ORGANIC CHEMISTRY







FRONTIERS

REVIEW

View Article Online View Journal | View Issue



Cite this: Org. Chem. Front., 2022, 9,

A brief overview of classical natural product drug synthesis and bioactivity

Gen Li,†a Mingliang Lou†a and Xiangbing Qi ** ** and Xiangbing Qi ** ** and Xiangbing Qi ** ** and Xiangbing Qi ** and Xiangbi

Traditional medicines consisting of compounds derived from natural organisms have been used for human health care worldwide since ancient times. Since the last century, huge numbers of bioactive natural entities with diverse chemical scaffolds have been discovered, and some have been explored as clinical medications to treat various diseases. The advent of modern technologies has promoted the discovery of natural product-based pharmaceutical agents. The synthesis of natural products not only paves the way to confirm their molecular structures but also offers the structural modification opportunity to rationally optimize the drug-likeness parameters and evaluate the bioactivity of analogs. By providing a brief overview of a miscellaneous collection of complex natural products synthesis and the efforts of the structure-activity relationship, the present report aims to highlight the impact of chemical synthesis in natural product generation, diversification, bioactivity evaluation, and natural product-based drug development.

Received 8th September 2021, Accepted 18th November 2021 DOI: 10.1039/d1qo01341f rsc.li/frontiers-organic

Introduction 1.

For millennia, nature has provided an invaluable source of traditional medicines from a variety of plants, animals, microorganisms, and minerals for humans to treat a wide spectrum of diseases. Since around 2600 BCE the successful use of natural

^aNational Institute of Biological Sciences (NIBS), 7 Science Park Road ZGC Life Science Park, Beijing 102206, China. E-mail: qixiangbing@nibs.ac.cn

medicines and remedies, particularly plant-originated substances, has been well documented in a large number of countries. The World Health Organization (WHO) estimated in 1985 that more than 4000 million people in the world relied mainly on traditional medicines for their primary health care.² Since the 19th century, developments in the fields of phytochemistry and analytical chemistry have enhanced humans' ability to examine natural materials more deeply and to identify pure bioactive constitutes to further leverage the application potential of natural products.

Beginning with the isolation of morphine, the first active principle from the opium poppy, by German pharmacist



Gen Li

Gen Li was born in Xinjiang, China. He received his B.Sc. (2004)Chemistry from in Xinjiang University and his Ph. D. (2011) degree in Chemical Biology from Peking University thesupervision under Professor Xin-Shan Ye. After a seven year research stage at Xinjiang Technical Institute of Physics and Chemistry, CAS, as an assistant professor and associate professor, he joined Qi's group at the National

Institute of Biological Science (NIBS) at the beginning of 2019. His research interests focus on the total synthesis of natural products, and medicinal chemistry.



Mingliang Lou

Mingliang Lou was born in Anhui Province, China. He received his BS degree in Biology from Nankai University in 2014. In 2020, he received his Ph.D. degree in Medicinal Chemistry from Peking Union Medical College under the direction of Professor Xiangbing Qi. Currently, he works in the National Institute of Biological Sciences (NIBS) and focuses on natural product synthesis and drug development.

^bTsinghua Institute of Multidisciplinary Biomedical Research, Tsinghua University, Beijing 100084, China

[†]These authors made equal contribution.

Fig. 1 Structures of morphine, cocaine, quinine, penicillin G, and artemisinin.

Friedrich Sertürner in 1806, and followed by its commercialization as an analgesic and sedative by Merck in 1827, several active natural products were subsequently purified, such as cocaine and quinine from plant coca and cinchona, respectively (Fig. 1).4 However, not until the last century did drug discovery-based natural products come of age due to the scientific breakthroughs in modern chemistry and medicine. The natural products pool was enriched with broader chemical structures and diverse spectrum bioactivities. Compared with conventional synthetic molecules, naturally generated products are characterized by more complex skeletons with a higher molecular mass and a larger number of sp³-hybridized



Xiangbing Qi

Xiangbing Qi was born in Shandong, China, and received his Ph.D. in chemistry and biochemistry from UT Southwestern Medical Center, Dallas, in 2005. After postdoctoral training at the University of Illinois Urbana-Champaign and medicinal chemistry research UTSouthwestern Medical Center, Dr Qi joined the faculty at the National Institute of Biological Sciences, Beijing, in 2013 and he is currently the Director of the

Chemistry Center. The central theme of the research program is focused on the interface of synthetic chemistry, chemical biology, natural product total synthesis, and medicinal chemistry.

atoms, which can be advantageous for the exploration of small molecule and biomolecule interactions.

Despite the successful discovery of multiple drugs based on original natural products, such as morphine and artemisinin (Fig. 1), they still suffer from source and availability challenges. More importantly, the low success rate of the natural entity in approved drugs emphasizes the importance of structural optimization to improve natural products' drug-like characteristics during drug development. With the development of new synthetic strategies and methodologies, more complex natural products have been assembled by efficient total synthesis or semisynthesis. The complementation of total synthesis strategies with semisynthetic transformations has been applied widely in the diversity- and bioactivity-oriented synthesis of drug-like entities.

Natural products and their synthetically modified analogs have historically made a remarkable contribution to the development of new drugs. A number of reports and books have reviewed the development of natural product-based drugs, 1,5-13 including antibiotic drugs, 14-16 antitumor agents, 17-20 analgesics, ^{21,22} and antiparasitic drugs. ^{23–25} All the reviews mentioned above emphasize that original natural products play significant roles in the drug discovery and development process. This is also confirmed further in the recent survey provided in serial reviews from Newman and Cragg, 5,26-29 which reported that only 463 purely synthetic drugs accounted for 24.6% of 1881 approved new chemical entities between 1981 and 2019. The rest of the new drugs on the market were either derived from the original natural structure or inspired by the rigidly complex skeletons of natural entities; 427 naturally occurring products (N, Newman and Cragg's category) and their structural derivatives (ND) were approved over almost 39 years.

Even though these reviews summarized nicely the natural product-based new drug development, druggability-related synthesis was not intensively tackled. We proposed to provide a brief overview of a miscellaneous collection of complex natural products synthesis and the endeavors of structure-activity relationships (SAR) studies to highlight the impact of chemical synthesis in natural product medicinal chemistry. Given the large number of approved natural product drugs in history, only the representative complex natural products whose chemical synthesis played an important role during the related drugs development are selected here. We admit that some natural products drugs, such as penicillin, vancomycin, ciclosporin, and rapamycin, are also classical and complex, but these are not covered due to limited space. Herein, the total synthesis of eight natural entities that were approved as drugs is presented according to the structural features, and the roles that total synthesis played in drug discovery and development are overviewed. The SAR studies are also discussed extensively.

Morphine and morphine derivatives

Morphine (1), isolated in 1806 by a German pharmacist, Friedrich Sertürner, is used primarily to treat both acute and

chronic moderate-to-severe pain. 30-34 It functions through selective binding to the μ -type opioid receptor in the brain and spinal cord.35,36 Many natural analogs and semisynthetic derivatives of morphine have been isolated or synthesized until now, some of which are presented in Fig. 2. The primary source of morphine and its congeners is isolation from the

> morphine (1): $R^1 = H$ oripavine (7): $R^2 = H$ codeine (6): R1 = Me thebaine (8): $R^2 = Me$ 'OH oxymorphone (10): $R^3 = H$ hydromorphone (9) oxycodone (11): R3 = Me HO naloxone (12): R4 = morphine (1) naltrexone (13): R4 =

Fig. 2 Morphine and related compounds, and their ring lettering and carbon numbering system.

opium poppy, which is greatly affected by climate and political instability. 32,37 Although the morphine biosynthesis pathway has been elucidated and the laboratory scale production of morphine in yeast has been realized, major challenges remain to be addressed before the realization of practical strategies for industrial-scale production. 38,39 Therefore, total synthesis may be a plausible surrogate for extraction in the production of morphine and its derivatives.

The structure of morphine was first proposed by Robinson in 1925, and this structure was supported by Gates' first total synthesis of morphine in 1952 and later by X-ray analysis in 1954. 40-43 As shown in Fig. 2, there are several intriguing structural features of morphine: a rigid pentacyclic bridged/fused ring structure (designated A, B, C, D, and E), five contiguous stereocenters (C5, C6, C9, C13, and C14) including one quaternary stereocenter (C13), an ether linkage between C4 and C5, two hydroxy groups (C3 and C6), a basic tertiary amine moiety, and a double bond between C7 and C8, which make its total synthesis a challenging task. The numerous studies on the synthesis of morphine alkaloids have led to over 40 synthesis routes, which have been summarized in several recent reviews. 31,33,34,37,44,45 Selected representative routes for catalytic asymmetric total synthesis of morphine alkaloids will be discussed in the following section.

Catalytic asymmetric total synthesis of morphine alkaloids

2.1.1 Overman's total synthesis of (-)-morphine. Overman and co-workers completed the first catalytic asymmetric synthesis of (-)-morphine (1).46 In their synthesis, enantioselective reduction of ketone 14 with catecholborane in the presence of (R)-oxazaborolidine catalyst (15) and subsequent condensation of the alcohol with phenyl isocyanate produced 16 (Scheme 1). The latter was transformed into 18 through two

Scheme 1 Overman's synthetic route

key operations: catalytic dihydroxylation of the terminal double bond and introduction of an allylsilane group, in which the desired C14 (following the numbering in morphine) stereocenter was established. Condensation of 18 with a tetrasubstituted A ring aldehyde 19 through the Mannich reaction generated ring D in 20. An intramolecular Heck reaction within 20 and subsequent removal of the benzyl group gave the tetracyclic compound 21 concomitant with the construction of the crucial quaternary C13 stereocenter. Ring E was constructed through epoxidation of the double bond and an epoxide ring-opening cyclization sequence. The pentacyclic compound 22 was transformed into (-)-morphine (1) following Rice's method.47

2.1.2 White's total synthesis of (+)-morphine. In White's synthesis, 48 the phenanthrene ring system (ring A/B/C) was constructed first, and this ring system was used as a platform for the sequential introduction of the dihydrofuran ring E and piperidine ring D. The first chiral center (C9) was established through Rh-catalyzed enantioselective hydrogenation of the α,β -unsaturated ester 23 in the presence of (4R,5R)-MOD-DIOP 24 (Scheme 2). The following intramolecular Friedel-Crafts reaction and Robinson annulation provided the phenanthrene nucleus 28. DBU-promoted cyclization yielded the tetracyclic compound 29, which was transformed into the diazo ketone 30. A Rh(II)-catalyzed carbenoid C-H insertion reaction was used to construct the challenging C13 all-carbon stereocenter in 31. The cyclopentanone was transformed into the piperidine ring through Beckmann rearrangement, which completed the synthesis of the pentacyclic core skeleton 32. Several functional group manipulations successfully led to the synthesis of (+)-morphine.

2.1.3 Trost's total synthesis of (-)-morphine. Trost's asymmetric total synthesis of (-)-morphine (1) features a series of Pd-catalyzed reactions. 49-51 By employing Pd-catalyzed asymmetric allylic alkylation of allylic ester 35 with 2-bromovanillin 34 in the presence of chiral ligand 36, the aryl ether 37 with 87% to 88% ee and the desired stereo configuration at C5 was obtained (Scheme 3). Then, a Pd-catalyzed intramolecular Heck reaction was applied to construct the ring E accompanied by the establishment of the quaternary stereocenter C13 (38 to 39). Another intramolecular Heck reaction on 40 smoothly led to tetracyclic compound 41, which was transformed into allyl alcohol 42. The final piperidine ring D was constructed through an intriguing visible light-promoted hydroamination reaction. It should be noted that intramolecular Heck reactions were also used in Hudlicky's asymmetric total synthesis of (-)-codeine (6) to construct ring E and ring B, wherein the chiral starting material was derived from enzyme-catalyzed enantioselective dihydroxylation β-bromoethylbenzene. 52,53

2.1.4 Tu's total synthesis of (-)-morphine. In Tu's total synthesis of (–)-morphine (1),⁵⁴ chiral spiropyrrolidine (SPD) organocatalyst 44-catalyzed asymmetric Michael addition within 43 and subsequent p-toluenesulfonic acid-promoted Robinson annulation were utilized to construct the A/C/E ring system with the establishment of the quaternary stereocenter C13 (Scheme 4). Introduction of an aldehyde group on 45 and subsequent polyphosphoric acid (PPA)-catalyzed Friedel-Crafts type cyclization led to the tetracyclic compound 46. A Wharton reaction was used to transform α,β-unsaturated ketone 46 to allylic alcohol 47. Introduction of the sulfonamide through a

Scheme 2 White's synthetic route.

Scheme 3 Trost's synthetic route.

Scheme 4 Tu's synthetic route.

regioselective Mitsunobu reaction and configuration inversion of C6 hydroxy group via an oxidation/reduction process led to 48, which was ready for the introduction of the final piperidine ring D. A lithium 4,4'-di-tert-butylbiphenylide (LiDBB)-promoted hydroamination reaction and a subsequent demethylation operation successfully led to (-)-morphine (1).

2.1.5 Dong's total synthesis of morphine alkaloids. Dong and co-workers developed a deconstructive strategy for the asymmetric total synthesis of morphine (Scheme 5).55 In their synthesis, the phenanthrene ring system in 51 was efficiently constructed through Rh-catalyzed "cutand-sew" transformation between a sterically hindered trisubstituted alkene and benzocyclobutenone within 49 in the presence of the chiral bidentate phosphine ligand 50. The chiral C13 and C14 centers with the desired stereo configuration were also set up. Then, BBr3-mediated ether bond cleavage successfully yielded compound 53, which was a suitable substrate for the introduction of the sulfonamide group in 54. The piperidine ring was constructed through sodium naphthalenide-mediated removal of the Ts group and subsequent radical cyclization. Selective demethylation on one of the two MeO ethers using NaSEt led to compound 55, which was a common precursor for the synthesis of morphine (1), codeine (6), and thebainone A.55

Scheme 5 Dong's synthetic route.

