

Design, Synthesis, and Biological Evaluation of Diminutive Forms of (+)-Spongistatin 1: Lessons Learned

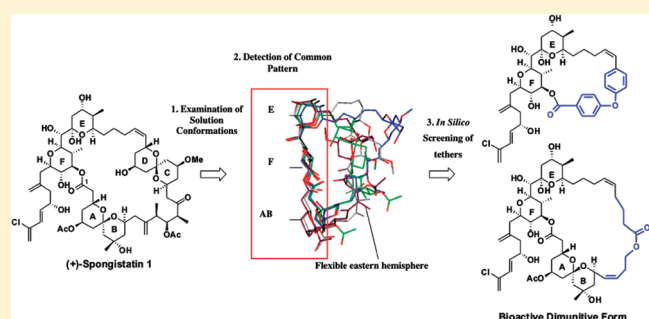
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S Supporting Information

ABSTRACT: The design, synthesis, and biological evaluation of two diminutive forms of (+)-spongistatin 1, in conjunction with the development of a potentially general design strategy to simplify highly flexible macrocyclic molecules while maintaining biological activity, have been achieved. Examination of the solution conformations of (+)-spongistatin 1 revealed a common conformational preference along the western perimeter comprising the ABEF rings. Exploiting the hypothesis that the small-molecule recognition/binding domains are likely to comprise the conformationally less mobile portions of a ligand led to the design of analogues, incorporating tethers (blue) in place of the CD and the ABCD components of the (+)-spongistatin 1 macrolide, such that the conformation of the retained (+)-spongistatin 1 skeleton would mimic the assigned solution conformations of the natural product. The observed nanomolar cytotoxicity and microtubule destabilizing activity of the ABEF analogue provide support for both the assigned solution conformation of (+)-spongistatin 1 and the validity of the design strategy.



INTRODUCTION

The spongistatins (1–9, Table 1), independently reported by the Pettit,¹ Fusetani,² and Kitagawa³ laboratories in the early 1990s, comprise a unique family of marine macrolides that embody remarkably complex structures possessing extraordinary antimitotic activity. Members of this family, in particular spongistatins 1 and 2 [(+)-1 and (+)-2, Table 1], rapidly attracted broad interest in the chemical and biological communities, given their potential as cancer chemotherapeutic agents. Spongistatin 1, recognized at the time as one of the most selective cytotoxic agents known, has an average IC₅₀ value of 0.12 nM against the NCI panel of 60 human cancer cell lines.^{1,4} The initially observed *in vitro* cell growth inhibition activity was subsequently explored by Hamel's group for tubulin polymerization, competitive microtubule assembly, and turbidity/aggregation.⁵ Their results revealed that (+)-spongistatin 1 (1) competitively inhibits tubulin binding of maytansine and rhizoxin, as well as GTP exchange;^{5,6} (2) noncompetitively inhibits tubulin binding of dolastatin 10, halichondrin B, and vinblastine;^{5,6} (3) does not induce formation of GTP-independent microtubule aggregates;⁶ and (4) inhibits formation of the Cys-12-Cys-201/211 cross-linking on tubulin.⁷

On the basis of these results, Hamel and co-workers proposed a “polyether” binding site on β -tubulin for the spongistatins located near the *vinca* domain,⁸ distinct from but in the vicinity of the “peptide” and “*vinca*” sites, where dolastatin 10/phomopsin

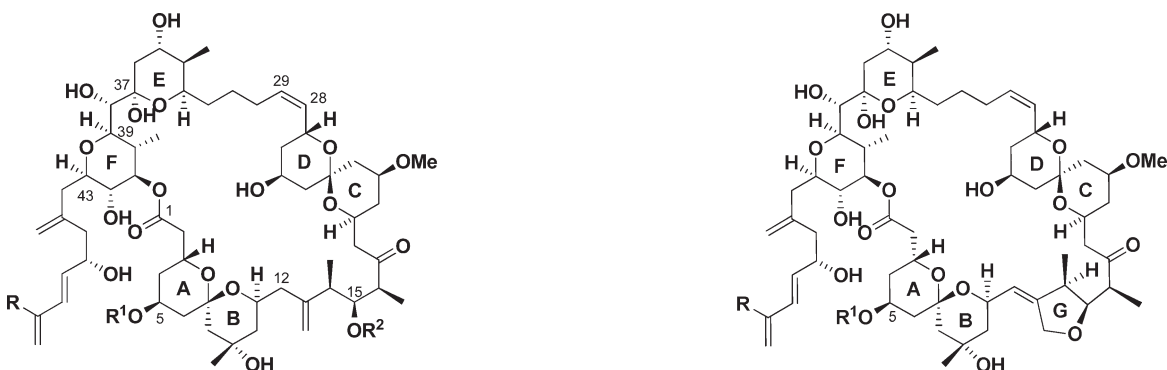
A and vinblastine/vincristine respectively are known to bind.⁶ The development of the spongistatins as possible clinical agents, however, has not progressed given their low availability, notwithstanding seven total syntheses of (+)-spongistatin 1⁹ and four total syntheses of (+)-spongistatin 2.¹⁰

Following completion of our first-generation total syntheses of (+)-2 and (+)-1, respectively we focused on material advancement to deliver more than 1 g of synthetic (+)-1, via what in the end comprised an effective fourth-generation synthetic route.^{9b,11} In addition to defining the limitations and scalability issues along the synthetic route, a significant outcome of this synthetic campaign was the development of a one-pot, multi-component dithiane union tactic,¹² based on the earlier efforts of the Tietze laboratories.¹³ This tactic has now evolved into a new synthetic paradigm termed Type I and Type II Anion Relay Chemistry (ARC)¹⁴ for the rapid construction of polyketide¹⁵ and alkaloid¹⁶ natural and unnatural products.¹⁷

With sufficient material for preclinical development in hand (ca. 1 g), we turned our attention to identifying the architectural subunits of the spongipyran skeleton responsible for the subnanomolar cytotoxicity, with the goal of initiating structure–activity relationship (SAR) studies. Notwithstanding the optimized (+)-spongistatin 1 route, construction of random analogues

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Table 1. Inhibition of Cell Growth of L1210 Murine Leukemia Cells and Average GI₅₀ Values of 60 Different Cancer Cell Lines^a


spongistatin	L1210	avg.	spongistatin	L1210	avg.
1 [(+)-1]; R = Cl, R ¹ = R ² = Ac	0.03	0.13	5 (6); R = Cl, R ¹ = H	0.20	0.12
2 [(+)-2]; R = H, R ¹ = R ² = Ac	2.0	0.85	7 (7); R = R ¹ = H	2.0	1.00
3 (3); R = Cl, R ¹ = H, R ² = Ac	1.0	0.83	8 (8); R = H, R ¹ = Ac	3.0	0.23
4 (4); R = Cl, R ¹ = Ac, R ² = H	0.10	0.10	9 (9); R = Cl, R ¹ = Ac	0.3	0.04
6 (5); R = R ² = H, R ¹ = Ac	0.80	1.10			

^a Data compiled from ref 1. GI₅₀ values are in nM.

appeared unattractive. We were, of course, well aware of the remarkable success of the commercial drug Halaven (eribulin), based on the marine natural product halichondrin B, developed by the Kishi and Eisai Inc. laboratories. We asked the question, *Is the entire structure of the large, flexible polyketide skeleton of the spongistatins a prerequisite for biological activity?*

In this, a full account, we report the design, synthesis, and biological evaluation of two diminutive forms of (+)-spongistatin 1. Importantly, computational methods developed during this venture led to (1) a general strategy to identify the binding regions of flexible small-molecule *macrolide* natural products and, more specifically, (2) a general design strategy that permitted the simplification of the polyketide structure of the spongistatins while maintaining biological activity.

