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Enantioselective total synthesis of (–)-colchicine, (+)-demecolcinone and metacolchicine: determination of the absolute configurations of the latter two alkaloids†

Bo Chen,^{‡a} Xin Liu,^{‡ac} Ya-Jian Hu,^a Dong-Mei Zhang,^b Lijuan Deng,^b Jieyu Lu,^a Long Min,^a Wen-Cai Ye^{ib} and Chuang-Chuang Li^{ib* a}

Here, we describe a concise, enantioselective, and scalable synthesis of (–)-colchicine (9.2% overall yield, >99% ee). Moreover, we have also achieved the first syntheses of (+)-demecolcinone and metacolchicine, and determined their absolute configurations. The challenging tricyclic 6-7-7 core of colchicinoids was efficiently introduced using an intramolecular oxidopyrylium-mediated [5 + 2] cycloaddition reaction. Notably, the synthesized colchicinoid **23** exhibited potent inhibitory activity toward the cell growth of human cancer cell lines (IC₅₀ = ~3.0 nM), and greater inhibitory activity towards microtubule assembly than colchicine, making it a promising lead in the search for novel anticancer agents.

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Introduction

Colchicine (**1**), an alkaloid natural product, was the first tubulin-destabilizing agent to be reported in the literature (Fig. 1a). The effects of this compound against a variety of indications have been investigated owing to its remarkable antimitotic activity. Furthermore, colchicine (**1**) has been used to treat several diseases, including acute gout, familial Mediterranean fever, and chronic myelocytic leukemia.¹ Colchicine (**1**) has also been used as a neurotoxin in animal models of Alzheimer's disease and epilepsy.² The development of novel colchicinoids with enhanced antitumor properties continues to pose a significant challenge to drug discovery scientists.³ From a structural perspective,⁴ the unusual 6-7-7-membered ring system of colchicinoids **1–4**, as well as the stereocenter at C-7 and the aR-configured stereogenic axis defined by the pivot bond joining the A and C rings, represents a formidable synthetic challenge. The regioselective construction of the highly oxidized tropolone C ring and the stereoselective installation of the C-7-acetamido group undoubtedly represent two of the more challenging features of any prospective colchicinoid synthesis.⁵ It is noteworthy that (+)-demecolcinone (**3**), which was the first naturally occurring dextrorotatory colchicinoid to

be reported, contains a constrained azabicyclo[3.2.1]octane (tropane) framework that is unprecedented in nature.⁶ Nevertheless, the relative configuration of naturally occurring (+)-demecolcinone (**3**) was proposed only based on the energy-minimized representation of the molecule and the postulated biosynthetic pathway, owing to the lack of direct and conclusive evidence. The absolute configuration of (+)-demecolcinone (**3**) has not been determined through X-ray crystallographic analysis or the exciton chirality circular dichroism method.

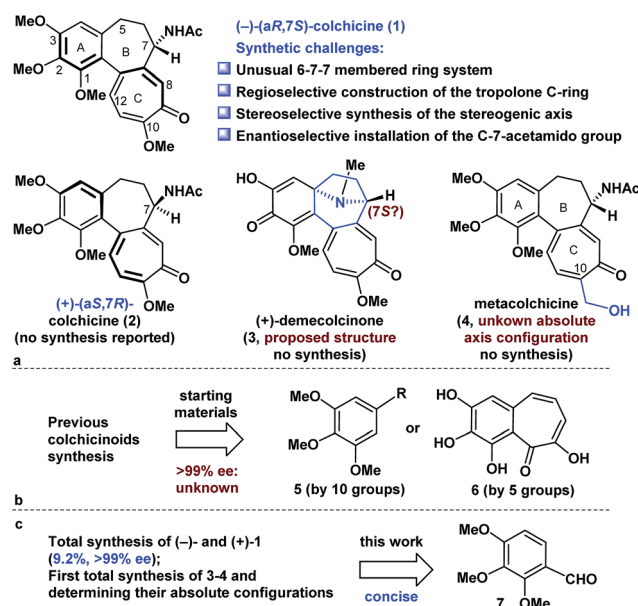


Fig. 1 Selected colchicinoids and a summary of their syntheses.

