

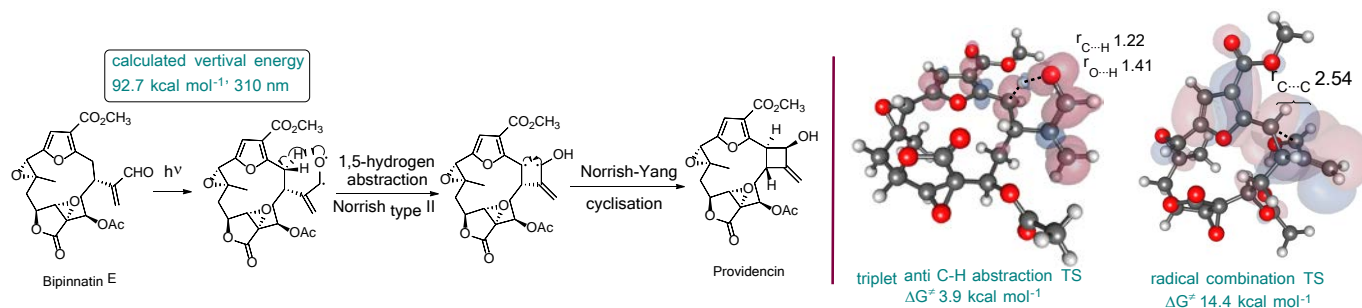
# Biosynthesis of Providencin: Understanding Photochemical Cyclobutane Formation with Density Functional Theory

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Supporting Information Placeholder



**ABSTRACT:** The unique structure of furanocembranoid natural product providencin has stimulated biosynthetic hypotheses, especially concerning the formation of its cyclobutanol ring. One such hypothesis involves a photochemically-induced Norrish-Yang cyclisation in bipinnatin E. We have used computation to assess the feasibility and the stereochemical outcome of this proposed biosynthetic transformation. Density functional theory calculations reveal that the proposed Norrish-Yang cyclisation in bipinnatin E is possible, and that the stereoselectivity of this step is consistent with that of the natural product.

Isolated from Caribbean sea plume *Pseudopterogorgia* *giakallos*, providencin **1** belongs to the family of furanocembranoid natural products, which contain a 14-membered macrocycle, a 2,3,5-trisubstituted furan, and a butenolide ring as common features, e.g. bipinnatin E (**2**) and bipinnatin J (**3**).<sup>1</sup> Compared to other furanocembranes, the most intriguing part of this **diterpenoid** is the methylene-cyclobutanol fused at the C1 and C2-positions. Moreover, the *trans* arrangement of the  $\Delta$ -7,8 epoxide, together with the rigid angle between C2 and C7 appendages around the furan ring, make the macrocycle highly strained. The relative configuration was determined by X-ray crystallography; however the absolute configuration is still unknown.<sup>1</sup> Providencin exhibits modest activity against human breast, lung and CNS cancer cell lines. The biological profile and structural complexity of this marine **diterpenoid** have inspired several synthetic approaches towards the construction of the macrocycle ring as well as the furanyl-cyclobutanol moiety.<sup>2-6</sup> Nevertheless, the total synthesis of providencin has not yet been accomplished. The synthetic challenge lies in its structural features: 1) the highly strained 14-membered macrocyclic structure - it has been observed that it is almost impossible to build a

Dreiding model without breaking any bond;<sup>2</sup> 2) the highly functionalized cyclobutanol moiety, with three contiguous

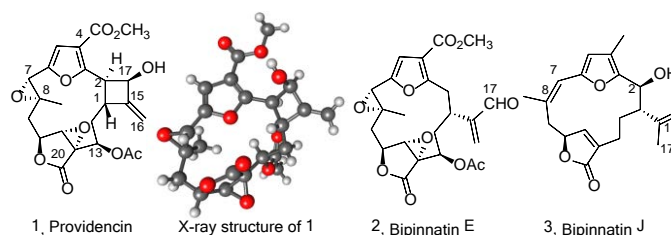


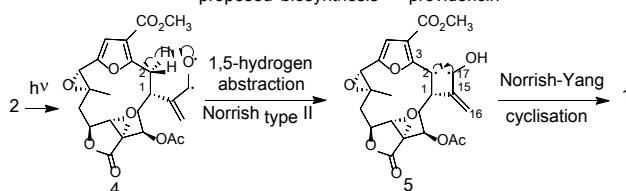
Figure 1. Providencin **1**, bipinnatin E (**2**) and J (**3**)

stereo centers in addition to the exo-methylene substitution.