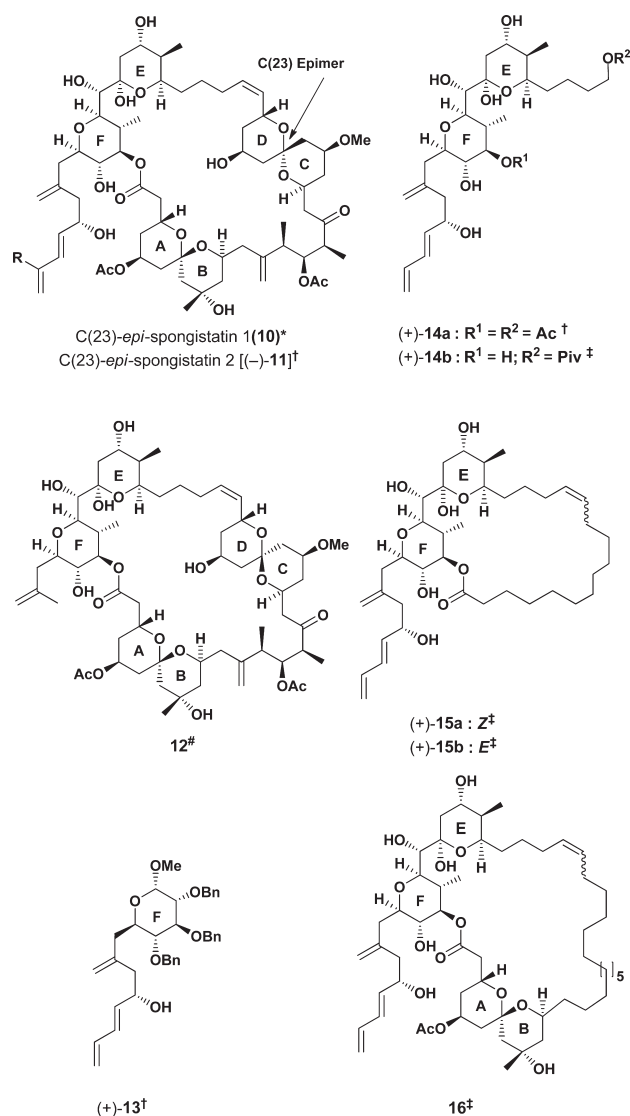
■ SPONGISTATIN ANALOGUES: EARLY CONGENERS

Members of the spongistatin family (1–9, Table 1) possess an array of architecturally complex structural features, including a 31-membered macrolactone ring endowed with 24 stereocenters, two [6,6] spiroketals (the AB and CD rings), and one bis-tetrahydropyranylmethane moiety (E and F rings) complete with a diene side chain differentiated by the acetate substitution pattern at C(5) and C(15) and, in the case of (+)-spongistatins 6–9, by an additional tetrahydropyran ring (G ring) inscribed in the tether linking rings B and C. The cell growth inhibitory activity against the NCI-60 DTP human tumor panel for the natural congeners averaged at the sub-nanomolar level. Structural features for optimal activity comprise the chlorine substituent on the side chain (1 vs 2; 4 vs 6; and 6 vs 7) and an acetate at C(5) (1 vs 3; and 6 vs 9), while structural elements not critical for maintaining activity include the C(15) acetate and ring G (1–5 vs 6–9). Although these observations provided useful insights, our analogue design criteria also benefited initially from synthetic intermediates and analogues that emanated from the Kishi,^{9a,b} Paterson,¹⁸ and Heathcock¹⁹ laboratories as well as our own.²⁰

For example, in Kishi's synthesis of (+)-spongistatin 1,^{9b} as well as in our synthesis of (+)-spongistatin 2,^{10b,e} unexpected epimerizations that occurred at the C(23) center of the CD spiroketal led to congeners **10** and (–)-**11** (Figure 1) that possessed modest tumor cell growth inhibition against several cancer cell lines [avg. GI₅₀ of 0.2 μM for (–)-**11**].^{20c} Subsequently, Paterson and co-workers constructed an analogue lacking the diene side chain (**12**) that significantly attenuated activity, highlighting the requirement for this structural unit.¹⁸ Later we prepared a series of glucose-based mimetics of the F and EF rings possessing the (+)-spongistatin 2 side chain [(+)-**13** and (+)-**14a**].²⁰ Although modest tumor cell growth inhibition activity was observed with (+)-**13**, subsequent unpublished results suggested a different mode of action.^{20a,21} More recently, Heathcock and co-workers reported a series of (+)-spongistatin 2 analogues, including (+)-**14b**, which did not show any activity at the tested concentrations.¹⁹ The loss of activity led Heathcock to prepare macrolide congeners **15** and **16**, employing polymethylene tethers to replace the ABCD and CD structural segments. Observation of only modest activity (GI₅₀ values of 4.8 and 4.6 μM against HCT116 human colon tumor cell lines, respectively) led Heathcock to conclude that the CD ring may comprise a structural moiety required to maintain the tumor cell growth inhibition.

■ ANALOGUE DESIGN: AN EF RING CONGENER POSSESSING A ROTATIONALLY RESTRICTED ABCD RING SYSTEM

The lessons learned from analogues (+)-**13**–**16**, in conjunction with the activity observed for the C(23) *epi*-congeners [**10** and (–)-**11**], led us to the hypothesis that the western perimeter of the spongistatins constituted a significant component of the recognition domain. We reasoned that the linker (CD) was not important and might be replaced by a less complex unit, as long as effective control over the conformation of the western perimeter was maintained (i.e., the dihedral angles between rings E and F). Similar



* Kishi et al. ref. 9a, 9b; # Paterson et al. ref. 18;

† Smith et al. ref. 20; ‡ Heathcock et al. ref. 19

Figure 1. Spongistatin analogues.

simplifying design tactics have been employed by the Kishi,²² Wender,²³ Taylor,²⁴ and Romo²⁵ laboratories to construct biologically active halichondrin B, bryostatin, epothilone, and pateamine analogues, respectively.

To design a suitable tether that would orchestrate the C(37)–C(39) dihedral angles linking the spongistatin E and F rings, we turned to molecular modeling. We were cognizant of the solution conformation reported by Kitagawa²⁶ during his elegant structural assignments of 5-desacetylaltohyrtin A (spongistatin 3, 3). To establish a baseline on the solution conformation of (+)-spongistatin 1 [(+)-1], we carried out repetitive Monte Carlo conformational searches on (+)-1 until no additional families of conformers were detected. Computations utilizing the MMFF force field²⁷ with the GB/SA solvation model²⁸ revealed that the energy ordering of the conformational populations changed significantly in different solvents.

According to our calculations, in chloroform, (+)-1 adopts a “flat” conformation in which the surface area of the molecule is

maximized by an increase in number of intramolecular hydrogen bonds (Figure 2).^{24d} In water, a “twisted” conformer is preferred in which the oxygen atoms are oriented toward the solvent. When DMSO and acetonitrile solvent models are employed, the lowest energy conformer possesses a “saddle” shape. Interestingly, the orientation of the EF rings in the Kitagawa solution structure did not resemble any of these low-energy conformers. We reasoned that the derived Kitagawa structure might well have comprised a *virtual* conformation, resulting from constrained conformational searches.