Structure-activity relationships of morphine derivatives

As the prototypical opioid agonist, morphine (1) has been intensively studied, which led to various types of morphinerelated drugs. Representative pentacyclic morphine-based drugs are listed in Fig. 3. There are several structural features that are important for morphine's biological function: the C3 phenolic hydroxy group, C6 allylic hydroxy group and C7-C8 double bond, the B/C cis-fused ring system, the substitution groups at C14, and the amine group. Detailed SAR studies have been summarized previously⁵⁶ and a brief discussion is provided in the following paragraph.

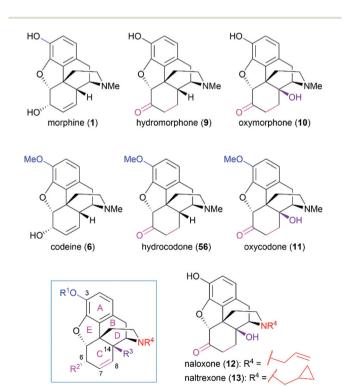


Fig. 3 Representative morphine-related drugs.

The C3 phenolic hydroxy group is essential for the binding of morphine with the μ receptor. Therefore, codeine (6), the C3 methoxy ether derivative of morphine, shows 10-fold lower potency than morphine.⁵⁷ However, codeine possesses greatly enhanced oral bioavailability. A similar relationship is observed between codeine-derivatized hydrocodone (56) and morphine-derivatized hydromorphone (9), which are the same as oxycodone (11) and oxymorphone (10).56 These codeinederivatives function as prodrugs and must undergo CYP2D6mediated demethylation of their corresponding phenolic methyl ethers before binding to μ receptors. 56

The C6 allylic hydroxy group is engaged in a weak hydrogen bond between morphine and its receptor.⁵⁸ However, removal of the hydroxy group increases the molecular lipophilicity and therefore leads to a 10-fold increase in activity. 59 If the alcohol is oxidized to ketone, a 3-fold decrease in activity is observed; further hydrogenation of the C7-C8 double bond led to hydromorphone (9) with more flexibility in ring C, which shows a 6-fold increase in potency compared with morphine.⁵⁶ Introduction of a β-hydroxy group at C14 transforms hydromorphone (9) into oxymorphone (10), which shows a 2- to 3-fold increase in receptor affinity when compared with hydromorphone (9).

The essential amino nitrogen functions in a protonated form by forming an electrostatic bond with the receptor. Substituents at this nitrogen atom have a great influence on its biological function. A methyl group is commonly observed in morphine-derived agonists; however, naloxone (12) and naltrexone (13), with allyl or cyclopropylmethyl group, respectively, are pure antagonists.

Tetracyclines

Following the discovery of chlortetracycline (57) by Duggar in 1945,⁶⁰ several other natural tetracyclines—oxytetracycline (58),⁶¹ tetracycline (59),⁶² and demeclocycline (60),⁶³—were discovered in the following years (Fig. 4). The broad-spectrum

Fig. 4 Structures of natural tetracyclines and the core structure of tetracyclines.

activity and low side effects of tetracyclines make them one of the most important classes of antibiotics. ⁶⁴ It is well established that tetracyclines function by binding to the 30S ribosomal subunit, which prevents the association of aminoacyltRNA with the bacterial ribosome and therefore inhibits bacterial protein synthesis. ⁶⁵ Over 70 years of widespread use has resulted in the emergence of bacterial resistance to tetracyclines. There are three main mechanisms responsible for tetracycline resistance: (1) efflux pumps reduce the intracellular tetracycline concentration by exporting tetracycline from the cell; (2) ribosomal protection proteins protect ribosomes from tetracycline inhibition *via* various mechanisms; and (3)

protection enzymes inactivate tetracycline through modification and degradation. 64,66-68 The growing problem of tetracycline resistance calls for much more investigation on the synthesis of tetracycline derivatives for the development of higher potency antibiotic drugs.

In the early 1950s, the structures of chlortetracycline (57) and oxytetracycline (58) were first determined by scientists from Pfizer with the help of Woodward. 69-71 Subsequent studies showed that the tetracyclines share one common nucleus, sancycline (61), which is composed of four linearly fused 6-membered rings (designated A, B, C, and D) with four stereocenters. A combination of different substitution groups, for example, chloride, hydroxy, and methyl, with the different substitution positions on the nucleus could lead to various tetracyclines with more structural complexity. The tetracyclines are frangible under an array of conditions: (1) the C6 hydroxy group undergoes dehydration in acidic media; (2) the C4 dimethylamino group epimerizes in mildly acidic conditions; and (3) the B and C rings undergo a retro-Dieckmann reaction in basic media. 72,73 The important biological functions and the complicated structural features of tetracyclines have attracted much attention of synthetic chemists.

3.1 Total synthesis of tetracyclines

3.1.1 Woodward and Shemyakin's total synthesis. One decade after they elucidated the structures of chlortetracycline (57) and oxytetracycline (58), Woodward and co-workers completed the first total synthesis of the prototypic tetracycline antibiotic, 6-demethyl-6-deoxytetracycline (61, sancycline). Starting from methyl 3-methoxybenzoate (62), they constructed the four fused rings one by one in a D to A direction through sequential Friedel–Crafts acylation and Claisen-/Dieckmann-type condensation reactions (Scheme 6). At the final stage of their synthesis, C12a was selectively oxidized with O_2 in the

Scheme 6 Woodward's total synthesis of sancycline.

Scheme 7 Shemyakin's total synthesis of tetracycline.

presence of CeCl₃ to construct the hydroxy group, which gave the desired product sancycline.

Not long after Woodward's synthesis of sancycline, Shemyakin reported their synthesis of tetracycline.⁷⁷ Starting from juglone (68), they developed a synthetic strategy similar to that of Woodward (Scheme 7). It should be noted that the C6 hydroxy group was installed using a photooxidation method.⁷⁸

3.1.2 Muxfeldt's total synthesis. In 1965, Muxfeldt and coworkers reported their total synthesis of sancycline (61).⁷⁹ Still performed in the D to A direction, their synthesis utilized 1-chloro-2-bromomethyl-4-methoxybenzene (72) as a starting material and the D/C ring fused compound 75 was synthesized (Scheme 8). Condensation of 75 with methyl N-t-butyl-3-oxoglutaramate (76) in the presence of sodium hydride led to the tetracyclic product 77. Notably, their strategy constructed the A and B rings in one step utilizing a cascade reaction consisting of intramolecular Michael addition and Dieckmann condensations. Further exploration of this strategy led to their achievement of the total synthesis of oxytetracycline (58).80

3.1.3 Stork's total synthesis. In Stork's total synthesis of tetracycline, 81 the C5a stereocenter was established using C6 hydroxy group-directed radical cyclization (78 to 79, Scheme 9), and the C4a stereocenter was correctly constructed in a substrate-controlled stereoselective Michael reaction to generate 80. Dieckmann cyclization of 80 in the presence of potassium hydride smoothly led to the pentacyclic product 81, which could be transformed into tetracycline (59) with hydrogenolytic opening of the isoxazole ring and incorporation of the C12a hydroxy group. It should be pointed out that when ring B was constructed first, trials to synthesize ring A were unsuccessful. Therefore, the authors proposed that ring A was formed prior to ring B in the transformation of 80 into 81.

3.1.4 Tatsuta's total synthesis. There was no asymmetric total synthesis of tetracyclines until Tatsuta's seminal report on the completion of (-)-tetracycline in 2000.82 Starting from 82, which was derivatized from D-glucosamine, the functionalized ring A (83) was synthesized first (Scheme 10). The subsequent regio- and stereo-selective [4 + 2] cycloaddition reaction between 83 and the diene 84 successfully led to the A/B ring-fused bicyclic intermediate. The following tandem Michael-Dieckmanntype reaction between 85 and 86 constructed the ring C with the concomitant introduction of the ring D. In contrast to previous synthetic strategies, Tatsuta's synthesis builds the tetracyclic framework in the ring A to ring D direction.

3.1.5 Myers' total synthesis. In Myers' total synthesis of the FDA-approved drug (-)-doxycycline (94), the functionalized ring B 88,83 derivatized from benzoic acid, was transformed into compound 89, which was a precursor of the tricyclic compound 90 with A/B-fused rings (Scheme 11). Like that in Stork's synthesis,81 Myers and co-workers used a functionalized isoxazole as a precursor for the sensitive groups on ring A. Following a strategy similar to that used in Tasuta's total synthesis of (-)-tetracycline, 82 the cascade Michael-Dieckmann reaction between 91 and 92 successfully yielded the pentacyclic compound 93, which could be efficiently transformed into (-)-doxycycline (94).

Scheme 8 Muxfeldt's total synthesis of sancycline

Scheme 9 Stork's total synthesis of tetracycline

Scheme 10 Tatsuta's total synthesis of (-)-tetracycline.

Scheme 11 Myers' total synthesis of (-)-doxycycline.

Previous X-ray crystallography studies on the interaction between tetracycline (59) and the 30S ribosomal subunit indicated that the D ring is suitable for modifications to generate more potent antibiotics.65 Therefore, Myers' late-stage construction of ring C with the concomitant introduction of structurally varied forms of ring D is a promising strategy to synthesize tetracycline derivatives with modified D ring. With continuing optimization, this synthetic strategy is now suitable for industrial-scale application.⁸⁴⁻⁸⁸ What's more, utilizing Myers' method, countless tetracycline derivatives, which are difficult to synthesize or are inaccessible by semisynthesis, can be constructed. Fig. 5 shows three representative tetracycline deriva-

Fig. 5 Representative tetracycline derivatives synthesized based on Myers' strategy.

tives with different substitutions, the 7-azatetracycline 95, the pentacycline 96, and the fluorocycline 97, among which 97, the first fully synthetic tetracycline antibiotic, was approved in 2018.89 To date, more than 3000 tetracycline derivatives have been synthesized based on Myers' strategy by a company named Tetraphase. 90 These tetracycline derivatives with diverse modifications make more detailed SAR studies possible.

3.2 Structure-activity relationships of tetracycline derivatives

Since the application of chlortetracycline (1) as the first broadspectrum antibiotic, extensive efforts have been devoted to the SAR studies of tetracyclines, through natural tetracyclines to semisynthetic derivatives and then to fully total synthetic candidates. The tetracycline drugs can be roughly divided into four generations and representative members are presented in Fig. 6.

Naturally isolated tetracyclines are the main members of the first-generation drugs. These natural drugs share a common pharmacophore and comprise different substitution groups at C5, C6, and C7. Several structural features are important for their bioactivities. Maintenance of the four linearly fused six-membered carbocyclic skeleton and the α stereochemical configurations of C4, C4a, C5a, and C12a are essential for their high antibacterial activity. 64,91,92 Chelation with metal ions has a great influence on tetracyclines' activity, and the keto-enol systems (C1-C3 and C11-C12) and the carboxamide (C2), which are the sites of interaction with ions, are necessary. 64,91 Alkylation of the carboxamide (C2) generally decreases activity, whereas suitable substituents can increase the water solubility, as in the case of the prodrug lymecycline. 64,92 Removal of the C4 dimethylamino or its replacement with higher alkylamino groups reduces activity.⁹¹ Furthermore, esterification of the C12a hydroxy group diminishes the activity.

However, as mentioned above, the C6 hydroxy group in these natural products is labile and undergoes dehydration in

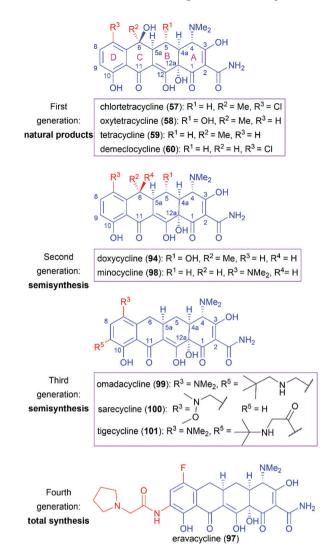


Fig. 6 Representative members of approved tetracycline drugs.

acidic media to produce anhydrotetracycline, which shows nephrotoxicity.93 Therefore, removal of the C6 hydroxy group and further modifications through semisynthesis led to two second-generation drugs, doxycycline (94) and minocycline (98), which are more stable while maintaining similar or even higher antibacterial activity. 94,95

The widespread use of tetracycline antibiotics has brought about serious resistance problems, which has led to the call for more-effective drugs. Investigations on C9-substituted tetracyclines led to the discovery of one glycylcycline named tigecycline (101), which is a derivative of minocycline (98).96 Tigecycline (101) shows high potency on tetracycline-resistant strains that utilize the efflux and ribosomal protection mechanisms of resistance.⁹⁷ However, tigecycline (101) is only available in an injectable formulation.91 Further modifications led to the discovery of omadacycline (99), which is a C9-aminomethylcycline. Compared with tigecycline (101), omadacycline (99) shows improved oral bioavailability while maintaining activity against tetracycline-resistant bacteria. 98 Another C7aminomethylcycline, sarecycline (100), was approved as a narrow-spectrum tetracycline for the treatment of acne.

The success of the second- and third-generation tetracycline drugs indicates the potential role of the tetracycline pharmacophore in developing new antibiotics. However, the use of semisynthesis, as opposed to total synthesis, for the modification of tetracyclines has inherent limitations. The power of total synthesis is demonstrated by the approval of eravacycline (97), 89,99 the first fully synthetic tetracycline drug. We believe that total synthesis will play a much more important role in the campaign against growing antibiotic resistance.