^aDepartment of Chemistry, South University of Science and Technology of China, Shenzhen 518055, China. E-mail: ccli@sustc.edu.cn^bCollege of Pharmacy, Jinan University, Guangzhou 510632, China^cInstitute of Chinese Medical Sciences, University of Macau, Macao, China

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‡ These authors contributed equally to this work.

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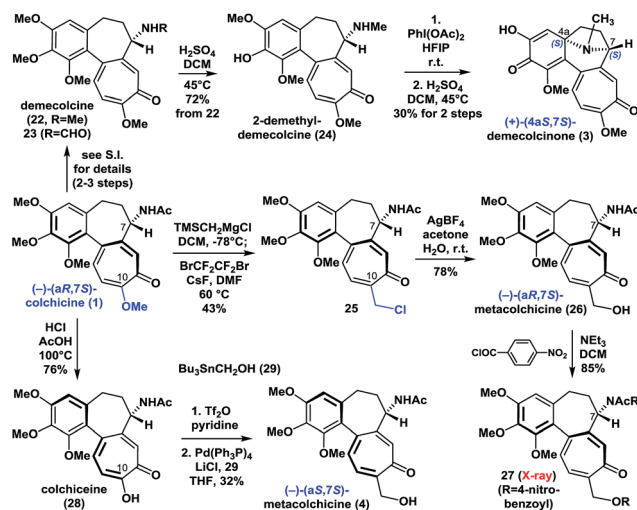
65% yield. 1,2-Addition of the organolithium reagent prepared by treatment of furfuryl alcohol (13) with BuLi to 12 provided ketone 18 in 92% yield (4.0 g scale). Subsequent condensation of ketone 18 with Ellman auxiliary 11 using $\text{Ti}(\text{OBu})_4$ (2.0 equiv.) in THF, followed by reduction of the resulting *tert*-butanesulfinyl (*t*-BS) ketimine with DIBAL-H afforded 10a (R = *t*-BS) with very high diastereoselectivity. Treatment of 10a with HCl and methanol in the same pot resulted in clean cleavage of the sulfinyl group to give the corresponding amine, which was acetylated *in situ* in the presence of NaHCO_3 to give 10 (R = Ac, >99% ee) in 78% yield (one-pot, 3.0 g scale, see the ESI for details†). Oxidative rearrangement of 10 using 3-chloroperoxybenzoic acid (*m*CPBA), followed by acetylation of the anomeric hydroxyl and intramolecular [5 + 2] cycloaddition of acetoxypyranone 9 under our optimized one-pot reaction conditions, gave tricyclic 6-7-7 core-containing 8 in 52% yield (1.0 g scale), which was confirmed by X-ray crystallography.

Next, we continued with our proposed total synthesis of colchicine (1) from 8. After extensive experimentation, we found that treatment of 8 with iodine in a mixture of pyridine and DCM, afforded α -iodoenone 19 in 88% yield. Reduction of the ketone group in 19 followed by *in situ* chemoselective methylation of the resulting alcohol afforded 20 in 76% yield (1.0 g scale). A palladium-catalyzed cross-coupling reaction of bis(pinacolato)diboron with 20 followed by oxidation under mild conditions ($\text{H}_2\text{O}_2/\text{H}_2\text{O}$) gave ketone 21 with 78% yield (1.2 g scale). Finally, double elimination of the oxa-bridge in 21 using a slightly modified version of Cha's procedure^{8c} in the presence of TMSOTf and Me_2EtN proceeded smoothly to complete our total synthesis of (–)-1 in >99% ee. We also achieved the first synthesis of (+)-(a*S*,7*R*)-colchicine (2, *ent*-1), using Ellman auxiliary *ent*-11 for condensation with ketone 18, according to a similar sequence to that shown in Scheme 1. Notably, this route provided facile access to a total of 1.1 g of (–)-1, thereby highlighting the robust nature of this chemistry.