To understand further the solution behavior of (+)-spongistatin 1, molecular dynamics (MD) simulations were performed using water as solvent, wherein changes in the dihedral angles of the macrocycle were analyzed. The flexible regions of (+)-1 were identified by examining (1) the dihedral angle distributions of each C_{sp}³–C_{sp}³ bond (Figure 3A) and (2) the long-range coupled torsional changes during the dynamics simulation. This analysis was performed exploiting polar coordinate graphs, in a fashion similar to Taylor's epothilone studies,²⁴ wherein the coupled torsions were identified by analyzing the correlation in the torsional changes via principal component analysis. As illustrated in Figure 3A, torsional bonds shown in red have rigid torsions, those in green have flexible torsions, and those in blue have torsions with intermediate flexibility. In Figure 3B, the numbers indicate bonds where the torsions change together, corresponding to correlated movements. As can be seen, tethers of the CD ring are flexible, with the dihedral angle pairs labeled with the same number changing together, thereby permitting movement of the CD ring. Taken together, the results of this multivariate analysis reveal (1) that the western perimeter, including ABEF rings, has a preferred conformation wherein each torsional angle has a preferred value and (2) that reduction of a large, flexible molecule to a single solution structure derived by NMR constraints in general is not feasible.

In support of these observations, MD simulations constrained by NMR-derived distance and torsion values are known to generate average conformations; however, the conformations generated are often of high energy due to the constraints applied. Equally important computational methods, relying solely on molecular mechanics force fields, suffer from inaccurate energy ordering of flexible organic molecules, due to the additive nature of parametrization errors. To circumvent these issues, we devised a computational hybrid strategy wherein the NMR-derived interproton distances and torsional angles are used to determine the most populated families of conformations. This effort led to the development and implementation of a software program termed Distribution of Solution Conformations (DISCON). The thrust of this software package is to define a combination of conformers that overall would provide the best match for the time-averaged torsional angles and interproton distances obtained by NMR. Contrary to methods developed earlier,²⁹ DISCON utilizes hierarchical clustering of conformations based on NMR variables for feature selection to avoid “overfitting,”³⁰ as well as a genetic-algorithm-based global optimization strategy to avoid local minima, two common problems often observed in NMR-based structure optimization. Details and availability of this software package for the deconvolution of NMR observables over an ensemble of solution conformations will be provided elsewhere.³¹

From this analysis, we deduce that the low-energy conformations determined in water and chloroform (Figure 4a–d) define the major solution conformations of (+)-spongistatin 1. The two major calculated conformations comprise twisted

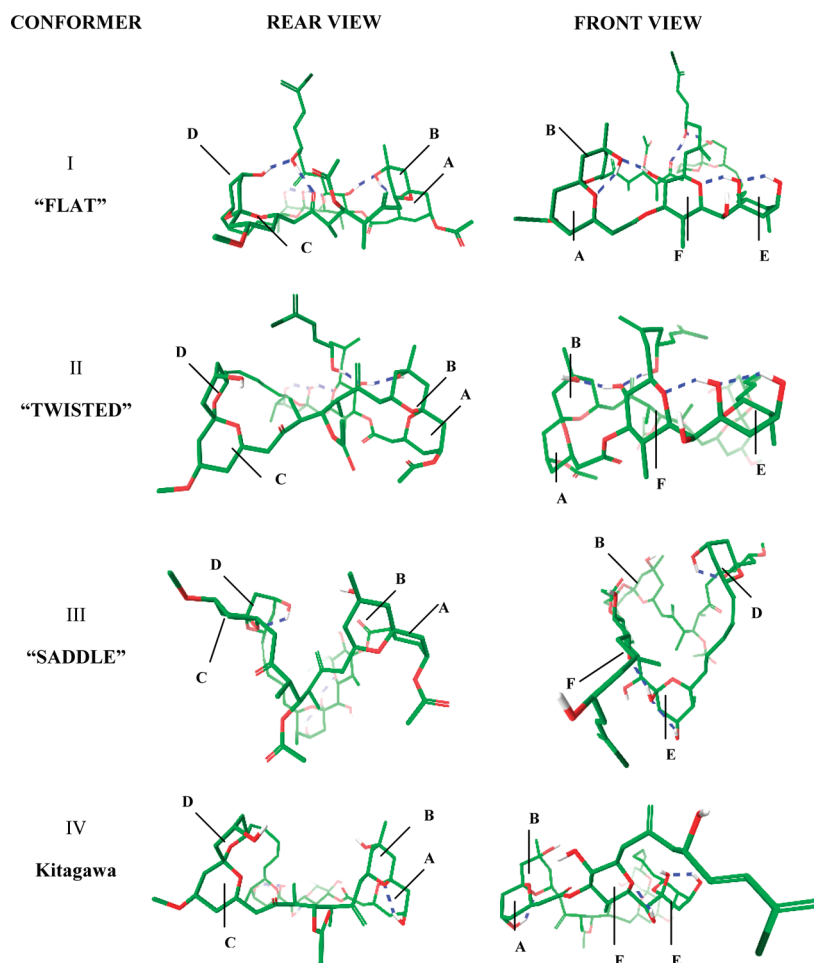


Figure 2. Front and rear 3D views (see Supporting Information section for stereoview images) of the lowest energy conformations (I–III) calculated by conformational searches employing different solvation models and Kitagawa solution structure (IV). Blue dashed lines represent hydrogen bonds. Rings A–F are shown in each conformation.

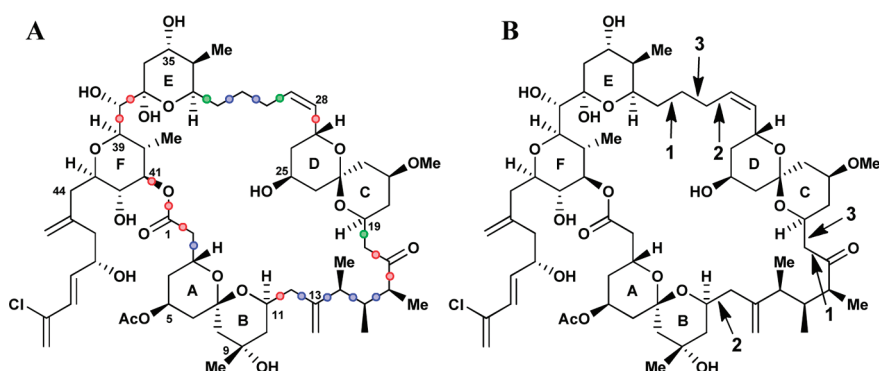


Figure 3. Flexibility of macrocyclic dihedral angles in (+)-spongistatin 1. (A) Red, rigid torsions; green, flexible torsions; and blue, torsional angles with intermediate flexibility. (B) Numbers indicate bonds pairs where the torsions change together, corresponding to long-range movements during the simulations.

and flat forms (57% and 13%, respectively), as illustrated in Figure 4a,b.