4. Macrolides

The macrolides, a series of macrocyclic lactones with 14-, 15and 16-membered rings and decorated with one or two sugar moieties are primarily isolated from different Streptomyces spp. The first macrolide and wide-spread antimicrobial agent, erythromycin (102), was isolated in 1952 from the culture broth of Saccharopolyspora erythera, and its structure features, a highly substituted 14-membered lactone bearing 10 chiral carbon atoms, and configuration were successively defined by chemical and X-ray diffraction studies. 100-102 Inspired by its excellent and broad-spectrum activity against Gram-positive pathogens, the mechanism by which erythromycin binds to the 23S RNA of the 50S ribosomal subunit to hamper the exit peptide tunnel and inhibit protein synthesis was elucidated and confirmed further by determining the crystal structure of erythromycin bound to the 50S ribosomal subunit. 103,104 Since erythromycin was launched as Ilosone to treat bacterial infections of the respiratory tract, skin, and soft tissues, the problem of varying levels of resistance to macrolides has been uncovered; resistance was characterized as resulting from the expression of efflux proteins in Gram-positive pathogens, 105,106 a change in erythromycin binding site in mutated ribosomal proteins, 107 and bio-modifications of macrolides by methylase^{108,109} and esterases.^{110,111} In addition, erythromycin is unstable, especially in acidic environments, due to ketal formation between the C6 and C12 hydroxy groups and the C9 ketone; this undesired side reaction alters the structural properties and leads to low safety and bioavailability. 112 To effectively address these problems and improve the drug-likeness potential of natural macrolides, total synthesis and semisynthesis strategies for the macrolides and their structural analogs have been explored to develop novel lead structures with new molecular features and mechanisms of actions.

4.1 Total synthesis of macrolides

4.1.1 Woodward's total synthesis of erythromycin A. Only three groups in the world, including Woodward and Martin's teams, have completed the total synthesis of erythromycin A and B. The first route to erythromycin A (102) was reported by Woodward and co-workers using dithiohemiacetal as starting material, which was followed by 12 conversion steps to generate a common intermediate (103). Then, 103 was converted to

intermediate 104 and aldehyde 105, followed by aldolization in the presence of mesityllithium as a base, yielding diastereomeric aldols (106), which underwent a series of reactions, including deprotection, protection, and oxidation, to generate aldehyde 107. The coupling of 107 with the enolate tert-butyl thiopropionate exclusively generated the "Cram" product with an undesired stereochemistry at C2. The desired stereochemistry at C2 was subsequently obtained in the presence of t-BuLi, to yield 108, which was transformed into a single secoacid derivative (109) decorated with 2-pyridyl-thioester as a precursor of macrocyclization. Next, 109 was subjected to Corey's method of lactonization to afford 110, and this was followed by ten steps including two innovative glycosidation manipulations using D-desosaminide 112 and L-cladinoside 114 as Königs-Knorr glycosyl donors, which finally led to the production of erythromycin A (102) in 48 steps (Scheme 12). 113-115

4.1.2 Martin's total synthesis of erythromycin B. A decade after Woodward's approach to erythromycin A synthesis was reported, the formal synthesis of erythromycin A was reported by Oishi's group and a highly efficient strategy for semisynthesis of erythromycin A from its natural aglycon was accomplished by Tatsuta using altered glycosidation strategy. In 1997, Martin completed the total synthesis of erythromycin B (116), a congener of erythromycin A lacking a hydroxy group at C12 (Scheme 13). 116 Based on known trihydroxy intermediate 118 prepared according to a reported method, synthesis of ketone 119 was achieved via interconversions of protecting groups and oxidation. This step was followed by aldolization with aldehyde 120 to generate intermediate 121. A macrocyclization precursor, seco-acid (122), was prepared from 121 with highly stereoselective aldolization as a critical conversion step. Using excellent Yamaguchi's macrolactonization, synthesis of the macrolide core skeleton (123) was achieved with a high yield. After multiple steps of protecting group transformations accompanied by two steps of glycosidation using Woodward's protocol, erythromycin B (116) was obtained in 23 steps.

4.1.3 Kang's total synthesis of azithromycin. The approved drug azithromycin (124), a semisynthetic analog, is prepared from erythromycin A in four steps. Considering its clinical use as a first-line antibiotic, Kang and co-workers used a novel synthetic plan instead of using erythromycin A as the precursor to construct azithromycin (124) (Scheme 14).117 According to the strategy, the triol (125) was converted to alkanolamine intermediate (126) in 11 steps using desymmetric mono-benzoylation and epoxide chemistry as key reactions. The eastern building block (130) with a carboxylic acid, was obtained by homologation, desymmerization, crotylation, and reduction and glycosidation using the known chiral moiety (127) as starting material. The alkanolamine (126) was coupled with an aldehyde by reductive amination under hydrogenation conditions, and amino methylation was completed in the presence of formaldehyde to give seco-acid (131). These and the following steps, including Yamaguchi macrocyclization, glycosidation, and deprotection, yielded azithromycin (124) in a total of 18 steps.

Scheme 12 Woodward's synthesis of erythromycin A.

Scheme 13 Martin's synthesis of erythromycin B.

Structure-activity relationships of macrolide derivatives

Since the discovery of erythromycin as the first macrolide with broad-spectrum antibacterial activity, the development of more potential novel antibiotic candidates has been spurred, leading to the preparation and testing of hundreds of macrolide derivatives. As mentioned above, spiroketal formation between the ketone at C9 with hydroxy groups at C6 and C11 under acidic conditions contributes to the low stability of erythromycin. To avoid this unpleasant side effect, a more acidresistant and powerful derivative with a 6-O-methyl group was developed by Ōmura and co-workers in 1984; six years later

Scheme 14 Kang's total synthesis of azithromycin.

this derivative, clarithromycin (133), was approved for clinical use (Fig. 7). 118 Meanwhile, a 15-membered ring of azamacrolide was furnished through Beckmann rearrangement of erythromycin A oxime at C9. This new strategy from Lazarevski's group led to a novel chemical scaffold with excellent stability and activity; this chemical was approved as a new drug in 1988, named azithromycin (124). In addition, the C3-cladinose moiety was shown to be unnecessary for protein contact. 119 The success of the marketed drug telithromycin (134) showed that its ability to bind to the ribosome was improved by the introduction of a C11/C12 carbamate ring with an extended aryl substituted alkyl chain, and binding was also improved by oxidation of the C3 hydroxy group to give C3 ketone without sugar. 120-122 However, a more effective antibiotic is urgently needed because of the serious drug resistance problem induced by the widespread use of the antibiotic. The development of more drug-like candidates for clinical applications, such as cethromycin (135) and solithromycin (136), is currently underway.

5. Vinca alkaloids

Vinblastine (137) and vincristine (138) are characteristic members of the biologically important vinca alkaloids (Fig. 8). Along with these two alkaloids, derivatives of these alkaloids, including vindesine (139), vinorelbine (140), and vinflunine (141), have been widely applied as well-known antitumor drugs. These molecules function through binding to tubulin, which disrupts microtubule function and results in mitotic arrest and apoptosis. 123-127

Vinblastine and vincristine were isolated from the leaves of Catharanthus roseus in 1958 and 1961, 128-130 respectively, and

Fig. 7 Derivatives of erythromycin A.

Fig. 8 Representative vinca alkaloids and related derivatives used as antitumor drugs and their ring lettering and carbon numbering system.

their absolute configuration was determined by X-ray crystallography in 1965. 131 Both vinblastine and vincristine possess an upper velbanamine subunit and a lower vindoline or vindoline-derived subunit. For vinblastine, the lower vindoline subunit contains a pentacyclic skeleton (ABCDE rings), which bears six contiguous stereocenters on ring C with two quaternary chiral centers. The upper velbanamine subunit is composed of four rings, including one 9-membered ring C'. There are three stereocenters in the upper subunit, including the C16' quaternary chiral center resulted from the linkage between the two subunits. The pharmaceutical importance combined with their intriguing structural features make this kind of alkaloid attractive synthetic targets, and great endeavors from the synthetic community in the past several decades have resulted in several elegant total syntheses of vinblastine and vincristine. These strategies are discussed in the following section.

Total synthesis of vinca alkaloids

5.1.1 Potier and Kutney's total synthesis of (+)-vinblastine. Inspired by the proposed biogenetic pathway for the synthesis of vinblastine, Potier and Kutney developed a pioneering strategy for direct coupling of catharanthine (142) with vindoline which are the precursors of (144), vinblastine (Scheme 15). 132-134 In their synthetic design, catharanthine (142) was first oxidized to N-oxide 143, which underwent fragmentation to give bis-iminiumion 145 through a TFAA-promoted Polonovski reaction. Then, the coupling of 145 with vindoline (144) successfully produced iminiumion 146 with the desired stereoconfiguration at C16'. Careful examination of reaction parameters showed that the C16' diastereoselectivity was temperature- and concentration-dependent: low temperature and high concentration gave better diastereoselectivity. 132

Potier and Kutney applied different strategies for the transformation of iminiumion 146 into vinblastine (137). In Kutney's one-pot operation, a 1,4-reduction of 146 with the carefully designed reductant 147 yielded enamine 148, which was successfully oxidized with O2 in the presence of FeCl3 to establish the C20' hydroxy group. 134 Potier's 1,2-reduction of iminiumion 146 with NaBH₄ produced anhydrovinblastine 149, which is a naturally occurring vinca alkaloid. Hydrogenation of the C15'-C20' double bond and oxidation gave 150. The following Polonovski reaction transformed 150 into enamine 148. Installation of the C20' hydroxy group was achieved through Tl(OAc)3 mediated oxidation followed by reduction with NaBH₄.¹³³

5.1.2 Kuehne's total synthesis of (+)-vinblastine. Kuehne's enantioselective synthesis of vinblastine commenced with the construction of the chiral aldehyde 152 through catalytic asymmetric Sharpless epoxidation (Scheme 16). 135,136 A Pictet-Spengler reaction between 152 and indoloazepine 151 produced the bridged azepines 153, which underwent benzylation and subsequent rearrangement via proposed intermediate 154 to give tetracyclic compound 155 with 1:1 dr. It should be noted that the diastereoselectivity of this transformation could be improved to 4:1 dr when 152 was condensed with 162 which contained a chiral naphthylethyl group. Chlorination of 156 with tert-butyl hypochlorite provided imine compound 157, which coupled smoothly with vindoline (144) in the presence of AgBF₄ to give 158 with the desired stereoconfiguration at C16'. KBH₄-mediated reductive ring-opening of 158 provided tricyclic compound 159, which possessed the 9-membered ring C'.

The final D' ring of the upper unit was constructed by heating the precursor tosylate 159 in MeOH at reflux for 48 h. It is worth noting that no cyclized product was found when 159 was heated in toluene at reflux for two weeks. Hydrogenolytic debenzylation followed by removal of the TMS group led to a higher energy atropisomer of vinblastine, which was transformed into vinblastine (137) by heating at reflux in toluene. An alternative reaction sequence including the epoxide intermediate 161 can also transform 159 into vinblastine (137).

Scheme 15 Potier and Kutney's synthetic route.

5.1.3 Magnus' total synthesis of (+)-vinblastine. In Magnus' total synthesis of vinblastine (137), condensation of L-tryptophan (163) with diester ketone 164 produced the tetralactam **165** through Pictet–Spengler (Scheme 17). 137 Aldol condensation of 166 with chiral aldehyde 167 provided 168, which could be converted to 169 with the required C14' stereo configuration after dehydration, desulfurization, and hydrogenation. When 169 was treated with ClCO₂CH₂C₆H₄NO₂-p, carbamate formation followed by fragmentation led to delocalized carbocation 170, which was trapped by vindoline (144) to give the coupled product 171. Their study showed that the stereochemistry of the newly generated C16' chiral center was influenced by temperature, reaction solvent, and nucleophilicity of the aromatic partner. The best diastereoselectivity could be achieved in a ratio of 84:16 (C16' S/R) when the reaction was run in CH₃CN/H₂O (15:1) at −15 °C. Hydrolysis of 171 followed by mono-oxidation of the diol and subsequent cyclization through reductive amination provided vinblastine (137).

5.1.4 Fukuyama's total synthesis of (+)-vinblastine. Fukuyama's total synthesis of vinblastine started with the enantioselective synthesis of the lower (-)-vindoline (144) subunit (Scheme 18). 138,139 Ring-opening of 7-mesyloxyquinoline (173) with thiophosgene followed by reduction with NaBH₄ provided isothiocyanate 174, which was transformed into thioanilide 175. Following the protocol developed in their laboratory, 175 was converted to indole compound 176 through radical-mediated cyclization. Further elaboration of

176 led to 177, which was transformed into pentacyclic compound 178 upon hydrolysis in the presence of TFA followed by removal of DNS protection and subsequent cyclization. Further manipulations including dehydration to construct the C14-C15 double bond, and introduction of the C16 and C17 hydroxy groups successfully provided (-)-vindoline (144). The synthesis of the upper subunit also relied on a radicaltype indole synthesis method developed by Fukuyama and coworkers to provide epoxide 180, which was a precursor of the eleven-membered-ring compound 181. When treated with tertbutyl hypochlorite, 181 was smoothly converted to chloroindolenine 182, which was coupled with (-)-vindoline (144) in the presence of TFA to yield 183 with the desired stereoconfiguration at C16'. It is worth noting that the diastereoselectivity of this transformation was very high and no other stereoisomer was found. The final D' ring was synthesized through deprotection of the tertial alcohol, removal of the Ns protection, and subsequent cyclization. This strategy was also applied to the synthesis of (+)-vincristine and vinblastine derivatives in the Fukuyama laboratory. 140,141

5.1.5 Boger's total synthesis of (+)-vinblastine. Boger's concise synthesis of vinblastine utilized a powerful cascade cycloaddition reaction developed in their laboratory. 142,143 When heated at a high temperature, the advanced intermediate oxadiazole 184 underwent an inverse electron demand Diels-Alder reaction to give 185, which was transformed into the bridged cycloadduct 187 after the loss of N2 and subsequent 1,3-dipolar cycloaddition (Scheme 19). It should be

Scheme 16 Kuehne's synthetic route.

Scheme 17 Magnus' synthetic route.