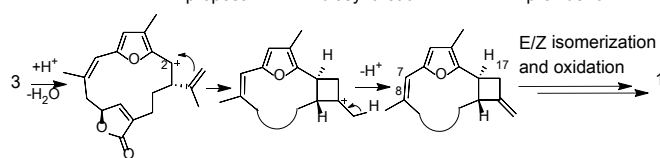
Biosynthetic proposals for the genesis of Providencin from other simple furanocembranes, i.e. bipinnatin J and E, which were also found in *Pseudopterogorgia*,<sup>7</sup> have been reported.<sup>2,6</sup> Pattenden proposed a Norrish-Yang cyclisation<sup>8,9</sup> forms the furan methylene cyclobutanol moiety in Providencin (**Scheme 1a**), based on a photochemically-mediated intramolecular C-H insertion reaction from the  $\alpha$ ,  $\beta$ -unsaturated aldehyde in bipinnatin E.<sup>6</sup> Although the

Norrish-Yang cyclisation is rare in natural product biosynthesis<sup>10–15</sup>, photochemical reactions play an important role in natural product biosynthesis.<sup>16</sup> Many cyclobutanes are thought to arise from photochemical reactions.<sup>17</sup> Mulzer proposed that the cyclobutane ring in providencin arises from a cationic cyclization in C-2 dehydroxylated bipinnatin J, followed by C7-C8 *E/Z* isomerization and further oxidation at C17 to generate the C17 hydroxyl group in providencin.<sup>2</sup>

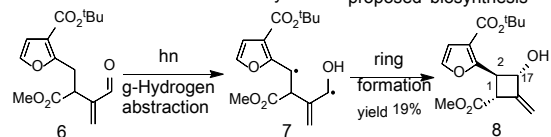
Scheme 1a: Pattenden's proposed biosynthesis of providencin



Scheme 1b. Mulzer's proposal for the biosynthetic formation of providencin



Scheme 2: Pattenden's model study for the proposed biosynthesis of providencin



Pattenden and coworkers performed a synthetic model study of the proposed Norrish-Yang cyclisation in bipinnatin E (**2**).<sup>6</sup> Irradiation of a solution of  $\alpha$ ,  $\beta$ -unsaturated aldehyde **6** in benzene in a pyrex photoreactor using light from a 400 W medium pressure Hg lamp, with regular monitoring by <sup>1</sup>H NMR spectroscopy, led to methylenecyclobutanol **8** through the anticipated Norrish-Yang reaction. The isolated yield was 19%. The conversion of **6** into **8** demonstrates the plausibility of photochemically-mediated cyclobutanol formation as found in the structure of providencin **1**.<sup>6</sup> However, the stereochemistry of **8** has a *trans*-1,2-*trans*-2,17-relationship around the cyclobutane ring, in contrast to the *trans*-1,2-*cis*-2,17-relationship in providencin. It was anticipated that the conformational bias imposed by the macrocyclic ring of the furanocembrane precursor, such as bipinnatin E, may influence diastereoselectivity towards the product, following a photochemical C-H insertion reaction similar to the conversion of **6** to **8**.<sup>6</sup> Therefore, further understanding of the nature of this reaction in the origin of providencin is necessary.