The minor conformers (Figure 4c,d) are also similar to Figure 4b, with the major difference being the orientation of the CD ring. Pleasingly, when the four major conformations are overlaid, a striking feature appears: the western hemispheres,

including the ABEF rings, have a common conformation (Figure 4e). This preference can be understood by earlier conformational studies of the bis-tetrahydropyranylmethane system reported by Hoffmann and co-workers³² (Figure 4f), wherein the preferred “skew” conformation minimizes the *syn*-pentane interactions between the E and F ring systems. This

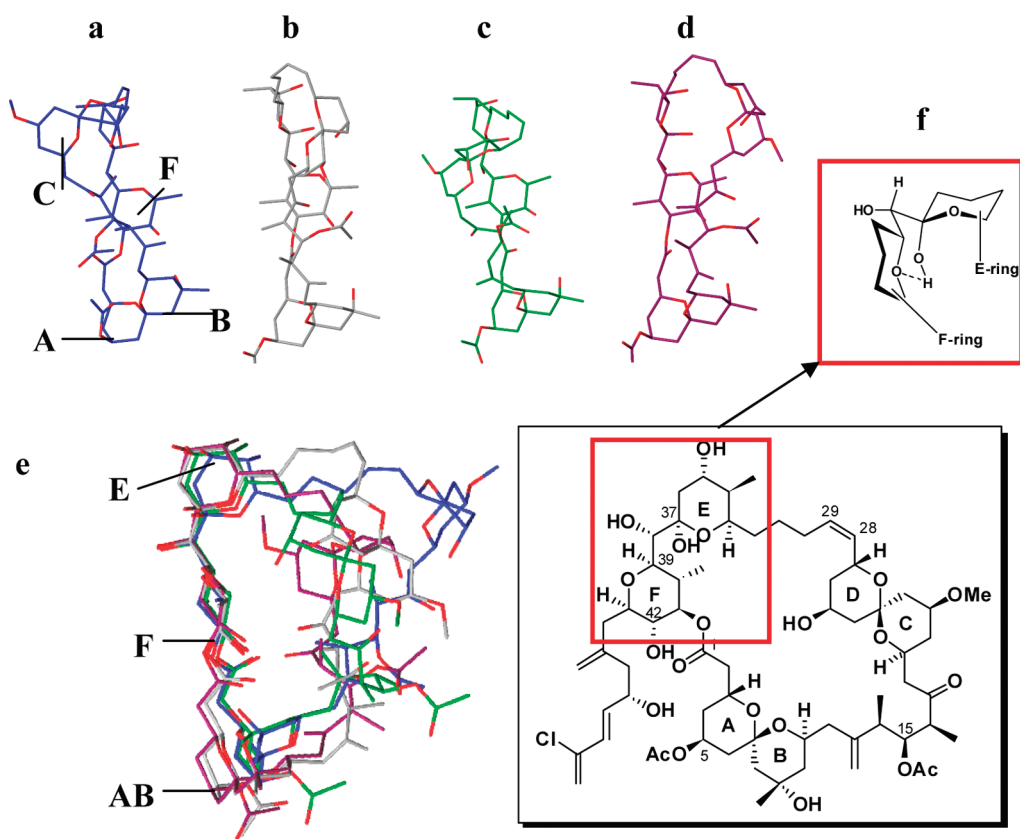


Figure 4. (a–d) Calculated major solution conformers. (e) Overlay of these conformations, showing the conserved ABEF ring conformations, where *syn*-pentane interactions (f) are important in fixing the relative conformations of E and F rings.

conformational arrangement of the E and F (+)-spongistatin 1 rings in turn facilitates an intramolecular hydrogen bond between the E ring hydroxyl and the F ring pyran oxygen (cf. the hydrogen bonds in Figure 2I,II).

The striking conformational similarities derived from the earlier MD simulations and subsequent DISCON analysis, in conjunction with the biological data from the *epi*-C(23) spongistatin congeners **10** and (–)-**11**, provided further support for the hypothesis that the ABEF portion of the spongistatin scaffold comprises the binding/small-molecule recognition domain that interacts with β -tubulin.

To explore this scenario, we first turned to analogues possessing only the E and F ring systems. *In silico* screening of a variety of linkers led to the biaryl ether tether in **17** (Scheme 1, red), which would be suitable to orchestrate the relative conformational

twist of the E and F ring systems, to access the proposed conformation of the western perimeter.

To test our conformational design strategy, we turned to synthesis. From the synthetic perspective, **17** was easily reduced to three fragments: phosphonium salt **18**, allylstannane **19**, and aldehyde **20** (Scheme 1). The route was designed to provide synthetic flexibility for SAR studies by a late-stage installation of the diene side chain, exploiting intermediates in hand from our earlier spongistatins syntheses. One of the attributes of developing preparative-scale syntheses of architecturally complex natural products possessing significant bioregulatory properties is the subsequent ready availability of advanced intermediates for SAR studies. Fragments (+)-**18** and aldehyde **20** were thus envisioned to be united via a Wittig reaction. Following macro-lactonization, side-chain attachment would be achieved via the elegant glucal–epoxide union protocol developed by Evans et al.³³ during the first total synthesis in the spongistatin area. Aldehyde **20** was particularly appealing, given Ullmann coupling protocols.³⁴

We began the synthesis of **17** with (+)-**21**, from our second-generation (+)-spongistatin 1 synthesis.^{9f} We envisioned that (+)-**21** could be readily converted to the corresponding F-ring dihydropyran (+)-**22** by treatment with trifluoromethanesulfonic anhydride (Tf₂O), followed by elimination of TfOH (Table 2). In the event, use of lithium diisopropylamide (LDA) provided an inseparable mixture (ca. 1:1) of (+)-**22** and a compound whose structure was tentatively assigned as **23**. Reasoning that a bulkier base might favor the desired compound, use of KHMDS resulted in a 3:1 mixture of (+)-**22** and **23**, albeit

Scheme 1. Retrosynthetic Analysis of EF Ring Analogue 17

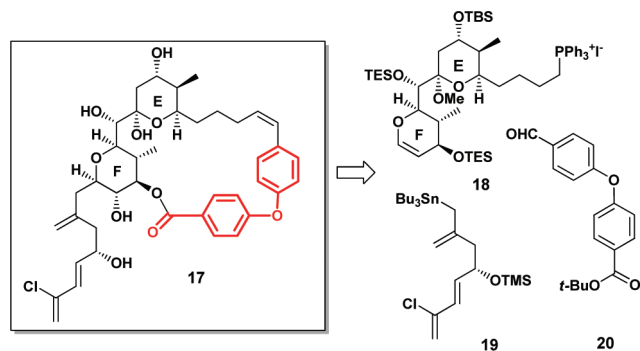
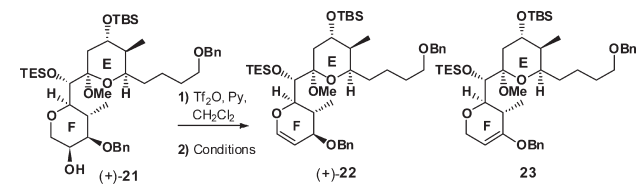


Table 2. Selectivity in the Preparation of F Ring Pyran

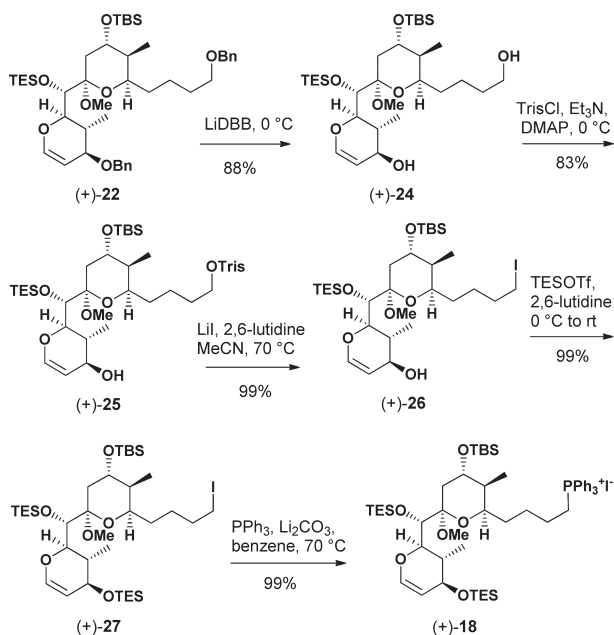


entry	base	temp (°C)	yield (%)	ratio (+)-22:23
1	LDA	−45 to −20	56	1:1
2	KHMDS	−45 to −20	35	3:1
3	NaHMDS	−78 to −30	72	12:1