Scheme 18 Fukuyama's synthetic route.

noted that this elegant cascade cycloaddition reaction constructed four C-C bonds and three rings concomitant with the establishment of the six stereocenters with high diastereoselectivity. Hydrogenolytic cleavage of the oxygen bridge in 188 followed by silyl ether cleavage and subsequent dehydration under Mitsunobu reaction condition successfully produced (-)-vindoline (144). The following Fe(III)-mediated coupling of (-)-vindoline (144) with catharanthine (142) and installation of C20' alcohol successfully produced vinblastine (137). According to their preliminary mechanistic studies, a plausible radical pathway was proposed for the coupling reaction: in the presence of FeCl₃, 142 was oxidized to radical cation 189, which underwent radical addition to the aromatic ring of vindoline at C10 to yield 190.144 Further oxidation for rearomatization and subsequent reduction of the iminiumion provided anhydrovinblastine (149). They also proposed another Fe(III)mediated radical pathway for the installation of a hydroxy group at C20'. 145 They applied this elegant strategy to the syntheses of vinblastine-related natural products analogs. 144,145

5.1.6 Jiang's total synthesis of (+)-vinblastine. Jiang's total synthesis of vinblastine started with inverse-electron-demand Diels-Alder cycloaddition of pyrone 191 with chiral enamide 192 to give bridged lactone 193, which constructed the C20 and C21 chiral centers (Scheme 20). 146 Further manipulations

on 193, including decarboxylation, reductive ring-opening, installation of the ester group, and oxidation, led to α,β-unsaturated ketone **194**. A challenging Fischer indolization on 194 for the introduction of the A/B rings and the C7 quaternary chiral center was optimized to give tetracyclic compound 196 with excellent diastereoselectivity and regioselectivity. Pd(0)-catalyzed installation of the ester group provided 197, which was transformed into pentacyclic compound 198 through a Pd/C-catalyzed one-pot cascade reaction including lactamization to construct ring D. Reintroduction of the C14-C15 double bond and reduction of the amide produced 199, which was an advanced intermediate in Fukuyama's total synthesis of (-)-vindoline (144). Coupling of vindoline with catharanthine (142) following Boger's protocol successfully provided vinblastine.143

5.2 Structure-activity relationships of vinblastine derivatives

Extensive SAR studies on vinblastine and related natural or synthesized derivatives led to the approval of several important antitumor drugs, some of which are listed in Fig. 8. However, the emergence of drug resistance caused by overexpression of the drug efflux pump phosphoglycoprotein (Pgp) calls for more potent vinblastine derivatives to be synthesized and evaluated in the future. Since several review papers have given a detailed summary of the SAR results, 147-150 we briefly discuss

Scheme 19 Boger's synthetic route.

Scheme 20 Jiang's synthetic route.

Fig. 9 SAR studies of vinblastine-related compounds.

research progress achieved in recent years, especially those in Boger's laboratory (Fig. 9 and Table 1).

Although substitution at the C12' position of ring A' in the upper velbanamine subunit (200 and 201) often led to reduced activity, substitution at the C10' gave promising candidates.

Table 1 Cytotoxicities of vinblastine and related derivatives to three carcinoma cell lines^a

Entry	Compound	IC_{50} (nM)		
		L1210 ^b	$\mathrm{HCT}116^b$	HCT116/VM46 ^l
1	Vinblastine (137)	6.0	6.8	6.0×10^{2}
2	200	50	65	7.3×10^{2}
3	201	3.3×10^{2}	2.2×10^{2}	5.0×10^{3}
4	202	0.70	0.80	80
5	203	6.2	7.6	7.2×10^{2}
6	204	50	50	7.8×10^{2}
7	205	6.3×10^{2}	6.2×10^{2}	$>1.0 \times 10^4$
8	206		60	6.0×10^{2}
9	207		6.0×10^{2}	$>1.0 \times 10^4$
10	208	_	90	7.5×10^{3}
11	209	40	7.5	4.4×10^{3}
12	210	_	0.82	5.3×10^{2}
13	211	5.9	2.8	80
14	212	0.51	0.60	7.5
15	213	60	70	8.3×10^{2}
16	214	6.3×10^{2}	6.7×10^{2}	7.4×10^{3}
17	215	$>1.0 \times 10^4$	$>1.0 \times 10^4$	$>1.0 \times 10^4$
18	216	6.5×10^{3}	5.8×10^{3}	$>1.0 \times 10^4$
19	217	5.7×10^{3}	7.4×10^{3}	$>1.0 \times 10^4$
20	218	55	80	9.0×10^{2}
21	219	5.6×10^{2}	80	7.8×10^{3}
22	220	7.2	6.5	4.5×10^{2}

 $[^]a$ Values given are IC $_{50}$ in nM and original activity data can be found in ref. 150–155. ^bL1210: murine leukemia cell line; HCT116: human colon cancer cell line; HCT116/VM46: resistant human colon cancer cell line, Pgp overexpression.

Applying their one-pot Fe(III)-mediated coupling and oxidation reaction, Boger and co-workers synthesized a series of vinblastine derivatives including the C10'-substituted ones. 151 These derivatives exhibited activity that correlated with the size and shape of the substituents: 10'-chlorovinblastine 203 matched the potency of vinblastine; 10'-fluorovinblastine 202 exhibited an 8-fold increase in activity against both a sensitive (HCT116) and a vinblastine-resistant tumor cell line (HCT116/VM46); in contrast, derivatives bearing the larger (SMe, 204) or rigidly extended (CN, 205) substituents were 10-100-fold less potent. According to the X-ray structure of vinblastine bound to tubulin¹²⁷ and the results of activity assays, the authors proposed that the 10'-fluorine substituent interacts with the protein at a hydrophobic site uniquely sensitive to steric interactions. 151

Initial substitution of the C20' hydroxy group with a hydrogen atom (206) or free amine (207) gave derivatives with less potency. 152,153 Acetylation of the free amine (208) resulted in activity improvement, and a derivative with the urea group (209) showed comparable potency with vinblastine. However, these derivatives exhibited a further decrease in activity against a resistant HCT116/VM46 cell line compared with vinblastine. Further substitution of the terminal nitrogen led to improved activity against both sensitive and resistant cell lines (210 and 211). Finally, they found that the isoindoline-substituted compound 212 exhibited a 10-fold increase in activity against the sensitive HCT116 cell line and an 80-fold increase in activity against the resistant HCT116/VM46 cell line.

Replacement of the C16' methyl ester with an ethyl ester (213), a cyano group (214), a primary carboxamide (215), or a hydroxy group (216) dramatically reduced the activity against both sensitive and resistant cell lines. 154 Derivatives (217-220) containing 6,6-, 5,5-, and 5,6-membered DE ring systems,

instead of the vindoline 6,5-DE ring, were synthesized. 155 Among these analogs, **220** and **218** matched the potency of vinblastine and C17-deacetoxyvinblastine, respectively. Studies also showed that the N1-methyl group, C20 ethyl group, C14–C15 double bond, and C11 methoxy group were all essential to vinblastine's cytotoxic activity. 150

6. Taxol

One of the most famous natural taxane products, taxol (221, paclitaxel) was first isolated from the stem bark of Pacific yew by Wall and Wani in 1966, and in 1971 its complex structure was elucidated by NMR techniques and X-ray crystal structural analysis as a diterpenoid with a unique 6-8-6 tricyclic carbon skeleton bearing 11 stereocenters, an unusual oxetane ring, and a β-phenylisoserine chain ester at C13 (Fig. 10). 156 Because of its potent anticancer activity, the worldwide academic and industrial research communities have developed different taxane agents for clinical use in the treatment of various cancers, including the marketed Taxol® and Anzatax®, as well as the synthetic structural derivatives under the tradenames Taxotere®, Jevtana®, and Abraxane®. In addition to its unique structure, taxol also demonstrates modes of action different from those of known anticancer agents. Mechanistic studies showed that during cancer cell mitosis, taxol diffuses through nanopores in the microtubule to bind the β-tubulin subunit on the inner surface of the microtubule, thereby inhibiting the microtubule depolymerization process and arresting the cell in the late G₂/M phase, which blocks the cell division and leads to apoptosis. Cancer cells are believed to have a higher cell division rate than "normal" cells; therefore, they are more sensitive to taxol. 157-159 As observed for other target-specific anticancer drugs, the mechanism of action discussed above and the long-term use as a first-line treatment have given rise to resistance against taxane anticancer agents. The actual resistance mechanisms are unclear, even though many of these mechanisms have been uncovered in different cell lines. For example, the over-expression of the mdr1 gene gives rise to the drug efflux of the membrane-bound P-glycoprotein. The abnormal expression of microtubule-associated proteins leads to drug resistance.

Scarcity of the natural resources of taxane entities hampers the development of taxane-derived anticancer agents, and scientists from academia and industry never stop procuring the taxane products on an industrial scale. Research groups

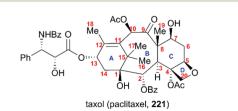


Fig. 10 The structure of taxol.

around the world have been working on the synthesis of taxanes and testing accessible synthetic strategies for the construction of the 20-carbon skeleton of the taxane family.

6.1 Total synthesis of taxol

6.1.1 Holton's total synthesis of taxol. In 1994, nearly three decades after the discovery of taxol (221), Holton's group reported the first total synthesis strategy using the readily available camphor derivative 222 as a starting material (Scheme 21). 160,161 This derivative underwent epoxidation and alcohol fragmentation followed by protection at C13 to construct the B ring of taxusin intermediate 223, which was subjected to the aldol condensation and a series of oxidations and reduction to give 224 with a ketocarbonyl at C2. With ketone 224 in hand, Chan's rearrangement in the cyclic system was applied to produce trans-fused hydroxy lactone 225 in the presence of LTMP at low temperatures. Then, multi-step transformations gave rise to intermediate 226, which was converted to the C ring-fused enol ester 227 through Dieckmann cyclization. Subsequent decarbomethoxylation and oxidation at C5 and the introduction of C4/C20 dihydroxys led to 228, which was further converted to the D ring-attached precursor 229. Additional functionality interconversions produced 7-BOM baccatin III 230, which was lithium alkoxided and treated with β-lactam 231 to successfully produce taxol (221) after deprotection of C7 BOM.

6.1.2 Nicolaou's total synthesis of taxol. Soon after Holton's report, significant work on the total synthesis of taxol from Nicolaou and co-workers was published. 162 In their strategy (Scheme 22), two pivotal intermediates, 234 and 237, with the ring A and ring B present in taxol, were both prepared by Diels-Alder cyclization between commercially available 232 and 233 and between 235 and 236, respectively; 234 and 237 were further decorated with functional groups. Then, they were subjected to Shapiro coupling to obtain a single diastereoisomer of hydroxy compound 238, which underwent epoxidation, regioselective reduction, deprotection, and oxidations to create dialdehyde 239. Subsequent optimal McMurry coupling resulted in the construction of the taxoid 6-8-6 ring skeleton 240 with 23% yield, and the configurations of the newly generated centers were confirmed by X-ray crystallographic analysis. After hydroboration-oxidation and interconversion of protecting groups, the important intermediate 241 with trihydroxy installed at C4, C5, and C20 was produced. During the next stage, the oxetane ring was built by sequential monosilylation of the primary hydroxy, triflation of the second hydroxy at C5, and acid treatment to produce a tetracyclic ring system. The allylic oxidation at C13 and benzoylation of C2-OH generated 7-TES-baccatin III 243, which was subjected to the protocol they previously reported using NaHMDS for C13 alkoxidation and consecutive treatment with β-lactam 231 to finally produce taxol (221). 163

6.1.3 Danishefsky's total synthesis of taxol. Different from the approaches previously reported, in which oxetane was constructed at a later stage, Danishefsky and co-workers prepared it at the beginning of their strategy using partially protected

Scheme 21 Total synthesis of taxol by Holton.

Scheme 22 Total synthesis of taxol by Nicolaou.

Scheme 23 Total synthesis of taxol by Danishefsky.

triol 246, which was obtained from a Wieland-Miescher ketone (Scheme 23).164 With the intermediate 247 in hand, additional functionalization led to the B ring precursor 248, which coupled with 245 derived from dione 244 to give rise to a single carbinol 249. After a series of conversions including epoxidation, hydrogenation, deprotection, and Wittig olefination, the precursor of cyclization 250 was obtained. In the next phase of synthesis, the intramolecular Heck reaction was employed to construct the tetracyclic skeleton 251, which was converted to the building block 243 used by Nicolaou through successive oxidation and protection reactions. By following previously described protocols for the final steps, the total synthesis of taxol was completed.

6.1.4 Wender's total synthesis of taxol. The synthesis strategy from the Wender team used verbenone 252, the air oxidation product of pinene, as the starting material (Scheme 24). 165,166 Reactions consisting of the major transformations of alkylation, photorearrangement, and nucleophilic addition produced the subunit tricyclic 253 as a supplier of the A-B ring system and source of chirality of the taxol core. Then, 253 was successfully converted to the A-B bicyclic ketone 254 via hydroxy-epoxide fragmentation induced by DABCO. After installation of the hydroxy at C1 and the alkyl with aldehyde group at C4, the intermediate 255 was subjected to intramolecular aldol cyclization in the presence of DMAP to construct the C ring. In the next phase, compound 256 underwent stereoselective bromination at C5 and dihydroxylation at C4/C20 to give rise to precursor 257, which directly closed to afford compound 258 with the D ring. After the protecting group interconversions, baccatin III (259) reacted with β-lactam 231 following previously reported procedures to produce the target taxol (221).

6.1.5 Kuwajima's total synthesis of taxol. In the strategy developed by Kuwajima's group (Scheme 25), 167,168 the starting materials 261 and 263, which were used as suppliers of the A ring/C2/C10 and C ring/C9, were prepared from compounds 260 and 262, respectively. In the presence of Mg(II) ions, the coupling reaction of 261 with the lithiated 263 yielded a single diastereoisomer, which was converted to the cyclization precursor 264 through the protection of the newly produced vicinal diol as a boronate. Several Lewis acids were optimized to induce cyclization, and, finally, TiCl₂(OiPr)₂ was selected as the most powerful one for the construction of the B ring. This step was followed by the removal of a protecting group, generating the tricarbocycle 265. After generation of hydroxy at C7 and ketocarbonyl at C4, cyclopropanation of the double bond between C3 and C8 was induced to give cyclopropyl ketone 266, which was exposed to the SmI₂/HMPA/methanol reductive system to smoothly produce 267 with a newly angled methyl group at C8. Functionalizations of 267 generated 268 as the precursor of oxetane, and in the presence of DBU, the tetracycle 269 was generated. In the final phase, Kuwajima and colleagues also applied previously reported protocols with some modifications to finish the total synthesis of taxol (221).