Enantioselective synthesis of (+)-demecolcinone (3) and metacalchicine (4)

With (–)-colchicine (1) in hand, we proceeded to investigate our proposed syntheses of the remaining colchicinoids (Scheme 2). Demecolcine (22, R = Me) and 23 (R = CHO) were rapidly prepared from colchicine (1) through a series of slightly modified procedures from the literature (see ESI†). The application of a modified version of Brossi's procedure²⁰ (H_2SO_4 in DCM at 45 °C) to 22 gave 2-demethyldemecolcine (24) in 72% yield. It is worth noting that compound 22 exhibited much better chemoselectivity towards *O*-demethylation than did colchicine (1) and 23 under the same conditions.

Our initial efforts to construct the 8-azabicyclo[3.2.1]octane structure in demecolcinone (3) involved the intramolecular oxidative dearomatization–azaspiroannulation reaction of 22 using a variety of hypervalent iodine sources²¹ in several different solvents. Unfortunately, none of these reactions afforded any of the desired products. However, after extensive experimentation, we found that treatment of 24 with $\text{PhI}(\text{OAc})_2$ in the highly polar solvent hexafluoroisopropanol at 25 °C



Scheme 2 Enantioselective synthesis of (+)-demecolcinone (3) and metacalchicine (4), and determination of their absolute configurations.

followed by demethylation completed our synthesis of (+)-demecolcinone (3). The ^1H and ^{13}C NMR spectra of synthetic (+)-demecolcinone (3), as well as its optical rotation, were identical to those of the natural product. Thus, the absolute configuration of naturally occurring (+)-3, which was the first naturally occurring dextrorotatory colchicinoid to be reported in the literature,⁶ was unambiguously established to be 4*aS*,7*S* based on our total synthesis. It is noteworthy that this study represents the first reported application of an oxidative dearomatization–azaspiroannulation strategy to the synthesis of a medically significant tropane structure.²²

We then continued to investigate the synthesis of metacalchicine (4), although we believed it would be challenging to regioselectively install the hydroxymethyl group at the C-10 position of colchicine (1). After a long period of exploration, we pleasingly found that treatment of colchicine (1) with (trimethylsilyl)methylmagnesium chloride, followed by halogenation in the presence of CsF and $\text{BrCF}_2\text{CF}_2\text{Br}$ in the same pot, gave 25. This novel process involved a series of sequential reactions, including the regioselective 1,8-conjugate addition of (trimethylsilyl)methylmagnesium chloride to colchicine (1) at C10, elimination of the C10-methoxyl group, and desilylation, bromination, and chlorination (see ESI†). It is noteworthy that the chlorine atom in 25 is derived from the (trimethylsilyl)methylmagnesium chloride. Later, chloride 25 was smoothly hydrolyzed with AgBF_4 in acetone and H_2O to afford (–)-(a*R*,7*S*)-metacalchicine (26) in 78% yield. The structure of 26 was determined by 2D-NMR spectroscopy and confirmed by X-ray crystallographic analysis of its derivative 27. Surprisingly, however, the ^1H and ^{13}C NMR spectra of 26 differed from those of the natural product of metacalchicine (4).⁷ The absolute configuration of naturally occurring metacalchicine (4) has only two possibilities: a*R*,7*S* or a*S*,7*S*. Thus, the absolute configuration of naturally occurring metacalchicine (4) was determined to be a*S*,7*S* according to our synthesis. To the best of our knowledge, naturally occurring (–)-(a*S*,7*S*)-metacalchicine (4) is the first compound to be identified that has a different absolute



Table 1 Cell growth inhibitory effects of selected synthesized colchicinoids

Compounds	IC ₅₀ ^a (×±SD) μM		
	A549	MDA-MB-231	LoVo
1	0.0710 ± 0.0110	0.0332 ± 0.0310	0.0087 ± 0.0023
2	>50	>50	>50
3	>50	>50	>50
4	0.4481 ± 0.1190	0.5752 ± 0.4881	0.4030 ± 0.0647
22	0.0276 ± 0.0106	0.0310 ± 0.0047	0.0346 ± 0.0032
23	0.0028 ± 0.0009	0.0032 ± 0.0003	0.0035 ± 0.0006

^a IC₅₀ values are expressed as the mean values ± S.D. from three independent experiments.

configuration from that of other colchicine-type natural products.