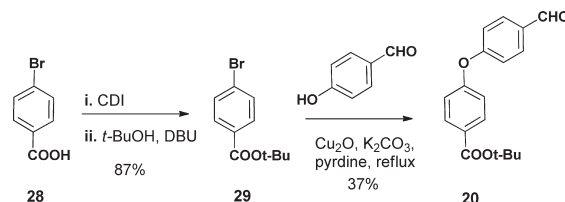
in modest yield (35%). Cooling the reaction mixture to -78°C led to exclusive formation of (+)-22, although the reaction did not proceed to completion. In an effort to increase the efficiency, we examined the use of the less basic NaHMDS. Pleasingly, significant improvement in selectivity (12:1) was observed to provide (+)-22 in 72% overall yield. With reliable access to glucal (+)-22, removal of the benzyl protecting groups employing lithium 4,4'-di-*tert*-butylbiphenylide (LiDBB) proceeded smoothly to provide diol (+)-24 in good yield (Scheme 2). Selective conversion of the primary hydroxyl to furnish the 2,4,6-triisopropylbenzenesulfonate (trisylate) (+)-25 initially proved problematic when the bulky base 2,6-di-*tert*-butyl-4-methylpyridine (DTBMP) was employed. The low reactivity of (+)-24 led us to screen a series of the bases. Triethylamine with DMAP at 0°C proved optimal, selectively furnishing (+)-25 in 83% isolated yield, which in turn was treated with LiI in the presence of 2,6-lutidine to generate iodide (+)-26. The secondary hydroxyl was then converted to the TES ether, followed by displacement of the iodide in (+)-27 with triphenylphosphine to furnish Wittig partner (+)-18 in nearly quantitative yield.

The requisite biaryl ether tether 20 (Scheme 3) was constructed via condensation of 4-bromobenzoic acid with *tert*-butanol, mediated by 1,1'-carbonyldiimidazole (CDI),³⁵ followed

Scheme 2. Preparation of (+)-18

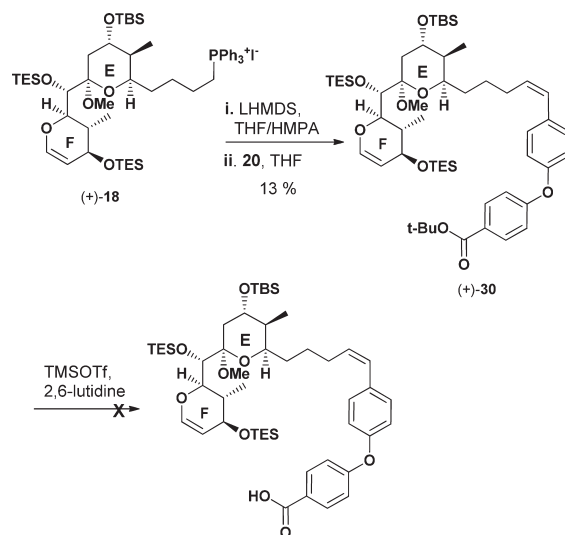


Scheme 3. Preparation of Biaryl Ether 20



by Ullmann coupling³⁴ with 4-hydroxybenzaldehyde to yield 20. While the efficiency of the last step was only modest, we moved forward to explore the viability of this tether.

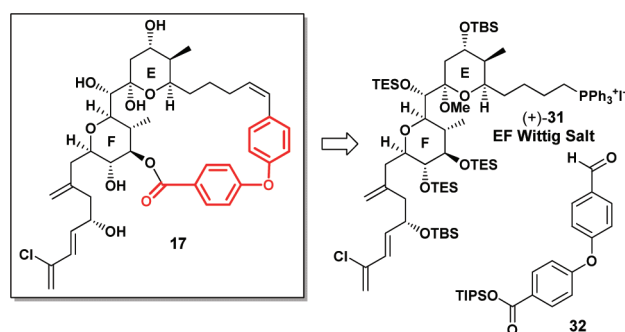
Wittig union³⁶ of (+)-18 with 20 provided the Z-olefin geometry, albeit in low yield (Scheme 4). With studies to improve both the Ullmann and Wittig unions underway, we

Scheme 4. Attempted Removal of *tert*-Butyl Group in (+)-30

proceeded with removal of the *tert*-butyl ester. Unfortunately, various conditions, including treatment of (+)-30 with TMSOTf, resulted only in decomposition, presumably due to the incompatibility of the enol ether with the required mild Lewis acidic conditions.

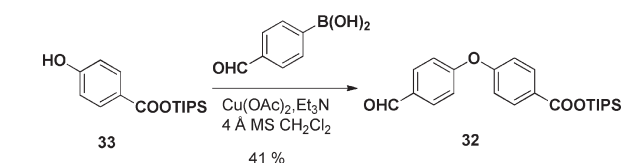
Concurrent with studies to access 17 from (+)-13, we also had access to EF Wittig salt (+)-31 (Scheme 5), again developed in

Scheme 5. Second-Generation Retrosynthesis of EF Ring Analogue 17



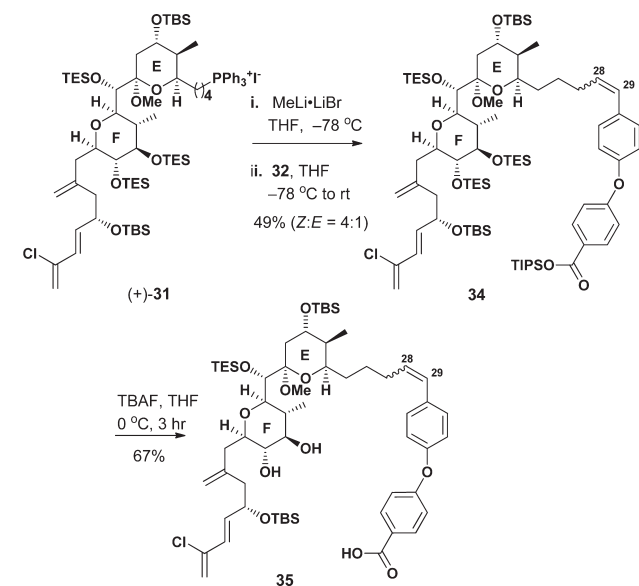
conjunction with our gram-scale synthesis of (+)-spongistatin 1.^{10e} A second-generation approach to the spongistatin EF analogue 17 was thus initiated on the basis of the availability of (+)-31. In the forward sense, we envisioned a biaryl linker now possessing a triisopropylsilyl (TIPS) group to protect the carboxylic acid, in view of our experience with the *tert*-butyl ester. Construction of this biaryl ether tether, employing the Evans-modified Ullmann aryl ether synthesis,³⁷ proved to be compatible with the labile TIPS ester (Scheme 6).