6.1.6 Mukaiyama's total synthesis of taxol. In 1999, Mukaiyama's group applied a novel approach to taxol using optically active polyoxy linear unit 272 to prepare the B ring of taxol at the early stage of the plan (Scheme 26). 169 The cyclic precursor 272 prepared from amino acid 271 by repeated intermolecular aldol reactions, was converted to the 8-membered ring smoothly in the presence of SmI2, and followed by dehydration, α,β-unsaturated cyclooctanone 273 was provided. Introduction of the alkyl aldehyde at C3 produced 274, which was converted into the desired bicyclic compound 275 by the intramolecular aldol reaction in the presence of NaOMe. After transformations of the functional groups and protections of newly formed hydroxy groups, diketone 276 was given and it subsequently was subjected to the intramolecular pinacol

Scheme 24 Total synthesis of taxol by Wender.

Scheme 25 Total synthesis of taxol by Kuwajima.

Scheme 26 Total synthesis of taxol by Mukaiyama.

coupling reaction to produce the desired pinacol 277 in the presence of low valent titanium. At the next stage, the protocols used previously were applied here to get taxol (221).

6.1.7 Baran's total synthesis of taxol. Except for the formal syntheses of taxol reported by Takahashi, 170 Nakada, 171 and Chida, 172,173 no synthetic approach was reported in the past two decades before Baran's work. The conventional convergent approaches to taxol proved not to be practical in the pharmaceutical industry; on the contrary, the relatively linear natural route used to synthesize taxol is efficient and the oxidation sequence of taxol biosynthesis could produce more natural analogs of taxol. 174,175 Inspired by the taxol biosynthetic approaches, including the generation of C-C bonds by the first type of enzymes in a "cyclase-phase" and installation of C-O bonds by the second type of enzyme in an "oxidase-phase", Baran's team presented a two-phase synthesis scheme for taxol. 176,177 As shown in Scheme 27, in the first "cyclase phase", 178 the cyclization precursor 280 was prepared on a decagram scale by the Lewis acid-modified organocopper 1,6addition of known compounds 278, 279, and a subsequent aldol reaction and oxidation. With mixture 280 in hand, the tricyclic intermediate 281 was generated in the presence of BF₃·OEt₂. At the beginning of the "oxidase phase", regioselective allylic oxidation was more feasible at C13 than at C10 in the presence of the Cr(v)-based oxidant. After bromination at C5, compound 283 experienced radical oxidation and nucleophilic substitution at C10 to generate 284, which was subjected to sequential elimination of C5 bromide, the addition of MeMgBr at C4, and reduction of the C2 and C13 ketone groups accompanied by TBS installation to give the oxidation precursor 285. The oxidant DMDO was employed to produce the epoxy-triol 286 using CHCl₃ as an optimal solvent, followed by TPAP-mediated oxidation of C2 to afford 287, which was reduced using Na/i-PrOH and exposed to triphosgene to produce carbonate 288. After sequential iodine-oxidation in the presence of TBAI/BF₃·OEt₂ and elimination and epoxidation by DMDO, 288 was converted to epoxy-taxane 289. After the reductive opening of epoxide, BOM protection, and Burgess dehydration of C4, allylic alcohol 290 was generated in moderate yield. The intermediate with a hydroxy group activated by MsCl at C5 was unstable and immediately exposed to

OsO₄ to obtain the oxetane precursor 291, which was subjected to oxetane ring formation to generate tetracyclic carbonate 292. Then, (PhSeO)₂O was applied to introduce a hydroxy group at C9 in the present of t-BuOK to deliver 293. Finally, synthesis of the target molecule 221 was achieved using previously reported protocols.

6.2 Structure-activity relationships of taxol derivatives

The discovery of taxol as one of the most potent anticancer agents spurred a large number of academic and industrial researchers to investigate the SAR for identifying lead compounds with drug-likeness potential. However, the complex structure of taxol, a heavily oxygenated polycyclic diterpenoid, presents a huge challenge for pharmaceutical studies that call for a substance-rich library. A number of studies have uncovered the still-incomplete SAR information about taxol. This information has been described by several reviews, 179-183 and is summarized here in Fig. 11.

Studies on the modifications at C1, such as removal and esterification, revealed that the C1-OH is not essential for the activity of taxol. 184 In addition, C7-OH, C9-C=O, and C10-OAc play insignificant roles in the anticancer activity of taxol. 185-188 SAR studies have shown that the benzoyl group at C2 is necessary for activity; 2-deoxy-taxol displayed decreased antitumor activity, 189 and 2-benzoate analogs did not lead to any remarkable improvement in biological activity in a microtubule assay or cytotoxicity test. 189 The acetoxy group at C4 was also demonstrated to be essential for activity, and the analog 4-deoxytaxol was inactive on several tumor cell lines. 190,191 However, oxetane plays a critical role in the bioactivity of taxol, and analogs in which the oxygen atom was replaced by nitrogen, sulfur, or other heteroatoms did not significantly enhance antitumor activity. 192,193

SAR studies revealed that the *N*-benzoyl-β-phenylisoserine side chain is essential for the antitumor activity of taxol, and this chain has attracted more extensive investigation than any other functional group of taxol. 194-198 A number of analogs with a modified side chain exhibit stronger activity than taxol, some of which are listed in Fig. 12, including two marketed drugs, docetaxel (Taxotere®, 1995) and cabazitaxel (Jevtana®, 2010).

Scheme 27 Total synthesis of taxol by Baran.

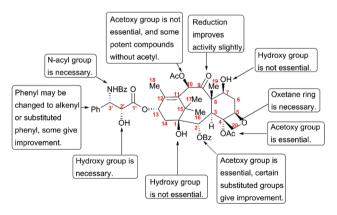


Fig. 11 Structure-activity relationships of taxol.

7. Artemisinin

(+)-Artemisinin (QingHaoSu) (Fig. 13, 5), the most used drug against malaria worldwide, was isolated from the traditional

Chinese medicine plant Artemisia annua L. in 1972 by Youyou Tu, who shared the 2015 Noble Prize for this discovery, 199 and (+)-artemisinin-based combination therapy is advocated by WHO. Artemisinin is a sesquiterpene lactone, and its structure is characterized by a unique 1,2,4-trioxane moiety, a pharmacophore that is a necessary endoperoxide bridge for the development of novel antimalaria drugs through semisynthesis of (+)-artemisinin. So far, pharmacological studies have not fully elucidated the exact mechanism of action of (+)-artemisinin, which is believed to be a pro-drug of carbon-centered free radical that plays a substantial role in antimalarial produced by cleavage of its endoperoxide ring inside erythrocytes. Upon bio-activation, the binding of (+)-artemisinin with low valent ions in hemin leads to reduction of the endoperoxide ring to produce an O radical, which subsequently generates a carbon free radical. This free radical is suggested to alkylate heme in the late stage of the parasite life cycle and react with reduced glutathione (GSH) in the early life-ring stage, after which cellular targets are selectively damaged.²⁰⁰⁻²⁰⁵ Due to its different mechanism of action, (+)-artemisinin shows higher efficacy

Fig. 12 Examples of potent taxol analogs.

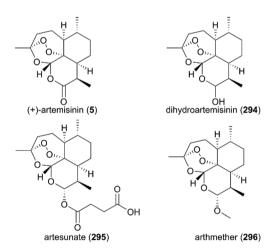


Fig. 13 (+)-Artemisinin and its derivatives.

against multi-drug-resistant malaria compared with the old antimalarial agent quinine and synthetic drug chloroquine. However, along with the widespread use of (+)-artemisininbased combination therapy in the world have come reports of resistance from different regions. The mechanism of artemisinin-resistance is unclear enough and mainly attributed to the C580Y mutation in Kelch13 (K13) of parasites, and associated

elevation of the lipid product phosphatidylinositol-3-phosphate (PI3P). It may influence host remodeling, functions of the apicoplast and food vacuole, all of which protect the parasites against artemisinin challenge. 206,207 Therefore, there is an urgent need to develop novel drugs with new chemical scaffolds and, of course, different mechanisms of action would be excellent. But (+)-artemisinin is still sourced from plants with low yield, which limits the development of more efficient drugs. Therefore, exploring chemical synthesis methods, especially for the total synthesis of (+)-artemisinin, is important for academia and the pharmacy industry.

7.1 Total synthesis of artemisinin

7.1.1 Schmid and Hofheinz's total synthesis of (+)-artemisinin. In the half-century since the discovery of (+)-artemisinin as an antimalarial drug with an unprecedented endoperoxide bridge structure, more than ten teams have completed its total synthesis. 208,209 In 1983, scientists from Roche, Schmid and Hofheinz, first reported a 13-step route to synthesize artemisinin using (-)-isopulegol (297) as a starting material (Scheme 28).210 The alkylated menthone 298 was obtained from 297 with a 6:1 diastereomeric ratio in five steps of protection, oxidation, and alkylation reactions. Following the nucleophilic addition of a substituted trimethylsilane reagent, accompanied by deprotection and oxidation, enol ether 299

Scheme 28 Total synthesis of (+)-artemisinin by Schmid and Hofheinz.

was generated after desilylation. With the key intermediate **299** in hand, photooxygenation was carried out at -78 °C, and the crude oxidation mixture was treated with acid; the target (+)-artemisinin (5) was achieved in 30% yield after crystallization.

7.1.2 Zhou's total synthesis of (+)-artemisinin. Three years later, Zhou and co-workers finished the total synthesis of (+)-artemisinin using R-(+)-citronellal (301) as raw material (Scheme 29). 211 R-(+)-citronellal (301) was subjected to two intramolecular cyclizations to produce α,β -unsaturated ketone 302, which was converted to aldehyde-ketone 303 via eight consecutive steps. Subsequently, enol ether 304 was afforded

5 steps 8 steps BnO (+)-citronellal (301) 302 3 steps O₂, hv MeO₂C MeO₂C 303 304 Н ó HCIO₄, OsO₄ o. MeO₂C (+)-artemisinin (5)

Scheme 29 Total synthesis of (+)-artemisinin by Zhou.

by protecting group manipulations and the condensation of aldehyde carbonyl with trimethyl orthoformate. Photooxidation of the methanolic solution of 304 in the presence of oxygen and Rose Bengal at -78 °C provided intermediate 305, which was hydrolyzed in 70% HClO₄ to give (+)-artemisnin (5) in 28% yield.

7.1.3 Avery's total synthesis of (+)-artemisinin. Soon after Zhou's work, Avery and co-workers reported their method for the total synthesis of (+)-artemisinin (Scheme 30).²¹² The known chiral sulfoxide 306 was used as the starting material, and five conversion steps gave rise to the diastereomerically pure silvl-acetate 307, which was converted to the cyclohexylacetate 308 by the exclusive Ireland-Claisen ester enolate rearrangement in the presence of lithium N-isopropylcyclohexylamide (LICA) as the base. Successive methylation, hydrolysis, and deketalization reactions generated the desired keto-acid 309. After exposure of the keto-acid 309 to ozone at -78 °C, the crude product was acidified with trifluoroacetic acid (TFA), and the synthesis of artemisinin was finished in a total of 12 steps. In 1992, Avery and co-workers completed the stereoselective total synthesis of 5 again, utilizing an optimized strategy with 10 steps.213

7.1.4 Ravindranathan's total synthesis of (+)-artemisinin. In 1990, Ravindranathan and co-workers utilized commercially available monoterpene (+)-car-3-ene as raw material to prepare the precursor enol ether 310, which was subjected to high temperature for 3 days to undergo an intramolecular Diels–Alder reaction, generating an epimeric mixture of ether 311 in a ratio of 3:2 (Scheme 31). After sequential oxidations, hydrolysis and methylation, the ketone-aldehyde 312 Zhou reported previously was obtained, and, followed by Zhou's protocol, the synthesis of (+)-artemisinin (5) was completed.

Scheme 30 Total synthesis of (+)-artemisinin by Avery.

Scheme 31 Total synthesis of (+)-artemisinin by Ravindranathan.

7.1.5 Liu's total synthesis of (+)-artemisinin. As shown in Scheme 32, 215 the total synthesis of (+)-artemisinin by Liu and co-workers began with a ZnCl2-catalyzed intermolecular Diels-Alder addition of (+)-enone ester 313, followed by photooxygenation in the presence of 5,10,15,20-tetraphenyl-21H,23Hporphine and oxygen to give enedione 314, which was subjected to protection, fragmentation of the cyclobutane ring, and a Wittig reaction to generate a mixture of enone ethers (315). After nine conversion steps, including deprotection, reduction, a Mitsunobu reaction, and selective hydroborationoxidation reaction, the ester 316 was obtained as a single stereoisomer. Finally, Schmid and Hofheinz's protocol for photooxygenation was applied to finish the preparation of 5.

Scheme 32 Total synthesis of (+)-artemisinin by Liu.

7.1.6 Bhonsle's total synthesis of (+)-artemisinin. In Bhonsle and co-workers' synthesis strategy (Scheme 33), 216 (-)-menthol (317) was used as the starting material to prepare 318 by Jones oxidation, α -formylation, condensation with ethyl acetoacetate, epoxidation, and regioselective reductions, along with chemoselective acetylation. With intermediate 318 in hand, photooxygenation in the presence of lead tetraacetate and I₂ was used to generate **319** as a major isomer. Next, alcohol **320** was obtained by consecutive saponification, oxidation, nucleophilic addition, and dehydration. Then, the natural product 5 was achieved using previously reported methods.