Quantum chemical calculations have been widely used to uncover new mechanisms,<sup>18,19</sup> as well as seeking theoretical support to biosynthetic proposals.<sup>20–22</sup> For instances, calculations have shown the oxidopyriliium-alkene cycloaddition in the biomimetic synthesis of intricarene<sup>23,24</sup> is feasi-

ble, with activation barriers around 20 kcal·mol<sup>-1</sup>.<sup>25</sup> Quantum chemical calculations at CASSCF/CASPT2 and DFT/TD-DFT level of theory also enabled the rationalization of complex reaction pathways involving multiple excited states in the photo-induced intricarene formation.<sup>26</sup> Transannular cycloaddition reactions of furanoxonium ions have been conducted both experimentally<sup>27,28</sup> and investigated computationally with density functional theory (DFT) in furanocembrane biosynthesis.<sup>29</sup> A recent DFT study of cyclobutane formation in Bielschowskysin biosynthesis predicted both thermal closure and photocycloaddition are possible, with the latter being more efficient.<sup>30</sup> Herein we report computational studies to assess the feasibility and stereochemistry outcome of the aforementioned Norrish-Yang cyclisation in the cyclobutanol formation in the biosynthesis of **1**.

**$n\text{-}\pi^*$  Photoexcitation and H-abstraction.** The aforementioned Norrish-Yang reaction is initiated by  $n\text{-}\pi^*$  excitation of the carbonyl group in ketones or aldehydes, *i.e.* the carbonyl group in the  $\alpha$ ,  $\beta$ -unsaturated aldehydes in bipinnatin E (**2**) (Scheme 1a). The key step is the intramolecular 1, 5-H migration from the  $\gamma$ -carbon atom (C2 in **4**) to the oxygen atom of the carbonyl group (Scheme 1a: **2**→**4**→**5**). As a result, a biradical intermediate is formed. This biradical species can then either cyclize to the corresponding cyclobutanol (Yang cyclization, **5**→**1**) or undergo fragmentation (Norrish type II cleavage).<sup>8,9</sup>

Upon photo-excitation, one electron of the oxygen lone pair is promoted to the  $\pi^*$  orbital of the C=O bond. The computed absorption bands for the singlet  $n\text{-}\pi^*$  excitation, calculated by time-dependent (TD)-DFT using CAM-b3lyp/6-311+G(d,p)<sup>31</sup> with a conductor-like polarizable continuum model (CPCM) of water, is 92.7 kcal·mol<sup>-1</sup>, this corresponds to wavelength of 310 nm. The vertical excitation energy is comparable to similar photo-excitations (98.1 kcal·mol<sup>-1</sup>) reported in literature,<sup>32</sup> and the calculated irradiation wavelength is in the range of experiment reported wavelength for aldehyde or ketone  $n\text{-}\pi^*$  excitation. For example, it is reported that aliphatic aldehydes exhibit a weak absorption band in the wavelength range 240–360 nm as a result of a symmetry forbidden  $n\text{-}\pi^*$  transition.<sup>33</sup> The R-band of acrolein is with low density and located in the range of 300–340 nm range.<sup>33,34</sup>

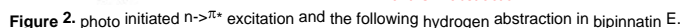


Figure 1 illustrates the reaction coordinate diagram for the C-H abstraction reaction. The diagram shows the energy profile from the reactant state (3TS<sub>1</sub>) to the product state (3TS<sub>2</sub>) and the intermediate state (1Int<sub>6</sub>). The energy difference between 3TS<sub>1</sub> and 3TS<sub>2</sub> is 6.1 kcal/mol. The reaction is labeled "anti C-H abstraction" and "Favored" in red, and "syn C-H abstraction" and "Disfavored" in blue. The diagram includes molecular models of the transition states and the intermediate, showing the C-H bond being abstracted and the C=O bond forming. The energy levels are indicated by the vertical axis, with 3TS<sub>1</sub> at 0 kcal/mol, 3TS<sub>2</sub> at 6.1 kcal/mol, and 1Int<sub>6</sub> at 12.2 kcal/mol.

straction TSs the distance between oxygen atom and the hydrogen atom is 1.41Å and 1.38Å respectively, and the distance between carbon atom and hydrogen atom is 1.22Å and 1.25Å. The resulting biradical intermediate **<sup>3</sup>Int\_4** and