Scheme 6. Preparation of 32



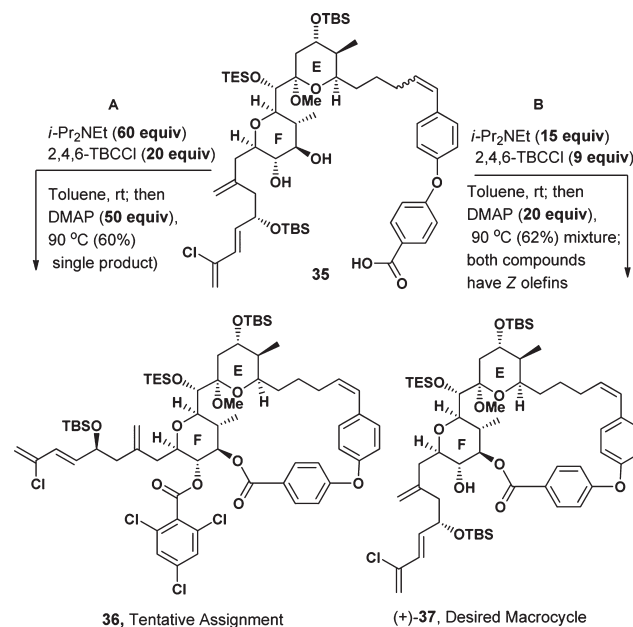
Union of 32 with EF Wittig salt (+)-31 was achieved upon deprotonation with MeLi·LiBr (Scheme 7). Addition of 32 provided 34 as in an inseparable mixture (4:1) of *Z*- and *E*-olefin isomers. The lack of complete *Z*-selectivity was surprising, given that the identical conditions in the gram-scale synthesis of (+)-spongistatin 1 led exclusively to the *Z*-olefin. The mixture of olefin isomers, without separation, was then subjected to TBAF (3 equiv) to effect deprotection of both the TIPS ester and the F-ring TES ethers; *seco*-acid 35, as a mixture of olefin *Z*:*E* isomers (4:1), was obtained in 67% yield.

Scheme 7. Preparation of Macrolactonization Precursor 35



The macrocyclization conditions that had proven successful in the preparation of (+)-spongistatin 1 [60 equiv of *i*-Pr₂NEt, 20 equiv of 2,4,6-trichlorobenzoyl chloride (TBCCl) in toluene at room temperature, then DMAP, 90 °C, Scheme 8A] were explored first. A single product containing a *Z*-olefin was generated, presumably due to the strain associated with incorporating an *E*-olefin into the macrocycle. The desired macrocycle (+)-37, however, was not the product, but instead trichlorobenzoate 36, tentatively assigned on the basis of NMR and MS

Scheme 8. Macrolactonization of 35

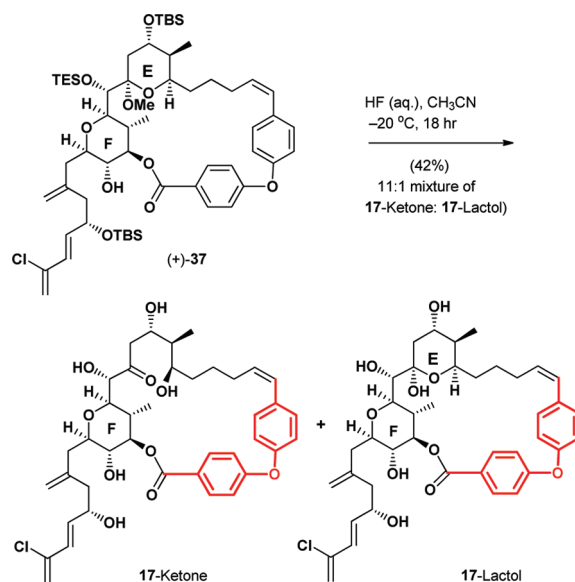


analysis. Presumably, following macrocyclization, the molecule adopts a conformation which facilitates acylation at C(42). Reduction of the amount of each reagent [15 equiv of *i*-Pr₂NEt, 9 equiv of 2,4,6-TBCCl, and 20 equiv of DMAP, Scheme 8B] furnished the desired macrocycle (+)-37, along with a minor inseparable impurity.

Global deprotection next yielded a mixture of products (11:1). Unfortunately, the major product was not the desired 17-Lactol (Scheme 9) but instead was identified as the *E*-ring-opened 17-Ketone, assigned on the basis of a ¹³C NMR carbonyl resonance at 211.4 ppm. For comparison, the hemiketal carbon resonance of (+)-37 appears at 100.5 ppm.

Isodesmic calculations,³⁸ employing 17-Ketone, (+)-1, and the corresponding C(1) *seco*-acids, were performed in a vacuum

Scheme 9. Global Deprotection of (+)-37 under Conditions Followed in the Gram-Scale Spongistatin 1 Synthesis



with the MMFF force field. The results revealed the increase in strain energy upon macrocyclizations of (+)-1 and 17-Lactol to be 5.8 and 45.2 kJ/mol, respectively. In a similar fashion, calculations employing the 17-Lactol and 17-Ketone revealed the release in strain energy upon opening of the pyran ring of 17-Ketone to be 11.2 kJ/mol. Although not definitive, these calculations suggest that, upon acidic global deprotection, the macrocyclic strain energy leads to opening of the E ring. We therefore decided to reinvestigate our design strategy.

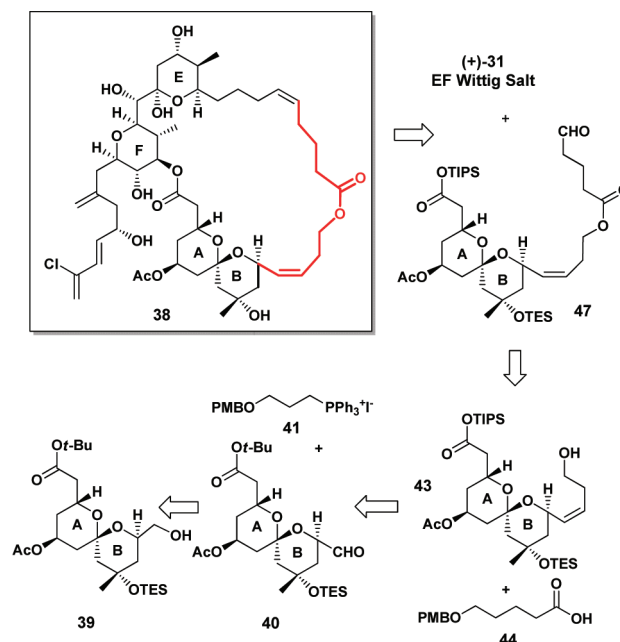
ADVANCED ANALOGUE DESIGN: LESSONS LEARNED LEAD TO A (+)-SPONGISTATIN CONGENER POSSESSING NANOMOLAR TUMOR CELL GROWTH INHIBITORY ACTIVITY

The design and synthesis of biaryl ether EF analogue 17-lactol led to an important lesson: consideration of ring strain energy is critical in the design of macrocyclic analogues. Although a linker may fix the overall conformation, the induced ring strain associated with the designed macrocycle may not be compatible. To rectify this issue, we turned to more flexible polymethylene tethers (Scheme 10). We also chose to incorporate an internal hydrogen bond acceptor in the form of a lactone carbonyl that might participate in a H-bond to the C(42)-hydroxyl on the F ring, in an effort to orchestrate the conformation of the (+)-spongistatin western binding/recognition domain. Our goal was to lower the overall flexibility observed in both our solution conformational studies of (+)-1 and the Heathcock cyclic analogues (15 and 16, Figure 1). Finally, we chose to include the AB spiro-ring moiety to expand the structural elements in an attempt to maximize tubulin binding.