Finally, the C10-methoxyl group of colchicine (**1**) was demethylated with high regioselectivity to give colchicine (**28**) in good yield. Triflation of **28**, followed by a Stille–Migita coupling reaction (Pd(PPh₃)₄, Bu₃SnCH₂OH (**29**)) furnished metacolchicine (**4**) in 32% overall yield. The ¹H and ¹³C NMR spectra and the optical rotation (synthetic: [α]_D²⁰ = −167 (*c* = 1.0, CHCl₃); natural: [α]_D²¹ = −160 (*c* = 0.39, CHCl₃)) of newly synthesized metacolchicine (**4**) were identical to those of the natural product. In this process (from **1** to **4**) clean inversion of axial stereochemistry was found, indicating that the inversion probably occurred during the conversion of **1** to **28** at 100 °C in the presence of acid. This approach could also be applied to the syntheses of structurally diverse analogues of colchicinoids bearing different groups at their C-10 positions and with different stereogenic axes. This would make it possible to carry

out structure–activity relationship (SAR) studies on these compounds.

Cell growth inhibitory activities of colchicinoids and tubule-targeting activities of **23**

The cell growth inhibitory activities of compounds **1–4** and **22–28** were determined in three human cancer cell lines, namely lung adenocarcinoma (A549), breast carcinoma (MDA-MB-231), and colon adenocarcinoma (LoVo) cells (Table 1, see ESI† for details†). Of the compounds tested in this study, compound **23** displayed the most potent inhibitory effects, with IC₅₀ values of 2.8, 3.2, and 3.5 nM against A549, MDA-MB-231, and LoVo cells, respectively.

Compound **23** was selected as a representative example to determine whether the synthesized colchicinoids acted as microtubule-targeting agents. As shown in Fig. 3a (see ESI†), **23** inhibited the polymerization of tubulin in a dose-dependent manner, thereby exhibiting similar behavior to that of colchicine (**1**). Microscale thermophoresis (MST) was used with purified recombinant tubulin to further confirm that **23** directly interfered with the assembly of tubulin monomers into microtubules. Colchicine (**1**) was also analyzed by MST as a positive control. The resulting MST measurements gave dissociation constants (*K*_d) of 0.5 ± 0.3 and 1.3 ± 0.4 μM for **23** and colchicine (**1**), respectively (Fig. 3b). Furthermore, pre-treatment of recombinant tubulin with **23** did not lead to any discernible difference in the binding affinity of colchicine (**1**) (1.9 ± 1.2 μM) for tubulin, which indicates that **23** binds to a different binding site on tubulin than does colchicine (**1**). It is noteworthy that the inhibitory activity of **23** towards the polymerization of tubulin and the binding affinity of **23** to tubulin were both more potent than those of colchicine (**1**) *in vitro*.

Microtubule-targeting drugs damage the microtubule structure of cells and induce cell cycle arrest in the G₂/M phase by disrupting the formation of the mitotic spindle required for mitosis.^{1d} We observed morphological changes in the microtubule structure of MDA-MB-231 cells after they had been exposed