7.1.7 Constantino's total synthesis of (+)-artemisinin. In 1996, Constantino and co-workers presented the total synthesis of 5, also using (-)-isopulegol (297) as the starting material (Scheme 34). 217 In their strategy, diketone was obtained through oxidations and alkylation, and after condensation, dehydration, and some of the procedures from Bhonsle's work, acid 323 was generated and synthesis of (+)-artemisinin was achieved using a photooxygenation and

Scheme 33 Total synthesis of (+)-artemisinin by Bhonsle.

Scheme 34 Total synthesis of (+)-artemisinin by Constantino.

additional oxidation by oxygen in the presence of trifluoroacetic acid.

7.1.8 Yadav's total synthesis of (+)-artemisinin. In 2003, Yadav and co-workers reported the total synthesis of natural product 5 in 11 steps using (+)-isolimonene **324** as a raw material (Scheme 35). In their strategy, an intermolecular radical reaction on the intermediate iodolactone **325** and a Wittig reaction on a ketone **327** were critical transformations. After deprotection and photooxygenation, synthesis of (+)-artemisinin was completed using previously reported procedures.

Seven years later, Yadav and co-workers altered the synthetic strategy by using (R)-(+)-citronellal **329** as a starting material instead of (+)-isolimonene **324** (Scheme 36).²¹⁹ Their new route involved asymmetric 1,4-addition, aldol condensation, ene reaction, and regioselective hydroboration as key steps; synthesis of target molecule 5 was finished in 11 steps.

7.1.9 Cook's total synthesis of (+)-artemisinin. In 2012, Zhu and Cook reported a highly efficient route for (+)-artemisi-

Scheme 35 Total synthesis of (+)-artemisinin by Yadav.

Scheme 36 Alternative total synthesis of (+)-artemisinin by Yadav.

nin synthesis in just five pots and a total of nine steps (Scheme 37). Their strategy began with the conversion of the simple and inexpensive starting material cyclohexenone 332 to α,β -unsaturated aldehyde 333 in five steps, including a one-pot conjugate addition/alkylation sequence and formylation of the sulfonylhydrazide in the presence of n-BuLi and N,N-dimethylformamide (DMF). A [4+2] annulation was explored for the installation of the six-membered lactone to generate enol ether 334. Different from the procedure that was previously reported, they applied a singlet oxygen generated from the decomposition of H_2O_2 by ammonium molybdate to oxidize the enol ether 334 in the final step. The conditions used for the final oxidative rearrangement to 5 were suitable for large-scale production.

7.1.10 Giannis' total synthesis of (+)-artemisinin and antipode (-)-artemisinin. In 2018, Giannis and co-workers reported the total synthesis of (-)-artemisinin and (+)-artemisinin utilizing S-(+)- and R-(-)-citronellene (335) as starting materials, respectively (Scheme 38). ²²¹ After five steps, conversion of S-(+)-citronellene to a triene mixture (336) was achieved; this mixture was subjected to a thermal Diels-Alder reaction to generate artemisinic acid derivatives 337 and 338. Then, the alcohols of 337 and 338 were treated with Martin sulfurane to generate α , β -unsaturated esters, which were reduced by NiCl₂/NaBH₄ and Li/NH₃ to produce esters 339 and 340, respectively. Subsequently, a previously reported procedure was applied to convert 339 and 340 to 9-desmethyl-(-)-artemisinin, which underwent methylation to produce (-)-artemisinin (341).

Although great advances in the total synthesis of artemisinin have been made by academic researchers over the past five decades, because of the complex structure of artemisinin and its derivatives and the drawbacks of the expensive starting material, long reaction routes, and excessive protecting group operations, no synthetic strategy is economically efficient enough for pharmaceutical industry-scale production to date. Therefore, more efforts are needed to design a total synthesis strategy that can be used by industry.

Scheme 37 Total synthesis of (+)-artemisinin by Cook.

Scheme 38 Total synthesis of (-)-artemisinin by Giannis

7.2 Structure-activity relationships of artemisinin derivatives

Since the discovery of artemisinin (5) as an excellent antimalaria agent, the structural analogs such as dihydroartemisinin (294), artemether (296), arteether (342), artesunic acid (295), and sodium artesunate (343) have been investigated (Fig. 14);

Fig. 14 The first-generation artemisinin derivatives.

this is because the poor physicochemical properties of artemisinin (6), including its low solubility in both oil and water, and short half-life $(t_{1/2})$, limits its first-line clinical use. A large number of artemisinin derivatives and analogs have been explored in systematic SAR studies as part of an effort to seek more potential drug candidates. 222-229

In 2008, the first resistance to artemisinin was reported in Africa. Because of its worldwide use for a long time, WHO recommends artemisinin-based combination therapy (ACT) with different mechanisms and longer half-life instead of monotherapy using artemisinin to avoid the development and spread of resistance to artemisinin. The first-generation artemisinin-derived analogs dihydroartemisinin (294), artemether (296), arteether (342), artesunic acid (295), and sodium artesunate (343) were prepared by semisynthetic modifications at C10 with an acetal linkage; these analogs have improved physicochemical properties compared with their parent molecule, such as higher water-solubility. They are recommended by WHO to be used in combination with therapies such as ASAQ Winthrop (sodium artesunate-amodiaquine), Eurartesim (dihydroartemisinin-piperaguine), and ASMO (sodium artesunate-mefloquine) for the treatment of malaria.

However, the drawbacks of analogs with the acid-sensitive acetal linkage include a short half-life and neurotoxicity. To overcome these drawbacks, the second-generation analogs without acetal groups were designed to improve the drug-like properties; in particular compounds 347 and 348 showed better activity (Fig. 15).²³⁰

The essential role of the peroxy bridge of artemisinin, which is the pharmacophore, in the development of antimalarials was established early after its discovery, and it was confirmed by the inactive analog deoxoartemisinin 349.231 The orientation of the peroxide bond in the chemical scaffold of analogs is also important. For example, the tricyclic analog 350, which has a flexible

Fig. 15 The second-generation artemisinin derivatives.

Fig. 16 Examples of artemisinin analogs.

Fig. 17 Examples of peroxy bridge-containing artemisinin analogs.

peroxide ring displayed weaker antimalarial activity compared with the parent molecule artemisinin (Fig. 16).²³²

A number of semisynthetic analogs with tricyclic 1,2,4-trixoanes in a rigid scaffold lacking the D ring of artemisinin were developed, and many of them exhibited potential antimalarial activities in lower nanomolar ranges, such as analogs 351, 352, and 353 (Fig. 17).

Most semisynthetic drug-like analogs with complex chemical scaffolds are probably not suitable for pharmaceutical industry-scale production. The natural product yingzhausu A (354) has an accessible endoperoxide ring and displays good antimalarial activity. Spurred by this finding, a large number of simple analogs with endoperoxides have been designed and some of them display good antimalarial activity and offer great potential as drug candidates (Fig. 18). 233,234

8. **Ecteinascidins**

Ecteinascidins are a family of marine natural products containing two or three tetrahydroisoquinoline subunits (Fig. 19). Many ecteinascidin members show antiproliferative activity against several tumor cell lines. 235,236 Among these bioactive ecteinascidins, ecteinascidin 743 (Et-743, trabectedin, 357) was approved in 2007 for the treatment of advanced soft-tissue sarcoma and ovarian cancer, and lurbinectedin (363), one synthetic ecteinascidin derivative, was approved in 2020 for the treatment of metastatic small cell lung cancer. 237,238 Et-743 was first isolated in 1990 from a Caribbean tunicate Ecteinascidia turbinate and its absolute stereochemistry was determined in 1992 with the help of the X-ray crystal structure of its N12-oxide

Fig. 18 Yingzhausu A and examples of analogs.

Fig. 19 Representative ecteinascidins and ecteinascidin derivatives and their ring lettering and carbon numbering system.

derivative. 239,240 Extensive studies of Et-743 show that this molecule functions by forming covalent adducts with DNA after binding to the minor groove of DNA and alkylating DNA at the N2 position of guanine. 241-243 The adducts can bend the DNA structure toward the major groove, 244,245 disrupt the interaction between DNA and DNA-binding proteins, 246 and induce doublestrand breaks (DBSs),²⁴⁷ all of which contribute to Et-743's antiproliferative activity on various cancer cell lines.

However, the natural source of Et-743 is limited: only 1.0 g of Et-743 can be isolated from about 1.0 ton of tunicate, which

Et-637 (366): R6 = Ac, R7 = H, R8 = NHAc

hampered its preliminary clinical studies. 235,236 In 1996, Corey and co-workers reported the first synthetic route for Et-743, which was initially used for its industrial production.²⁴⁸ A semisynthetic route relying on Corey's synthetic route was developed in 2000 for the commercial manufacture of Et-743. Apart from its biological function, Et-743 also has several structural features that make it a quite challenging but attractive synthetic target: (1) three tetrahydroisoguinoline moieties with different substitution patterns; (2) eight rings including one bridged 10-membered lactone ring containing a sulfur atom; (3) seven chiral centers including one spiro center; and (4) a labile carbinolamine functional group. Continuous efforts from the synthetic community have led to outstanding achievements in the total synthesis of Et-743 and we discuss them in the following section.

8.1 Total synthesis of Et-743

8.1.1 Corey's total synthesis of Et-743. Corey and coworkers achieved the first total synthesis of Et-743 and later (in 2000) they reported a more efficient synthetic route to the key

intermediate 373 (Scheme 39). 248,250 In their synthesis strategy, the chiral C13 (Et-743 numbering) stereocenter was established through enantioselective hydrogenation of the α,β-unsaturated ester 367 to generate the tyrosine derivative 368 with 96% ee. A similar approach was applied to the synthesis of another tyrosine derivative 370, which was transformed into the bridged lactone 371 through an intramolecular Mannich reaction. Condensation between the two tyrosine derivatives 369 and 371 and subsequent manipulation of the lactone led to lactol 372. Methanesulfonic acid promoted Mannich bisannulation of 372, leading to the successful construction of a bridged C/D ring system with establishment of the C11 stereocenter. Then, a Strecker reaction was used to construct amino nitrile 373, and subsequent manipulations led to pentacyclic compound 374 with the required methyl group on ring E.

Position-selective hydroxylation with benzeneseleninic anhydride followed by the introduction of the cysteine derivative 375 generated 376, which was ready for the construction of the challenging 10-membered bridged lactone ring F. The con-

Scheme 39 Corey's synthetic route.

struction of 377 from 376 involved a series of transformations: (1) activation of the tertial hydroxy group with the in situ-generated Swern reagent from Tf2O and DMSO, followed by addition of i-Pr₂NEt, which led to o-quinone methide; (2) Barton's basemediated removal of the Fm protecting group, and the subsequent nucleophilic addition of a sulfur ion to the quinone methide generating ring F; and (3) acetylation of the resulting phenoxide group with Ac₂O. Removal of the Alloc protecting group and the subsequent transamination with 378 generated the keto lactone 379, which was utilized for the introduction of the third tetrahydroisoguinoline moiety through a Pictet-Spengler reaction with 380. Removal of the MOM protecting group from the C18 phenolic hydroxy group and stereoselective introduction of the C21 hydroxy group completed the total synthesis of Et-743.

8.1.2 Fukuyama's total synthesis of Et-743. Fukuyama and co-workers reported their first-generation synthetic route to Et-743 in 2002 and continuous efforts in their lab led to a separate synthetic strategy in 2013. 251,252 In their first-generation synthetic design,²⁵¹ the C13 chiral stereocenter in the tyrosine derivative 382 was established through enantioselective hydrogenation, while a Mannich-type reaction between 383 and

chiral template 384 stereoselectively constructed the C1 chiral center (Scheme 40). A four-component Ugi reaction including 382, 386, p-methoxyphenyl isocyanide (387) and acetaldehyde led to 388, which was transformed into the enamide compound 389 containing ring C. An intramolecular Heck reaction was applied for the formation of the bridged C/D ring system in tetracyclic compound 390, which was converted to aldehyde 391 after several manipulations. Subsequent hydrogenolytic removal of the benzyl group led to spontaneous cyclization to form the pentacyclic compound 392 with the desired stereoconfiguration at C4. Protecting group manipulations followed by the introduction of the cysteine derivative provided 393, which was utilized for the installation of the crucial ring F. Selective removal of the acetyl group gave thiol, which cyclized to form the ring F upon exposure to TFA under high dilution conditions in TFE, presumably via the o-quinone methide intermediate. Subsequent acetylation of the phenolic hydroxy group gave rise to 394, which could be converted to Et-743 in a manner similar to that utilized in Corey's total synthesis.

In their optimized second-generation synthetic route, the diacetylated diketopiperazine 395, which was derived from

Scheme 40 Fukuyama's first-generation synthetic route.

OMe

Scheme 41 Fukuyama's second-generation synthetic route.

L-glutamic acid, was the only chiral source (Scheme 41).252 Perkin condensation of 395 with aldehyde 396 followed by protection of the amide and hydrogenation stereoselectively constructed the C13 chiral center. Selective reduction of the imide carbonyl group after hydrazinolysis of the acetyl group converted 397 to a suitable intermediate for construction of the bridged C/D ring system through Mannich-type reaction, which was performed in the presence of TFA. Tetracyclic enamide 399, which was derived from 398 with the installation of the methyl group on ring E and several other manipulations, was used as a platform for the introduction of ring A. A regio- and stereo-selective Heck reaction of 399 with a diazonium salt derived from amine 400 successfully led to 401. Manipulation of 401 led to the dialdehyde equivalent 402, which cyclized to form the pentacyclic compound 403 upon heating in m-xylene. Reduction of 403 with Red-Al gave oxazolidine 404, a suitable precursor for the introduction of the cysteine derivative to give 405, an analog of 393, which could be similarly converted to Et-743.

