthetic steps and more than twice the yield than those in the synthesis by Kita. The key intermediate 3r for the synthesis of FDM-A was also reported in the 4step synthesis in the work by Kita.^{3j,k} Notably, with dienophile 2r as a model substrate, a variety of FDM-A analogs (3s-3x) were afforded in moderate yields (42-53%) with excellent enantioselectivities (97: 3-99.5: 0.5 er). The absolute configuration of 3t was unambiguously confirmed from the results of singlecrystal X-ray diffraction analysis.

Recognizing tiny differences between prochiral substrates for the construction of symmetrical chiral centers is a challenging task in asymmetric organic synthesis.¹² To achieve an extremely symmetrical chiral structure with minimal differences, the chloro group of **1a** was replaced with a deuterium atom in our reaction, giving the interesting product **3y** in 65%



Scheme 3 Quasi-symmetrical spirocyclic structure with minor chiral distinction.

yield. As shown in Scheme 3a, the only difference to ensure the all-carbon guaternary chiral center was the difference between hydrogen and deuterium 6 atoms away from the chiral center. Using different enantiomers of the NHC catalyst, both enantiomers of 3y were obtained with similar enantioselectivities. Based on substitutions in compound 1 only slightly influencing the enantioselectivity (Scheme 1, products 3a-3h, 92%~94%) ee), 3y should be obtained in about 92% ee. However, the exact ee of 3y was not confirmed as no chiral-stationary column can separate the two enantiomers. No obvious signal was observed in the ECD spectra of the two enantiomers (Scheme 3b). Only optical rotation provided evidence to recognize the chirality of 3y. The optical rotations of 3y and its enantiomer each at various concentrations were measured, and good linear relationships were found ($R^2 > 0.99$), showing a reliable specific rotation of 0.6 (Scheme 3c). Our method thus can readily provide both enantiomers of the extremely weakly chiral compounds with minimal differences in remote position as challenging examples in chiral separation and chirality measurement.

To better understand the mechanism of the carbenecatalyzed hydroquinone formation reaction, a kinetic isotope effect (KIE) experiment was conducted and the result is shown



Scheme 4 Proposed mechanism.

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in Scheme 4a. The parallel KIE experiment revealed a secondary KIE $(k_{\rm H}/k_{\rm D} = 1.06)$, showing the breaking of the C-H bond to not be the rate-determining step (but probably a fast step). The postulated pathway of carbene-catalyzed hydroquinone formation reaction is illustrated in Scheme 4b. Briefly, the reaction starts with the formation of Breslow intermediate I via nucleophilic addition of the carbene catalyst to the aldehyde group. The following lactonization step forms the phthalidetype Breslow intermediate II (see ESI† for details of the phthalide-type Breslow intermediate),13 which has a resonance structure II'. The annulation may carry on via a stepwise Michael addition and Dieckmann condensation from intermediate II to intermediate IV (path a, Scheme 4b),¹⁴ or via a concerted [4 + 2] annulation from intermediate II' to IV (path b, Scheme 4b).¹⁵ During the annulation process, the chiral indane moiety favors the less steric hindered part and realizes the enantioselective control. The release of carbene catalyst from intermediate IV gives intermediate V and finally a rapid aromatization process with rapid C-H bond breaking gives the hydroquinone product 3a.

Conclusions

We have developed an NHC organocatalytic strategy for the enantioselective construction of spirocyclic hydroquinones bearing an all-carbon quaternary chiral center with small differences in a remote position. 3-Hydroxy phthalides were used as easily accessible starting materials and reacted with the prochiral spirocyclic dienophiles to afford the desired hydroquinone products in moderate to good yields with excellent enantioselectivities. Unlike the normal NHC organocatalytic model involving the Breslow intermediate, the annulation described in this manuscript involving the phthalide-type Breslow intermediate was found to favor the N-benzyl triazolium salts rather than the N-aryl ones. The N-benzyl triazolium salts were found to be easier to synthesize via simple alkylation of triazole, facilitating the building of a catalyst library for future investigations. With the help of this powerful method, the facile synthesis of FDM-A was achieved, shortening the synthetic route from 26 steps to 11 steps. Several analogs of FDM-A were readily synthesized as well. A more challenging quasisymmetrical spirocyclic structure with only differences of hydrogen and deuterium was also successfully synthesized to investigate the special performance of "weak" chirality.

Data availability

The data supporting this article have been included as part of the ESI.[†] Crystallographic data [for compounds **3a**, **3c** and **3t**] have been deposited at the CCDC [under CCDC 2294687 and 2394730–2394731] and can be obtained from https://www.ccdc.cam.ac.uk.

Author contributions

P. Ren conducted most of the experiments. Q. Zhao prepared the substrates for the synthesis of fredericamycin A.

Additionally, Y. R. Chi and T. Zhu conceptualized and directed the project, and drafted the manuscript with the assistance of all co-authors. All authors contributed to discussions.

Conflicts of interest

There are no conflicts to declare.

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