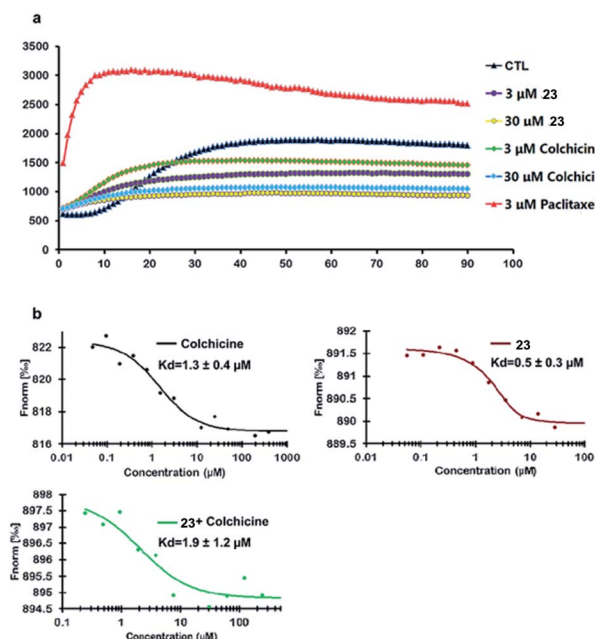


Fig. 3 Inhibition of the polymerization of tubulin by **23** and direct binding of **23** to tubulin *in vitro*.

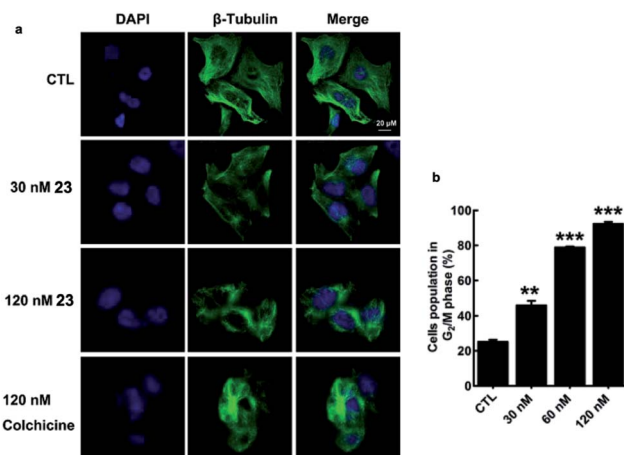


Fig. 4 Disruption of the microtubule structure and induction of G₂/M cell cycle arrest in MDA-MB-231 cells by **23**.



to **23** or colchicine (**1**) for 12 h. Treatment of MDA-MB-231 cells with **23** resulted in considerable disruption to their microtubule network with short microtubules. Similar effects were also observed for colchicine (**1**) (Fig. 4a). As shown in Fig. 4b, treatment of MDA-MB-231 cells with **23** led to a considerable dose-dependent increase in the number of cells in the G₂/M phase of the cell cycle, with values increasing from 25.19 ± 1.97% (CTL) to 46.03 ± 4.42% (30 nM), 78.85 ± 0.94% (60 nM) and 92.32 ± 2.08% (120 nM). These results therefore indicate that the activity of **23** stems from its effect on the depolymerization of the microtubules.

Conclusions

In summary, we have developed a concise and highly enantioselective synthesis of colchicine (**1**) (1.1 g, >99% ee) in nine steps²³ and 9.2% overall yield, without the need for protecting groups.²⁴ An intramolecular oxidopyrylium-mediated [5 + 2] cycloaddition reaction was used as a key step in this synthesis for efficient formation of the challenging tricyclic 6-7-7 core. In addition to achieving the most enantioselective total synthesis of colchicine (**1**) reported to date, we have also succeeded in describing the first reported asymmetric syntheses of (+)-demecolcinone (**3**) and metacolchicine (**4**), including the determination of their absolute configurations. Notably, demecolcinone (**3**), consisting of a novel and challenging azabicyclo[3.2.1]octane structure, was synthesized *via* an unusual oxidative dearomatization–azaspiroannulation reaction. It is noteworthy that an Ellman auxiliary was successfully used in the current study as a chiral directing group for the stereo- and enantioselective installation of the C7-acetamido group in a single step. Furthermore, the *in vitro* biological evaluation of compound **23** revealed that this material displayed the most potent inhibitory effects of all the compounds tested toward A549, MDA-MB-231, and LoVo cells (IC₅₀ = ~3.0 nM), as well as more potent inhibition of tubulin assembly than colchicine (**1**). These results therefore suggest that colchicinoid **23** could be used as a promising lead for the development of novel anti-cancer agents with improved properties.

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