8.1.3 Zhu's total synthesis of Et-743. Zhu's synthesis of Et-743 started with preparing the right tetrahydroisoquinoline moiety (D/E ring system) by condensation of chiral amino alcohol 406 with Garner's aldehyde (S)-407 (Scheme 42). 253 It is worth noting that the stereochemistry of the newly generated chiral center C11 was solely controlled by the absolute con-

figuration of 406. Assembly of 409 with racemic 410 gave coupled product 411 in 68% yield with its C1 epimer in 23% yield, the diastereoselectivity of which may be explained by an S_N1 mechanism. Construction of the piperazine ring C was achieved through oxidation of the hydroxy group of 412 followed by ZnCl2-catalyzed Strecker reaction to give 413. TFA-promoted cyclization of 414 concomitant with MOM protecting group removal successfully installed ring B in 415 with the desired stereoconfiguration at C4, which was similar to the ring B formation strategy utilized in Fukuyama's first-generation synthetic route. Removal of the acetate group followed by the introduction of the cysteine derivative led to 416, from which ring F was installed through a one-pot deprotection/ cyclization mediated by TFA. Compound 417 could be converted into Et-743 following Corey's protocol.

8.1.4 Danishefsky's formal total synthesis of Et-743. Starting with asymmetric reduction of the ketone moiety in 418 using Noyori transfer-hydrogenation conditions followed by a Mitsunobu reaction with diphenylphosphorylazide securely fashioned the C1 stereoconfiguration in 421 (Scheme 43).²⁵⁴ The left tetrahydroisoquinoline moiety (A/B ring) was formed from 421 through an acid-promoted Pomerantz-Fritsch reaction. Condensation of 422 with the chiral tyrosine derivative 423 afforded 424, which was converted into the cyclization precursor 425. Aldehyde 425 suc-

Scheme 42 Zhu's synthetic route.

Scheme 43 Danishefsky's synthetic route.

Scheme 44 Williams' synthetic route.

cessfully cyclized to give rise to the pentacyclic compound 426 through a CHF₂CO₂H-promoted vinylogous Pictet-Spengler reaction. DMDO-mediated epoxidation of the double bond in the advanced intermediate 427 followed by amide nitrogen atom-promoted epoxide ring-opening and subsequent reduction of the iminiumion produced compound 428 with

Scheme 45 Ma's synthetic route.

the desired stereoconfiguration at C3 and C4. It is worth noting that a large excess of sodium cyanoborohydride was required for the high yield of 428. Further manipulations on 428 led to 429, which was an intermediate in Fukuyama's total synthesis of Et-743.²⁵¹

8.1.5 Williams' formal total synthesis of Et-743. In Williams' synthetic route, 255 stereoselective condensation of the phenol 430 with (R)-Garner's aldehyde 431 in the presence of Ti (OiPr)₄ produced the anti-product 432 (Scheme 44). Condensation of the chiral amine 433 with ethyl glyoxalate led to an imine intermediate, which proceeded to form the left tetrahydroisoguinoline 434 with the desired C1 stereocenter established through an AIBN/Bu₃SnH-mediated radical cyclization. Removal of the acetonide from 436 with Dowex 50W-X8 cationic resin in methanol followed by desilylation and subsequent Swern oxidation produced 437. When treated with TFA, 437 successfully underwent a Pictet-Spengler reaction to give the pentacyclic compound 438, although with low regioselectivity on the E ring. Since several protecting group manipulations could transform 438 into 427, which was an advanced intermediate in Danishefsky's formal total synthesis, Williams and co-workers completed their formal total synthesis of Et-743.

8.1.6 Ma's total synthesis of Et-743. In 2019, Ma and coworkers reported an efficient total synthetic route to Et-743 and lurbinectedin, in which the left and right tetrahydroisoquinoline moieties were constructed from the common intermediate **439** (Scheme 45).²⁵⁶ Starting from the chiral amino alcohol **439**, which was prepared from Cbz-protected (*S*)-tyrosine, a precursor for the left tetrahydroisoquinoline moiety **440** was obtained through a Pictet–Spengler reaction. After oxidation of the

phenol 440 to quinone 441, the authors carried out detailed screenings for optimized conditions to form benzo[1,3]dioxole through a light-mediated remote C–H bond activation. They finally found that blue-LED irradiation of a THF solution of 441 could smoothly transform quinone 441 into phenol 442. It should be pointed out that this transformation was an essential step in their synthesis because the attempts to synthesize 442 from 443 through a direct Pictet–Spengler reaction under

Table 2 Cytotoxicities of Et-770 and related derivatives to three carcinoma cell lines^a

	Compound	IC ₅₀ (nM)			
Entry		HCT116 ^b	QG56 ^b	DU145 ^b	
1	Et-770 (360)	0.71	1.6	1.6	
2	Et-788 (449)	3.5×10^{2}	1.3×10^{3}	9.9×10^{2}	
3	450	54	1.6×10^{2}	1.0×10^{2}	
4	451	1.3	9.5	4.4	
5	452	0.50	1.8	1.2	
6	453	0.13	0.40	0.72	
7	454	0.014	0.071	0.087	
8	455	0.053	0.33	0.24	
9	456	0.010	0.12	0.041	
10	457	0.26	0.96	0.37	
11	458	0.31	1.7	0.62	
12	459	15	36	16	
13	460	13	36	24	
14	461	0.20	1.0	0.51	

 a Values given are IC $_{50}$ in nM and original activity data can be found in ref. 259–262. b HCT116: human colon cancer cell line; QG56: human lung cancer cell line; DU145: human prostate cancer cell line.

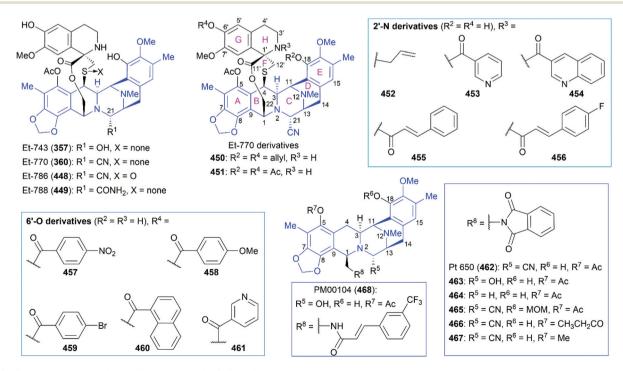


Fig. 20 Representative ecteinascidin derivatives for SAR study.

various conditions failed. Condensation of aldehyde 444 with amine 439 successfully provided 445, which contained the desired two tetrahydroisoquinoline moieties. TFA-promoted deprotection of 446 followed by Strecker reaction and BCl3mediated removal of the two benzyl groups produced amino nitrile 447, which could be transformed into Et-743 in a manner similar to that in Corey's synthetic route.

8.2 Structure-activity relationships of ecteinascidin derivatives

Although the reported total synthetic routes are not amenable to scaling-up for manufacturing ET-743 on an industrial scale, transformations developed in these studies provide access to

Table 3 Cytotoxicities of Pt 650 and related derivatives to four carcinoma cell lines^a

		$IC_{50}\left(nM\right)$	IC_{50} (nM)				
Entry	Compound	A-549 ^b	HCT116 ^b	A375 ^b	PC-3 ^b		
1	Et-743 (357)	1.0	0.50	0.15	0.70		
2	Pt 650 (462)	0.95	0.38	0.17	0.55		
3	463	1.0	0.61	0.20	0.53		
4	464	2.2×10^{3}	1.1×10^{3}	6.1×10^{2}	8.5×10^{2}		
5	465	2.3×10^{2}	74	66	98		
6	466	2.1	1.2	0.51	2.9		
7	467	3.1	1.4	0.55	3.1		

^a Values given are IC₅₀ in nM and original activity data can be found in ref. 257. ^b A-549: human lung cancer cell line; HCT116: human colon cancer cell line; A375: human melanoma cell line; PC-3: human prostate cancer cell line.

diverse ecteinascidin derivatives, some of which are listed in Fig. 20. These synthesized ecteinascidin derivatives make SAR studies of Et-743 possible and have led to the development of more anti-tumor molecules including the approved drug lurbinectedin (363) and PM00104 (468), which is in phase II clinical trials. We give a short discussion about the SAR studies on ecteinascidin derivatives in the following section (Fig. 20 and Tables 2, 3).

After achieving the first total synthesis of Et-743, Corey and co-workers synthesized a series of Et-743 analogs from a common intermediate, which led to the discovery of one simpler and more stable anti-tumor molecule named Pt 650 (phthalascidin, 462). 257 In 2002, Saito and co-workers isolated Et-770 (360), an analog of Et-743, from the pretreated Thai tunicate Ecteinascidia thurstoni with potassium cyanide. 258 Both Pt 650 and Et-770 showed antiproliferative activities on various cancer cell lines. Synthesis and evaluation of more synthesized analogs of Pt 650 and Et-770 revealed the structural features for high potent biological activities. 257-263

A cyano or a hydroxy group at the C21 position is essential and is responsible for alkylating DNA. Et-788 (449) with an amide group at C21 and 464 with hydrogen at C21 showed a significant decrease in activity compared with their corresponding parent compounds.^{257,261} Oxidation of the sulfur atom in Et-770 led to Et-786, which resulted in dramatically diminished cytotoxicity.²⁵⁹ Protection of the C18 and C6' phenolic hydroxy groups with allyl (450) or acetate (451) decreased cytotoxicity. 259,260 Protection of the C18 phenolic hydroxy group with MOM (465) also resulted in diminished antitumor activity.257

Fig. 21 Examples of natural products containing enediyne.

Modifications at the 2'-N position revealed that the 2'-N-allylated compound 452 showed activity similar to that of Et-770 (360), while several 2'-N amide derivatives including nitrogencontaining heterocycles (453 and 454) and cinnamovl derivatives (455 and 456) exhibited improved cytotoxicity. 260-262 Notably, 454 and 456, respectively, exhibited approximately 50and 70-fold higher cytotoxicity to the HCT116 cancer cell line than Et-770 (360).²⁶¹

Evaluation of the cytotoxicity of a series of C6' aromatic ester derivatives showed that 457 and 458 with a nitro group or a methoxy group on the benzene ring exhibited activities similar to Et-770 (360). However, derivatives with a bromo group on the benzene ring (459) or a naphthyl group (460) showed significantly decreased cytotoxicity. The incorporation of a nitrogencontaining heterocyclic ester group (461) resulted in a moderate increase in activity. For Pt 650 analogs, derivatives possessing a propionyl (466) or a methyl (467) protecting group on the C5 phenolic hydroxy group were slightly less effective than the parent compound Pt 650 (462). These results indicate the promising role of total synthesis in the future development of ecteinascidin-type and related natural products into drugs.

Calicheamicin γ_1^l

Calicheamicin $\gamma_1^{\rm I}$ (474) is a potent antitumor antibiotic that was isolated from Micromonospora echinospora ssp in 1986. One year later, its structure was defined as an enediyne with a bicyclo [7.3.1]tridec-9-ene-2,6-divne system, a unique N-O glycosidic linkage, and a methyl trisulfide moiety; other members of the calicheamicin family were also discovered (Fig. 21). 264-266 Several natural products containing the enediyne were also found in the 1980s, such as Neocarzinostatin chromophore (476),²⁶⁷ Esperamicin A1 (477), 268 and Dynemicin A (478) (Fig. 21). Calicheamicin $\gamma_1^{\rm I}$ displays excellent activity against Gram-negative and -positive bacteria at concentrations lower than 1 pg mL⁻¹. It demonstrated extreme toxicity to all cells, indicating that it has extraordinary potency against cancer cells. Investigation of its mechanism showed that calicheamicin γ_1^I targets and binds to the DNA minor groove, and then undergoes the Bergman cyclization after bioreductive activation to produce a biradical, 1,4-didehydrobenzene, which subsequently abstracts hydrogen atoms from the deoxyribose backbone of DNA to generate sugar radicals. The latter undergo oxygen-dependent reactions and finally

Scheme 46 Nicolaou's preparation of tetrasaccharide building block.

induce double-strand DNA breaks. 270-272 Because of its excellent activity and site-specific targeting, gemtuzumab ozogamicin (Mylotag®), a calicheamicin γ_1^{I} derivative, was approved by the FDA in 2000 as an antibody-drug conjugate (ADC) to treat CD-33positive myeloid leukemia. Unfortunately, 10 years later it was withdrawn from the market because of its serious side effects. In 2017, inotuzumab ozogamicin (Besponsa®), also derived from calicheamicin γ_1^{I} , was approved by the FDA as an ADC medication to treat relapsed or refractory CD22-positive B-cell precursor acute lymphoblastic leukemia. All in all, the fascinating structure and highly potent drug-likeness characteristics of calicheamicin $\gamma_1^{\rm I}$ have spurred chemists to meet the huge challenge of developing strategies for the total synthesis of calicheamicin $\gamma_1^{\rm I}$.

9.1 Total synthesis of calicheamicin $\gamma_1^{\rm I}$

9.1.1 Nicolaou's total synthesis of calicheamicin $\gamma_1^{\rm I}$. In 1992, Nicolaou and co-workers reported the first approach for the synthesis of calicheamicin $\gamma_1^{\rm I}$ using the strategy of splitting the parent molecule into two advanced intermediates, an oligosaccharide fragment and an aglycon precursor. 273-276 First, the building blocks 479 and 480 were glycosylated in the presence of AgClO₄ and SnCl₂ to produce α-disaccharide as the major product, which underwent selective deprotection followed by regioselective oxidation of C4-OH to generate hydroxy ketone 481. The active ketone was successfully condensated with hydroxylamine 482 using PPTS as a catalyst to produce trisaccharide 483. The introduction of thionoimidazolide instead of benzoyl generated 484, which proceeded the [3,3]-sigmatropic rearrangement and removal of the imidazoformyl to forge trisaccharide 485 for coupling with aryl saccharide 489. The latter was prepared by the glycosidation of saccharide unit 486 with fully substituted phenol 487 and subsequent interconversion of protecting groups. Photodeprotection of an o-nitrobenzyl glycoside of trisaccharide 490 generated lactol, and

Scheme 47 Total synthesis of calicheamicin γ_1^l by Nicolaou.

Schmidt's protocol was used to generate the trichloroacetimidate 491 from 490 (Scheme 46).