**1Int\_6** and **1Int\_7** (or **1Int\_9**, they differ in the different orientation of C17 hydroxyl group, **Figures 3** and **4**) are different in that, in **1Int\_6**, the C1-C15 bond is rotated to a position where the hydroxyl group can form a hydrogen bond with the C20 carbonyl oxygen of the epoxybutenolide ( $r_{O...H} = 1.99$ ), whereas, in **1Int\_7**, C1-C15 bond is rotated to a position where the same hydroxyl group is pointing away from the epoxybutenolide. Moreover, the presence of the very rigid 14-membered ring system with a *trans*  $\Delta$ -7,8 epoxide may also play a role in the energy difference between **1Int\_6** and **1Int\_7**. As a result, **1Int\_7** has a higher energy by 6.7 kcal·mol<sup>-1</sup>. Due to the steric hindrance in the region and the intramolecular bonding in **Int\_6**, we expect C1-C15 rotation for the interconversion between **Int\_6** and **Int\_7** is unlikely to happen without any significant barrier at room temperature. Therefore, due to the energy differ-

The reaction scheme illustrates the proposed mechanism for the formation of 1,2-diol 10 from 1,2-diol 6. The mechanism proceeds through several intermediates and transition states:

- 1,2-diol 6** (starting material) undergoes **cyclization** via **1TS-4** to form intermediate **1Int\_6**.
- 1Int\_6** undergoes **C-C rotation** to form intermediate **1Int\_8**.
- 1Int\_8** undergoes **cyclization** via **1TS-3** to form intermediate **1Int\_9**.
- 1Int\_9** undergoes **C-C rotation** to form intermediate **1Int\_10**.
- 1Int\_10** undergoes **cyclization** via **1TS-6** to form intermediate **1Int\_11**.
- 1Int\_11** undergoes **cyclization** via **1TS-5** to form the final product, **1,2-diol 10**.

The scheme also shows the direct cyclization of **1,2-diol 6** to **1,2-diol 10** via **1TS-5**.

**Radical recombination.** We set off to understand the four different radical combination reactions starting from singlet biradicals **1Int\_6** and **1Int\_7**. As shown in **Figure 4**, **1Int\_6** is perfectly set for the radical combination to form the desired stereochemistry in Providencin **1** through **1TS\_4**.

the activation Gibbs free energy was calculated as 14.4 kcal·mol<sup>-1</sup>. Alternatively, C15-C17 bond rotation in **1Int\_6** leads to **1Int\_8** (Figure 4), which was found to be the precursor of the C17-epimer of providencin, after overcoming activation free energy of 7.3 kcal·mol<sup>-1</sup> (relative to **1Int\_6**) in the radical combination step through **1TS\_3**. **1Int\_9** (with a free energy of 7.4 kcal mol<sup>-1</sup>) is found to link with a product with an activation free energy barrier of 12.9 kcal mol<sup>-1</sup> of which C2 and C17 centres are opposite to the ones in **1**. Meanwhile, **1Int\_10** leads to a C2 epimer of **1** after radical combination. The activation energy of this process is 6.8 kcal·mol<sup>-1</sup>.

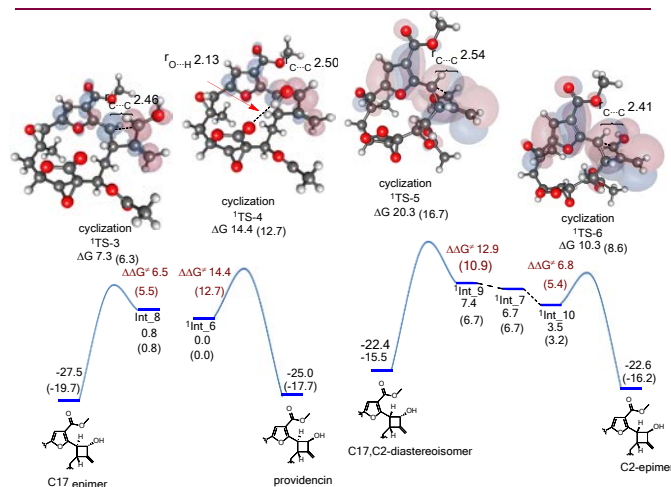


Figure 4. transition state geometry for radical combination step.