Iterative *in silico* conformational searches, employing a series of tethers of different lengths with hydrogen bond acceptors, led to structure 38 (Scheme 10, red), possessing a tether comprising a Z-olefin, in conjunction with a lactone moiety and a series of methylene units to mimic the span between ring B and the replaced CD spiroketal system. Importantly, the strain energy associated with the designed 29-membered diolide 38 (9.1 kJ/mol) was found to be nearly the same as the ring strain energy of (+)-spongistatin 1 (8.3 kJ/mol).

From the synthetic perspective, analogue 38 was envisioned to arise from advanced AB and EF ring intermediates, again

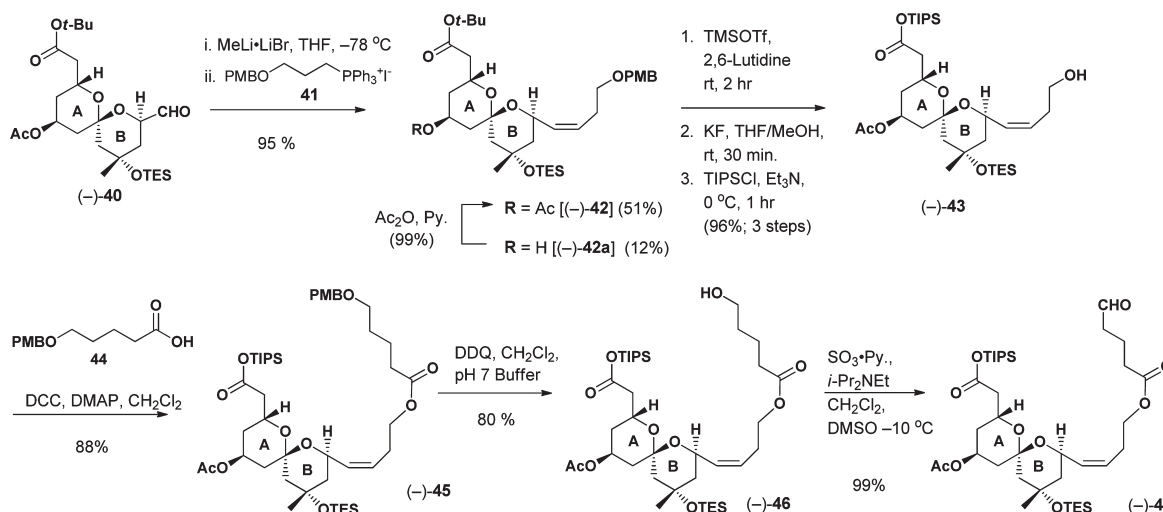
Scheme 10. Retrosynthesis of ABEF Ring Analogue



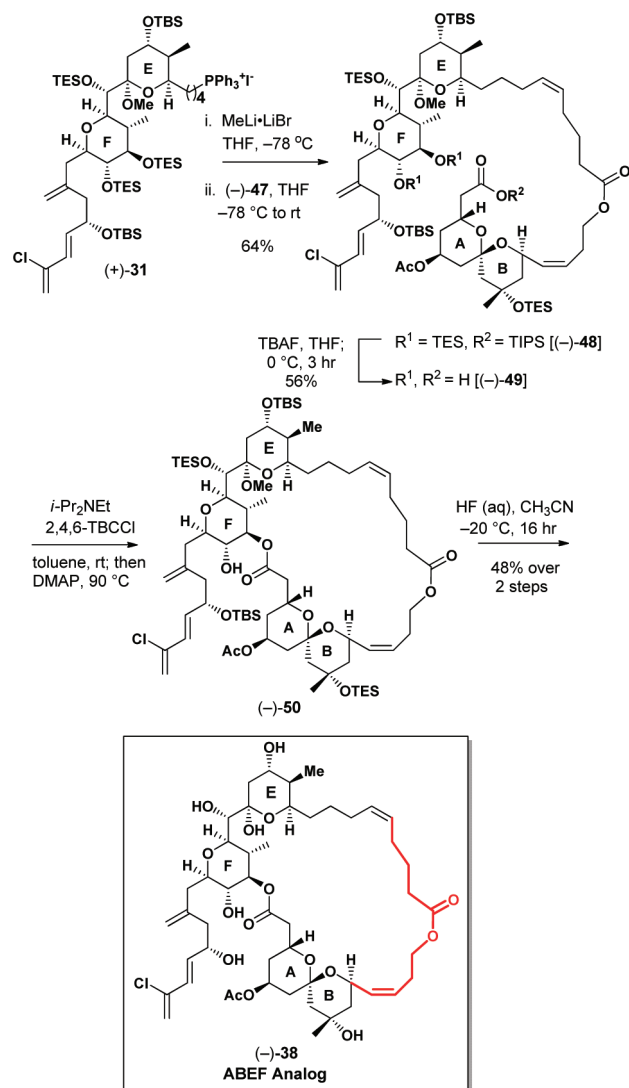
available from our gram-scale synthesis of (+)-spongistatin 1. That is, Wittig union involving EF phosphonium salt (+)-31 and now aldehyde 47, followed by macrolactonization and global deprotection, would lead to 38 (Scheme 10). Tether 47, in turn, would be constructed from three fragments: known AB ring aldehyde 40,³⁹ Wittig salt 41, and primary acid 44,⁴⁰ the latter derived from δ -valerolactone.

Toward this end, Wittig union of (–)-40 with 41 led to (–)-42 in 51% yield, along with the deacetylated congener (–)-42a, which was readily reacylated to furnish (–)-42 (Scheme 11); the overall yield was 63%. Given our earlier experience with (+)-30 (Scheme 4), the *tert*-butyl ester was converted to the TIPS ester by a three-step protocol (ca. 98% yield for the three steps). Known acid 44,³⁸ the remaining component, was then appended to (–)-43 via esterification. Removal of the PMB group followed by oxidation⁴¹ delivered the desired tether (–)-47 in good yield.

Scheme 11. Synthesis of (–)-47



Scheme 12. Preparation ABEF Ring Analogue (–)-38



Union of (–)-47 with EF Wittig salt (+)-31 proceeded smoothly to furnish (–)-48 (Scheme 12) in 64% yield, in this case exclusively as the *Z*-olefin. Removal of the TES and TIPS groups employing TBAF in THF next provided *seco*-acid (–)-49, which upon Yamaguchi lactonization⁴² followed by global deprotection employing HF in CH₃CN—conditions identical to those employed in our large-scale (+)-spongistatin 1 synthesis^{10e}—completed the construction of (–)-38, the ABEF analogue. The yield for the final two steps was 48%.

The solution conformation of (–)-38 was investigated by DISCON computational and NMR studies, similar in detail to those we employed to define the solution conformations of (+)-spongistatin 1. Three solution conformational families were found in CHCl₃ (Figure 5a–c), wherein the western ABEF perimeters of all conformers effectively recapitulate the western ABEF perimeter conformation observed in (+)-1. The major solution conformer (Figure 5a, 59%) incorporates two hydrogen bonds in addition to the H-bonds within the EF system. In accord with our design, the F ring hydroxyl makes a H-bond with the newly introduced lactone carbonyl, as well as the C(10) B ring hydroxyl, further fixing the conformation of the AB ring. The second major conformational family (Figure 5b, 33%)

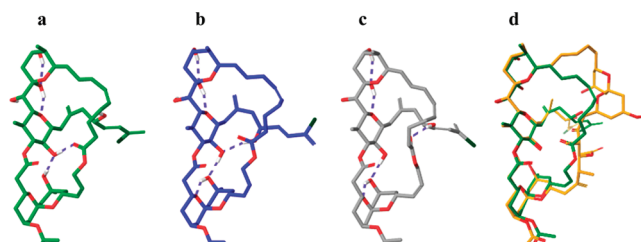


Figure 5. (a–c) Calculated major solution conformers of ABEF analogue (–)-38. (d) Overlay of the major solution conformation a with (+)-spongistatin 1 conformer seen in Figure 4b.

incorporates three additional H-bonds, in this case with the F ring hydroxyl selecting the C(48) hydroxyl [(+)-spongistatin 1 numbering] group on the side chain, rather than with the lactone carbonyl on the linker. Finally, the minor conformer family found (Figure 5c, 8%) incorporates a H-bond between the linker carbonyl and the C(48) [(+)-spongistatin 1 numbering] side-chain hydroxyl. Overall, the conformations are quite rigid at the ABEF ring juncture, giving the side chain a more preferential alignment compared to the (+)-1 structures. Overlay of the three conformations of (–)-38 (Figure 5a–c) with the identified solution conformations of (+)-1 (Figure 4a–d) reveals that they all possess a similar western conformation (Figure 5d).

