Abstract:

Hypocrellins and hypomycins are two subclasses of fungal perylenequinones with unique structural, biological, and photochemical properties. With the growing interest in these naturally occurring photosensitizers, more studies were warranted to better understand the structural relationships between these two subclasses of perylenequinones. In this study, the long-postulated biosynthetic precursor (7) of class B fungal perylenequinones was isolated and characterized from a Shiraia-like sp. (strain MSX60519). Furthermore, the electrochemical and chemical redox behaviors of hypocrellins and hypomycins were investigated under aerobic and anaerobic conditions. These studies served to define the structural relationship within hypocrellins (1–3), which was further supported by X-ray crystallography, and between hypocrellins and hypomycins (4–6). Chemical reductions of hypocrellins under anaerobic conditions identified the origin of hypomycin A (4), hypomycin C (5), and hypomycin E (6), which in turn served to confirm 4 and revise the absolute configurations of 5 and 6. Hypocrellins were shown to undergo reversible reduction and reoxidation under aerobic conditions, while in an anaerobic environment and longer time scale, the fully reduced form can, to some extent, undergo an intramolecular ring closing metathesis. This may impart a means of reductive pathway for self-protection against these phototoxins and explain the chemical diversity observed in the fungal metabolites.

Keywords: fungal metabolites | phototoxins | biosynthesis | redox behavior

Article:

***Note: Full text of article below***
Structural Diversity of Perylenequinones Is Driven by Their Redox Behavior

Zeinab Y. Al Subeh, Amy L. Waldbusser, Huzefa A. Raja, Cedric J. Pearce, Kin Lok Ho, Michael J. Hall, Michael R. Probert, Nicholas H. Oberlies, and Shabnam Hematian

ABSTRACT: Hypocrellins and hypomycins are two subclasses of fungal perylenequinones with unique structural, biological, and photochemical properties. With the growing interest in these naturally occurring photosensitizers, more studies were warranted to better understand the structural relationships between these two subclasses of perylenequinones. In this study, the long-postulated biosynthetic precursor (7) of class B fungal perylenequinones was isolated and characterized from a Shiraia-like sp. (strain MSX60519). Furthermore, the electrochemical and chemical redox behaviors of hypocrellins and hypomycins were investigated under aerobic and anaerobic conditions. These studies served to define the structural relationship within hypocrellins (1–3), which was further supported by X-ray crystallography, and between hypocrellins and hypomycins (4–6). Chemical reductions of hypocrellins under anaerobic conditions identified the origin of hypomycin A (4), hypomycin C (5), and hypomycin E (6), which in turn served to confirm 4 and revise the absolute configurations of 5 and 6. Hypocrellins were shown to undergo reversible reduction and reoxidation under aerobic conditions, while in an anaerobic environment and longer time scale, the fully reduced form can, to some extent, undergo an intramolecular ring closing metathesis. This may impart a means of reductive pathway for self-protection against these phototoxins and explain the chemical diversity observed in the fungal metabolites.

INTRODUCTION

Hypocrellins, a subclass of perylenequinones, have gained interest in photodynamic therapy as naturally occurring photosensitizers,1–4 with hypocrellin, its enantiomer (hypocrellin A), hypocrellin B, and shiraiachrome A as the main members of this class of fungal metabolites.5 We recently reported the isolation of ent-shiraiachrome A (1), the shiraiachrome A enantiomer, as a major constituent in the extract of Shiraia-like sp. (strain MSX60519).5,7 In addition, hypocrellin (2), hypocrellin B (3), and three hypomycins [hypomycin A (4), C (5), and E (6)] were isolated from the same fungus (Chart 1).6,7 Hypomycins are structurally similar to hypocrellins but with an additional six-membered ring and a less extended conjugated π-system, which is due to the absence of the Δ1(2) double bond (Chart 1).8,9 The biosynthetic gene clusters responsible for the biosynthesis of some fungal perylenequinones, i.e., cercosporin, hypocrellin A, and elsinochrome C, have been identified.10–12 However, the key biosynthetic steps involved in the generation of hypocrellins, such as the formation of the seven-membered ring, remain unresolved. Moreover, studies on the biosynthesis of hypomycins have never been reported.

In previous work, we also showed that exposure to light was important for the biosynthesis of both hypocrellins and hypomycins, with approximately 70- and 20-fold improvements in the production of hypocrellins (1–2) and hypomycins (4–6), respectively, when the fungi were grown under either continuous exposure to white LED light or under 12/12 h light/dark cycles as compared to complete darkness.6 Additionally, these two structural classes displayed potent cytotoxic activity against human skin melanoma cancer cells (SK-MEL-28) when exposed to light,6 and they are known to exhibit a broad-range of toxicities against bacteria and fungi.3,4,14,15 These interesting properties and findings have raised many questions, especially with respect to the biosynthesis of these perylenequinones. For instance, how does exposure to light not negatively impact the growth of the fungus? In other words, how do hypocrellin-producing fungi...
protect themselves against their own photosensitizers? In addition, is it possible that hypocrellins serve as the biosynthetic precursors for hypomycins, and if so, what are the requirements for converting hypocrellins into hypomycins?