Therefore, without considering the barriers in bond rotations in C1-C15 and C15-C17 bonds, radical combination leading to C17-epimer has the lowest barrier. And this, as mentioned above, is indeed the stereochemistry outcome observed in the model study carried by Pattenden group.<sup>6</sup> Therefore, based on these data, it is expected that the radical combination leading to the C17-epimer of providencin has the lowest activation barrier, i.e. 7.3 kcal mol<sup>-1</sup> (relative to **1Int\_6**), hence the C-17 epimer would be the favored product.

At this stage, several questions arise: (a) does the C=C stereochemistry at C7-C8 affect the stereochemical outcome of radical recombination? (b) Could it be possible that in nature, the Norrish-Yang cyclisation reaction happens at an earlier stage, prior to formation of the epoxides on C7-C8 and C11-C12? (c) does the macrocyclic constraint or the substitution pattern on the furan ring affect this reaction? To answer these questions, we investigated the cyclobutene-forming radical ring-closure of four different substrates (Figure 5). These substrates differ in the substitution of the furan ring and the geometry of the C7-C8 double bond. As shown in Table 2 in the supporting information, the TS leading to the C17 epimer (relative to providencin) is most stable for all five different substrates.

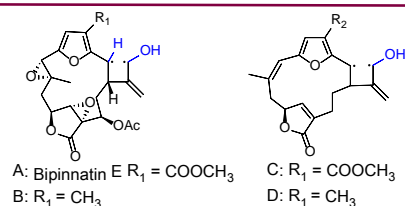


Figure 5. Other biradical substrate studied for radical combination step

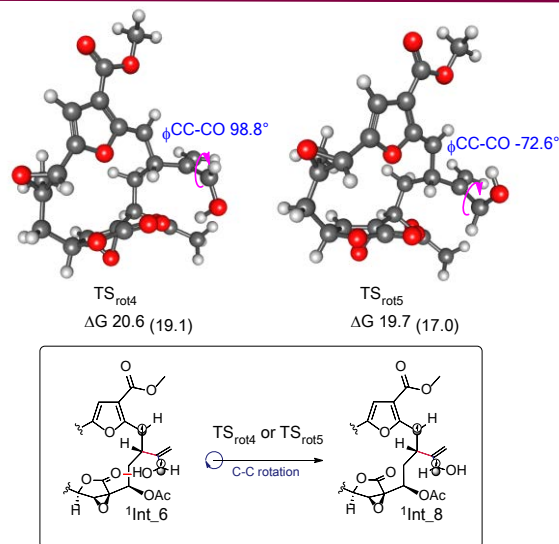
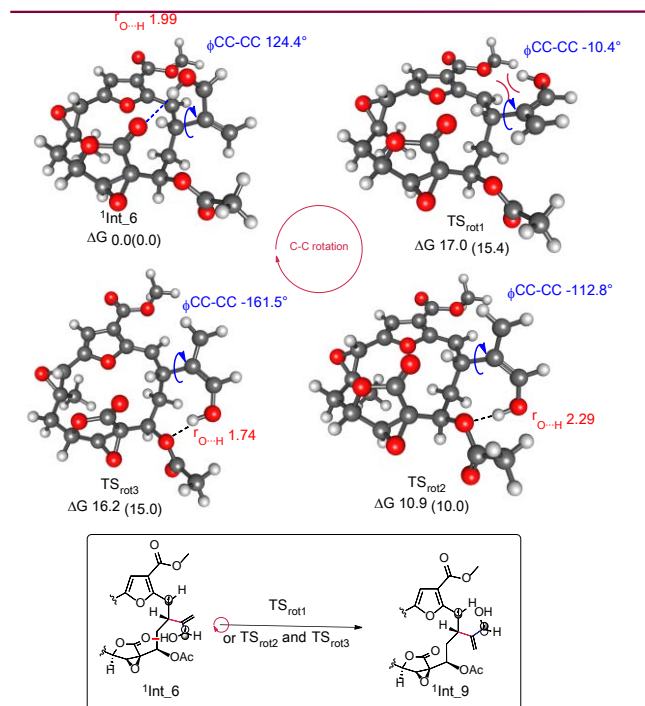