The critical synthesis of aglycon portion is to construct the cyclohexenone precursor of the bicyclic enediyne scaffold using the [3 + 2] nitrile oxide cycloaddition reaction (Scheme 47). The commercially available tetronic acid 492 was used to prepare the aldoxime precursor 493, which underwent an intramolecular 1,3-dipolar cycloaddition reaction using aqueous sodium hypochlorite solution in DCM as an oxidation system to generate isoxazoline 494. After removal of the benzoate group and the Jones oxidation of the resultant secondary hydroxy group, the ketone intermediate was produced. Subsequent addition of lithium(trimethylsilyl)acetylide and acetylation generated intermediate 495 with high stereoselectivity. After careful removal of MEM ether, the resultant secondary alcohol was subjected to Swern oxidation, generating keto isoxazole which was treated with methyl (triphenylphosphoranylidene)acetate followed by protecting group interconversions to generate the olefination product 496 as a single geometrical isomer. This product was subsequently coupled with (Z)-chloroenyne using a Pd(PPh₃)₄-CuI catalytic system to produce the protected enediyne 498. Cleavage of the isoxazole with molybdenum hexacarbonyl (Mo(CO)6), removal of the TMS group, and phthaloylation of amine gave rise to the

desired intermediate 499. The new secondary hydroxy group produced through exposure of 499 to potassium hexamethyldisilazide in toluene was mesylated and subsequently inverted by a neighboring ester group to furnish the lactone 500 in high yield. After exchanging protecting groups and double reduction using DIBAL/NaBH4, the required aglycon intermediate 501 was produced by benzoylation. With the tetrasaccharide donor 491 and aglycon 501 in hand, Schmidt glycosidation was applied to form the coupled product 502 as a single isomer. Then, the target molecule calicheamicin $\gamma_1^{\rm I}$ (474) was generated after multi-step transformations.

9.1.2 Danishefsky's total synthesis of calicheamicin $\gamma_1^{\rm I}$. Three years after Nicolaou's work, Danishefsky and co-workers also finished the total synthesis of calicheamicin $\gamma_1^{\rm I}$ (474). $^{277-281}$ As shown in Scheme 48 tetrasaccharide 516 was constructed from the coupling of two disaccharide building blocks 508 and 514. After deiodination and triflation, the first disaccharide building block 508 was prepared from disaccharide fragment 507, which was forged by the coupling of glycal 506 and 505 in the presence of collidine-complexed iodonium perchlorate. For the second disaccharide, glycosyl donor 509 and acceptor 510 were subjected to the Schmidt procedure to build the aryl glycoside 511 with excellent yield. Subsequently, 511 was converted to acyl chloride 512, which was then successfully coupled with

Scheme 48 Danishefsky's preparation of tetrasaccharide building block.

thiol 513 to provide disaccharide 514. Deprotonation of the carbamate nitrogen of disaccharide 514 using NaH as the base and subsequent alkylation with triflate 508 afforded tetrasaccharide 515, which was converted to glycosyl donor 516.

The synthesis of the aglycon unit started with the synthesis of unstable triol 518 from the commercially available benzoate 517; 518 was subjected to Becker oxidation to generate an epoxide and, following Dess-Martin oxidation, aldehyde 519 was obtained (Scheme 49). The aldehyde 519 was protected by treating with lithium N-methylanilide, and the keto group was attacked by the lithium acetylide generated from the parent enedivne to give adduct 520 after silvlation of the newly formed alcohol by utilizing trimethylsilyl triflate. Then, 520 was subjected to potassium 3-ethyl-3-pentoxide, and the core skeleton 521 was obtained. Protecting group interconversions, acetolysis, and oxidative cleavage were employed to convert 521 to 522. Exposure of 522 to a solution of sodium azide afforded the desired product, azidoenone 523, which underwent subsequent key transformations including reduction of azide, an intramolecular Horner-Wadsworth-Emmons reaction, reduction using NaBH₄, and a Mitsunobu reaction, generating trisulfide 524. Remarkably, the Schmidt protocol using silver triflate and molecular sieves was employed to couple trisulfide 524 with the tetrasaccharide donor 516 to forge 525 as a single isomer with moderate yield.

Not long after the discovery of calicheamicin γ_1^{I} , the structural analogs of calicheamicin, shishijimicin A (526)282 and namenamicin (527),²⁸³ were isolated from marine organisms (Fig. 22). Biological assays showed that they both displayed highly potent antitumor activity. Very recently, Nicolaou and co-workers finished the total synthesis of shishijimicin A (526)^{284,285} and namenamicin (527)²⁸⁶ using a protocol they developed previously, with some modifications. The details of

Fig. 22 The structures of shishijimicin A and namenamicin.

Scheme 49 Total synthesis of calicheamicin γ_1^l by Danishefsky.

these approaches are not discussed here. The derivatives and analogs of shishijimicin A were also constructed using the total synthesis strategy to explore the ADC payloads for clinical trials against tumors.287

9.2 Structure–activity relationships of calicheamicin $\gamma_1^{\rm I}$ derivatives

As mentioned above, since calicheamicin $\gamma_1^{\rm I}$ was discovered as a 10-membered enediyne with highly potent cytotoxicity against tumors, a number of synthetic analogs have been designed and used as ADC payloads in preclinical evaluations and clinical tests against tumors. According to the mechanism of action described, the enediyne subunit of calicheamicin $\gamma_1^{\rm I}$

that plays an irreplaceable role in the generation of DNA lesions through the Bergman rearrangement is unchangeable, and the allylic trisulfide serves as a trigger to induce cycloaromatization. Studies of the semisynthetic aglycon analogs revealed that the role of carbohydrate fragments is to facilitate DNA binding and confer target specificity. 270,288 To date, SAR research has mainly focused on the modification and simplification of oligosaccharide fragments or trisulfide groups.

The four calicheamicin $\gamma_1^{\rm I}$ analogs (Fig. 23), the natural products 471 and 472 and synthetic compounds 528 and 529, were all conjugated to CT-M-01 and the resultant derivatives were tested for therapeutic efficacy against MX-1 xenograft tumors. The results showed that compounds 472 and 528 with

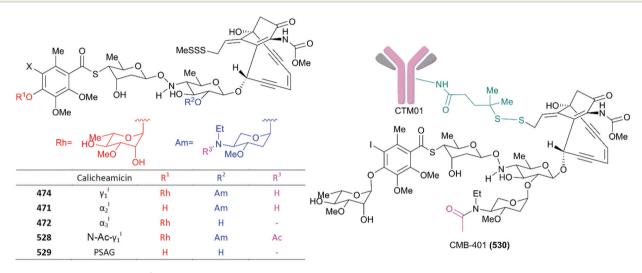


Fig. 23 Examples of calicheamicin γ_1^l derivatives(i).

Fig. 24 Examples of calicheamicin γ_1^I derivatives(II).

gemtuzumab ozogamicin (534) /inotuzumab ozogamicin (535)

Fig. 25 Structures of the marketed ADC drugs gemtuzumab ozogamicin and inotuzumab ozogamicin.

the DNA binding unit and the lack or modifications of amino sugar fragments showed stronger therapeutic efficacy than the parent molecule 474 and the derivatives 471 and 529, which missed the rhamnose at the DNA binding region. This result suggested that a rhamnose at the end of the binding region is required for ADC medications. ²⁸⁹ CMB-401 (530) was further developed using a more stable amide-disulfide-based linker to conjugate it with hCTMO1 based on the structure of compound 528, and this compound was tested in phase I and phase II studies against epithelial ovarian cancer. However, no patients had objective responses in the phase II investigation. Scientists speculated that the sterically hindered disulfide failed to release the drug from the ADC payload. ²⁸⁸

Calicheamicin $\theta_1^{\rm I}$ (Fig. 24, 531), a novel derivative of calicheamicin $\gamma_1^{\rm I}$, was developed using thioester to replace the methyl trisulfide and prepared through total synthesis by the Nicolaou group. After the test, calicheamicin θ_1^I was hypothesized to be a powerful biological tool for generating specific DNA lesions because of its highly potent and selective DNA cleaving, apoptosis-inducing, and cytotoxic properties.²⁹⁰ It was also discovered that calicheamicin θ_1^{I} (531) effectively suppressed liver metastases in a novel syngeneic model of murine neuroblastoma and CD19positive malignancies. ^{291,292} In some analogs, the allylic trisulfide was replaced by isopropyl disulfide or functionalized to improve the stability and solubility (Fig. 24, 532 and 533). 293,294 Gemtuzumab ozogamicin (Mylotag®, 534) and inotuzumab ozogamicin (Besponsa®, 535) are ADC drugs approved to treat myeloid leukemia and B-cell malignancies by the FDA in 2000 and 2017, respectively. Because of the important role that the linker plays in the effective release of the calicheamicin derivative from the conjugation product, allowing it to access the DNA target, a linker, 4-(4-acetylphenoxy)butanoic acid, was added to condensate drug and antibody. This conjugate showed stability in physiological buffer (pH 7.4) and the calicheamicin derivative was efficiently released inside lysosomes (pH 4.0) (Fig. 25).²⁹⁵

10. Conclusions

In this review, we discussed the total synthesis and SAR studies of eight representative families of natural products

applied in the treatment of various human diseases. Although just the tip of the iceberg, these examples demonstrate the essential role of total synthesis in natural product drugs' discovery: development of efficient transformations for natural products synthesis, especially the deep-seated structural modifications that are otherwise tough to achieve; and providing adequate and sustainable supplies of natural products and their analogs for clinical studies and use. From another standpoint, total syntheses of natural products, which often contain complex structural features, also promote the development of synthetic chemistry by inspiring the discovery of new reactions and reagents for rapid molecular assembly and structural diversifications. In combination with the development of other disciplines, such as synthetic biology and chemical biology, chemical synthesis will continue to play a profound role in functional mechanism elucidation, new biological target identification, and natural product medicinal chemistry.

Abbreviations

 $\begin{array}{lll} \mbox{AcOH} & \mbox{Acetic acid} \\ \mbox{AIBN} & 2,2'\mbox{-Azobisisobutyronitrile} \\ \mbox{BHT} & \mbox{Butylated hydroxytoluene} \\ \mbox{Boc}_2\mbox{O} & \mbox{Di-}tert\mbox{-butyl dicarbonate} \\ \mbox{BOPCl} & \mbox{Bis}(2\mbox{-oxo-}3\mbox{-oxazolidinyl})\mbox{phosphinic chloride} \\ \end{array}$

BPO Benzoyl peroxide
CSA Camphorsulfonic acid
DABCO 1,4-Diazabicyclo[2.2.2]octane
DBU 1,8-Diazabicyclo[5.4.0]undec-7-ene

DCM Dichloromethane
DEAD Diethyl azodicarboxylate
1,2-DFB 1,2-Difluorobenzene

DIPAMP (Ethane-1,2-diyl)bis[(2-methoxyphenyl)(phenyl)

phosphine]

DIPEA *N*,*N*-Diisopropylethylamine DMAP 4-(*N*,*N*-Dimethylamino)pyridine

DMDO Dimethyldioxirane
DME 1,2-Dimethoxyethane
DMF N,N-Dimethylformamide
DMSO Dimethyl sulfoxide

Dppp

1,3-Bis(diphenylphosphino)propane 2,6-Di-*tert*-butyl-4-methylpyridine

EDCI N-(3-Dimethylaminopropyl)-N'-ethylcarbodiimide

hydrochloride

HFIP Hexafluoroisopropanol HMPA Hexamethylphosphoramide

IBX 2-Iodoxybenzoic acid

KHMDS Potassium bis(trimethylsilyl)amide

LDA Lithium diisopropylamide

LICA Lithium N-isopropylcyclohexylamide

LiDBB Lithium di-tert-butylbiphenyl
LiHMDS Lithium bis(trimethylsilyl)amide
LTMP Lithium tetramethylpiperidide
m-CPBA meta-Chloroperoxybenzoic acid

MesLi Mesityllithium MS Molecular sieve

MsCl Methanesulfonyl chloride NaHMDS Sodium bis(trimethylsilyl)amide

NBS N-Bromosuccinimide

NMO 4-Methylmorpholine *N*-oxide
NTf₂ Bis(trifluoromethylsulfonyl)imide
Pd₂(dba)₂ Bis(dibenzylideneacetone)palladium
PhNTf₂ Bis(trifluoromethanesulfonyl)aniline
PMP 1,2,2,6,6-Pentamethylpiperidine
PPTS Pyridinium *p*-toluenesulfonate

Red-Al Sodium bis(2-methoxyethoxy)aluminiumhydride [Rh(COD) Chloro(1,5-cyclooctadiene)rhodium(1) dimer

 $Cl]_2$

Rt Room temperature

SARs Structure–activity relationships
TBAF tert-Butylammonium fluoride
TBAI tert-Butylammonium iodide

TBSOTf tert-Butyldimethylsilyl trifluoromethanesulfonate

TESOH Triethylsilanol

Tf₂O Trifluoromethanesulfonic anhydride

TFA Trifluoroacetic acid
TFAA Trifluoroacetic anhydride
TFE 2,2,2-Trifluoroethanol

TfOH Trifluoromethanesulfonic acid

THF Tetrahydrofuran

TMEDA Tetramethylethylenediamine
TMSCl Chlorotrimethylsilane
TMSCN Trimethylsilyl cyanide

TMSH Trimethylsulfonium hydroxide TMSimid 1-(Trimethylsilyl)imidazole

TMSOH Trimethylsilanol

TPAP Tetrapropylammonium perruthenate

TPP Triphenylphosphine
TsOH p-Toluenesulfonic acid

Author contributions

X. Q. conceived the study; G. L., M. L. and X. Q. prepared the manuscript; all authors discussed the contents and commented on the manuscript.

Conflicts of interest

The authors declare no conflict of interests.

Acknowledgements

This work was supported by the National Natural Science Foundation of China (NSFC) (Grant No. 21971018). We gratefully acknowledge the Beijing Municipal Government and Tsinghua University for financial support.

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