The mechanism used by hypocrellin-producing fungi to avoid self-toxicity has not been reported; however, evidence from fungi that produce the structurally related compound, cercosporin, suggested the ability of fungi to reduce and detoxify this photosensitizer (Figure S1).\textsuperscript{15−17} The reduced form of cercosporin has been shown to be a species with little photosensitizing capacity, resulting in diminished toxicity as compared to the parent cercosporin.\textsuperscript{19−21} Given the structural similarities between hypocrellins and cercosporin (Figure S1), it was conceivable that hypocrellin-producing fungi adapt similar mechanisms to protect themselves against these native photosensitizers.

A goal of this study was to better understand the biosynthetic relationships between hypocrellins and hypomycins via examining the origins of the structural diversity for these two subclasses of perylenequinones. To do so, we investigated their redox activities to probe the potential ability of hypocrellin-producing fungi to reduce and, hence, detoxify these perylenequinones. Here, we present the isolation and characterization of the long-proposed naturally occurring biosynthetic precursor of hypocrellins. Next, physical spectroelectrochemistry, were used to examine the electrochemical reversibility of the two one-electron reduction events of hypocrellins and the absorption profiles of the semiquinone (i.e., one-electron reduced) and hydroquinone (i.e., two-electron reduced) forms of ent-shiraiachrome A (1). In addition, chemical reduction of hypocrellins under aerobic conditions led to a suite of demethylated derivatives. Strikingly, under anaerobic conditions, hypocrellins were converted into hypomycins, and a series of demethoxylated derivatives were also observed. This anaerobic reduction study, along with the \textsuperscript{1}H NMR characterizations, comparisons of the intramolecular hydrogen bonds, and ECD calculations, led to revisions in the structures of hypomycin C (5) and E (6). Finally, this study also provided the first X-ray crystal structures of both ent-shiraiachrome A (1) and hypocrellin B (3).

Identification of the Potential Precursor of Hypocrellins. To secure larger quantities of hypocrellins, particularly ent-shiraiachrome A (1), cultures of strain MSX60519 were grown on rice medium with exposure to white LED light, replicating conditions reported earlier.\textsuperscript{16} The resulting extract (1.7 g) yielded 580 mg of 1, 23 mg of 2, and 0.5−5.0 mg of 3−6. In addition, a benefit of scaling the biosynthesis of fungal metabolites is the identification of other minor constituents,\textsuperscript{22,23} and in this case a new hypocrellin-like compound (7, 1.5 mg) was identified through high resolution electrospray ionization mass spectrometry (HRESIMS) and UV−vis absorption spectroscopy (Figures S2 and S3). The molecular formula of 7, as deduced from the HRESIMS data (measured m/z 547.1583 [M + H]\textsuperscript{+} vs calculated 547.1604), was similar to that reported for 1 and 2 (C\textsubscript{30}H\textsubscript{26}O\textsubscript{10}). However, the \textsuperscript{1}H- and \textsuperscript{13}C(\textsuperscript{1}H)-NMR data revealed only 13 proton and 15 carbon signals, suggesting a symmetrical structure (Table S1 and Figure S4). Examination of the 1D and 2D NMR spectra confirmed the characteristic conjugated pentacyclic core of perylenequinones in 7 (Figures S4−S8). However, the unsymmetrical seven-membered carbocyclic ring was missing.

■ RESULTS AND DISCUSSION

The questions of this study were probed via three interconnected phases. Initially, a scaled-up culture of fungal strain MSX60519 led to the isolation and elucidation of the suggested biosynthetic precursor of hypocrellins. Next, physical techniques, including cyclic voltammetry and UV−vis spectroelectrochemistry, were used to examine the electrochemical reversibility of the two one-electron reduction events of hypocrellins and the absorption profiles of the semiquinone (i.e., one-electron reduced) and hydroquinone (i.e., two-electron reduced) forms of ent-shiraiachrome A (1). In addition, chemical reduction of hypocrellins under aerobic conditions led to a suite of demethylated derivatives. Strikingly, under anaerobic conditions, hypocrellins were converted into hypomycins, and a series of demethoxylated derivatives were also observed. This anaerobic reduction study, along with the \textsuperscript{1}H NMR characterizations, comparisons of the intramolecular hydrogen bonds, and ECD calculations, led to revisions in the structures of hypomycin C (5) and E (6). Finally, this study also provided the first X-ray crystal structures of both ent-shiraiachrome A (1) and hypocrellin B (3).