Figure 6. Interconversion of **1Int\_6** and **1Int\_8**

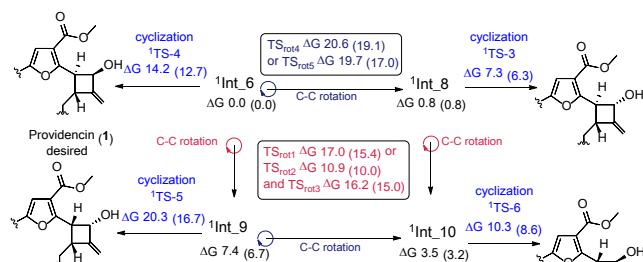
**Interconversion of biradical conformers.** Our discussions up to this point have assumed a Curtin-Hammett scenario in which biradical conformers interconvert more quickly than ring-closure. However, due to steric hindrance around C1-C15 and C15-C17 bonds preventing facile interconversion, we began to question this assumption. For instance, in order to access **Int\_8** from **1Int\_6** the C15-C17 bond needs to rotate (Scheme 3), an intermolecular H-bond must be broken and radical allylic conjugation disrupted. Calculated barriers for rotation in a clockwise or anti-clockwise fashion are 20.6 and 19.7 kcal mol<sup>-1</sup>, respectively. This is higher by 5-6 kcal mol<sup>-1</sup> than ring-closure (Figure 6). Rotation about the C1-C15 bond was also studied (Figure 7). Again, steric congestion and the energetic cost associated with having to break a stabilizing H-bond result in appreciable barriers for the interconversion of **Int\_6** to **1Int\_9**: this can occur in a single step (via **TS\_rot1** with barrier 17.0 kcal·mol<sup>-1</sup>) or sequentially (through **TS\_rot2** and **TS\_rot3** with barriers of 10.9 and 16.2 kcal·mol<sup>-1</sup>). Our calculations therefore suggest that a non Curtin-Hammett scenario can arise, in which **1Int\_6** is the only intermediate to undergo cyclization since conformational interconversion is severely restricted. The C-H abstraction step becomes stereodetermining, and is set by the geometry of **1TS\_1**.





**Figure 7.** Interconversion of  $^1\text{Int}_6$  and  $^1\text{Int}_9$

## Summary



**Scheme 4.** Summary of the computed biosynthesis of providencin

Following photoexcitation, irreversible hydrogen-abstraction in the triplet-state of bipinnatin E favors the formation of biradical intermediate  $^1\text{Int}_6$ . The presence of an intramolecular hydrogen bond and a congested steric environment stabilize this intermediate to such an extent that conformational interconversion via C-C rotation is prohibited and the Curtin-Hammett principle does not govern stereoselectivity. Radical recombination, forming the cyclobutane ring occurs readily, resulting in the correct stereochemistry of providencin. Substrates without the ability to prevent conformational equilibration of the biradical intermediate are predicted to form cyclobutanes epimeric at C-17, in accordance with the Curtin-Hammett principle. At this level of calculation, it is reasonable to conclude that the biosynthetic proposal for the formation of providencin from bipinnatin E through a Norrish-Yang cyclisation reaction is feasible in nature and providencin is the expected

product from the four possible diastereoisomers generated in the radical combination step. A synthetic work towards the synthesis of bipinnatin E for the biomimetic synthesis of providencin using Norrish-Yang cyclisation reaction is also carrying out in our laboratory. Progress will be reported in due course.

## Acknowledgements

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