BIOLOGICAL STUDIES

Not surprisingly, the first-generation EF ring analogue, isolated as a 11:1 mixture of structural isomers (17-Ketone and 17-Lactol) favoring the 17-Ketone, revealed low biological activity in the cell-based growth inhibition assay (Table 1). Pleasingly, however, the ABEF ring analogue (–)-38, demonstrated by NMR to comprise a good conformational mimic of the western perimeter of (+)-1, displayed nanomolar inhibitory activity (Table 3).⁴³ The question that now arose, given that a substantial proportion of the natural product had been deleted, became, *Does this analogue have the same mechanism of action as (+)-spongistatin 1?*

Table 3. Cell Growth Inhibition (IC₅₀, nM) with Spongistatin 1 and the Synthesized Analogues

	MDA-MB-435	HT-29	H522-T1	U937
(+)-1	0.0225	0.058	0.16	0.059
(–)-38	82.8	161.2	297.2	60.5
17	>1000	>1000	>1000	>1000

To answer this question, the effects of (+)-1 and ABEF analogue (–)-38 on cell cycle distribution were evaluated in a standard cell cycle analysis employing U937 lymphoma cells. As shown in Figure 6A, both (+)-spongistatin 1 (1) and analogue (–)-38 led to significant enrichment of cells in the G2/M phase of the cell cycle (G2/M peaks are indicated by arrows), consistent with the known mechanism of action of (+)-spongistatin 1 to target microtubule architecture. To ascertain further that analogue (–)-38 retains the same mechanism of action as (+)-1, both compounds were evaluated in an *in vitro* tubulin polymerization assay. As shown in Figure 6B, analogue (–)-38 inhibits tubulin polymerization in a dose-dependent manner, similar to (+)-1. The higher concentration (i.e., 30 μM) required for analogue (–)-38 to inhibit tubulin polymerization reflects the lower potency of the analogue as compared to the extraordinarily

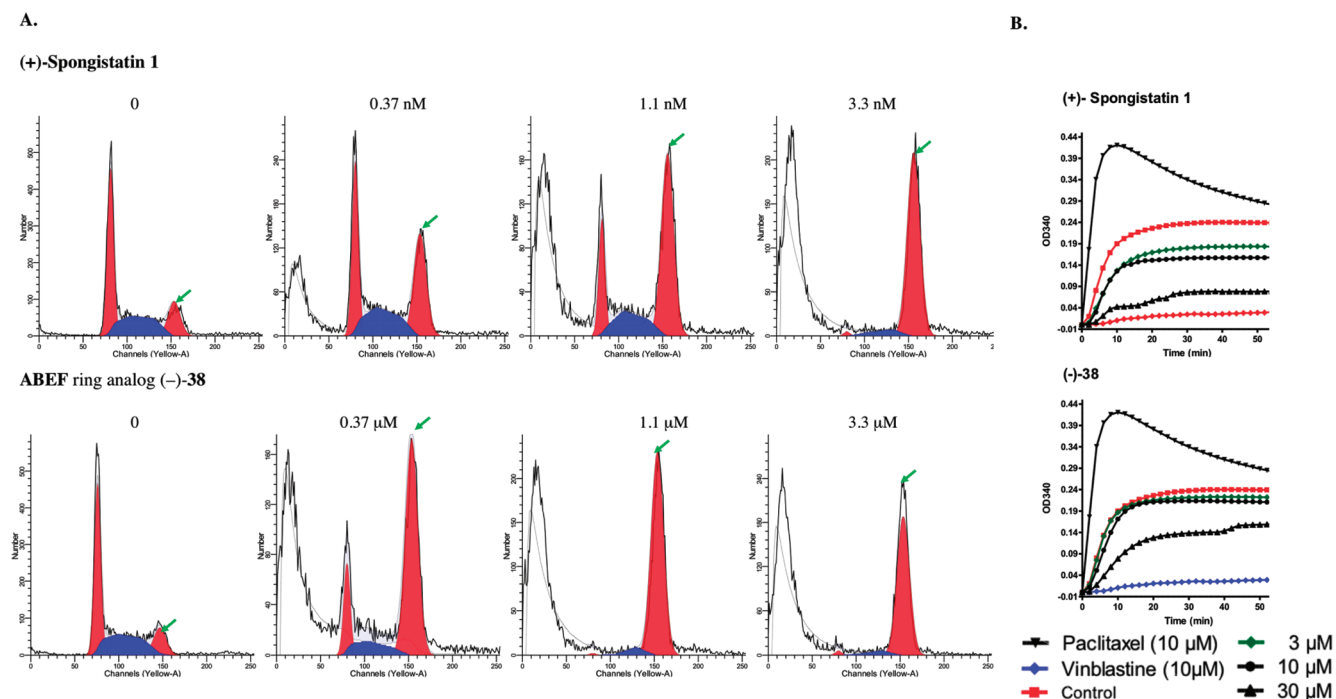


Figure 6. (A) G2/M cell cycle effects of (+)-spongistatin 1 and (–)-38. U937 cells were treated with either (+)-1 or the (–)-38 analogue for 18 h. Samples were subjected to flow cytometric cell cycle analysis. Data shown represent relative number of cells (Y axis) as a function of fluorescence intensity representing DNA content (X axis). (B) Inhibition of *in vitro* tubulin polymerization. An *in vitro* tubulin polymerization assay was carried out in the presence of indicated compounds and concentrations. Tubulin polymerization curves shown here represent the relative amount of tubulin polymer (OD340, Y axis) over time (X axis).

potent (+)-1. Vinblastine, a microtubule destabilizing agent, and paclitaxel, a tubulin stabilizing agent, were included as controls. Thus, like (+)-1, the designed analogue (–)-38 possesses significant microtubule destabilizing activity.

SUMMARY

The feasibility of designing a diminutive congener of (+)-spongistatin 1, (–)-38, wherein approximately one-third of the parent compound is replaced by a simplifying tether that retains significant microtubule destabilizing activity (ca. 60 nM for U937 cell lines), has been demonstrated. To achieve this goal, we introduced a new computational method (DISCON) that led to a design strategy based on the hypothesis that the pharmacophoric elements of biologically active natural products are likely to reside on conformationally rigid portions of the molecule. Significantly, this synthetic effort led to the elimination of some 30 steps compared to the synthesis of (+)-spongistatin 1.¹¹ Finally, a general strategy for the design of simplified bioactive analogues of large, flexible natural products has been developed and validated.

ASSOCIATED CONTENT

S Supporting Information. Spectroscopic and analytical data, selected experimental procedures, and complete ref 22. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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