Chart 1. Structures of the Hypocrellins (1−3) and Hypomycins (4−6) Discussed in This Studya

\begin{align*}
\text{ent-Shiraiachrome A (1)} & \quad & \text{Hypocrellin (2)} & \quad & \text{Hypocrellin B (3)} \\
\text{Hypomycin A (4)} & \quad & \text{Hypomycin C (5)} & \quad & \text{Hypomycin E (6)} \\
\end{align*}

\textsuperscript{a}Note: The absolute configurations of 5 and 6 are revised relative to what was proposed originally.\textsuperscript{6,7}
in 7, being replaced by 2-propanone side chains attached at C-1 and C-12 of the pentacyclic core (Chart 2). This compound was ascribed the trivial name prehypocrellin (7).

We propose that compound 7 represents a common intermediate in the biosynthetic pathway of compounds in the hypocrellin family. The structural similarity between 7 and the synthetic precursor used in the total synthesis of hypocrellin A, as described by O’Brien et al. (Chart 2),24 supports this conclusion. In fact, the structure of 7 was suggested as a plausible biosynthetic intermediate for a variety of fungal perylenequinones,24−27 although it has never been reported from nature. In this context, compound 7 may follow a variety of biotransformation paths to yield the structural diversity seen with perylenequinones, and the isolation of this biosynthetic intermediate further supports what had been hypothesized in the literature, that a common biosynthetic intermediate is responsible for the pentacyclic core and the common substituents for compounds in class B perylenequinones, including cercosporin-like compounds, elsinochromes, and hypocrellins.27 For instance, reduction of the ketone moieties at C-14 and C-17 could result in compounds like calphostins, phleichromes, and cercosporin (Scheme 1).

Alternatively, oxidative enol coupling could generate products like the elsinochromes, while diastereoselective aldol cyclization may generate hypocrellin derivatives (Scheme 1). Interestingly, compound 7 displayed no observable peaks in its Electronic Circular Dichroism (ECD) spectrum (Figure S20). This could be attributed to the presence of a racemic mixture of two possible atropisomers; a similar observation was reported previously for hypocrellin B (3) (Figure S20).6,28,29 In fact, a number of natural and synthetic perylenequinones have been shown to have relatively low barriers for interconversion between the two possible atropisomers. In the case of the hypocrellins (1−3), the additional seven-membered ring attached to the pentacyclic core lowers their barrier of atropisomerization, such that it can be observed at room temperature.9 The isolation of only a relatively small amount of 7 (1.5 mg), even from a scaled-up fungal culture, could be attributed to the high reactivity of this biosynthetic intermediate and its facile conversion into other perylenequinone derivatives.

**Electrochemical Study of Hypocrellins and Hypomycins.** To compare the redox behaviors of both hypocrellins (1−3) and hypomycins (4−6), cyclic voltammetry measurements were conducted in acetonitrile (CH₃CN) containing 100 mM of tetra-n-butylammonium hexafluorophosphate [(nBu)₄N][PF₆] as the supporting electrolyte under rigorous air-free conditions (Figure 1). All potential values are reported against the silver/silver chloride (Ag/AgCl) reference electrode. The voltammograms of the hypocrellin molecules (i.e., 1−3) in aprotic media showed two successive quasi-reversible one-electron reduction processes (Figure 1).

The first reduction event represents the formation of the semiquinone radical anion form (SQ•−) of hypocrellins (1−3) with half-wave potentials (E₁/₂) ranging from −0.46 V to −0.51 V (Figure 1 and Table S2). The second reduction event gave rise to hypocrellins in the dianion (Q²−) state with E₁/₂ ranging from −0.71 V to −0.76 V (Figure 1 and Table S2). Interestingly, the presence of the additional π-bond between

---

**Scheme 1. Compound 7 as a Common Precursor for Class B Perylenequinones**

---

**Chart 2. Structure of Compound 7 (A) as Compared to the Synthetic Precursor Used in the Total Synthesis of Hypocrellin A (B) as Reported by O’Brien et al.**

**A**

**B**

---

**Figure 1**

---

**Table S2**

---

**Notes**

---

**References**

---
single step two-electron two-proton process (e.g., for aqueous acetate buffer). The dianion form of 1, 3 and hypomycins (−E1/2) extended conjugated π-system in the pentacyclic core, their E1/2 values were more negative, ranging from −0.96 V to −1.05 V and −1.29 V to −1.36 V for the first and second reduction events, respectively (Figure 1 and Table S2). Additionally, the absence of the inductive electron-withdrawing effect of a methoxy group at C-2 in hypomycin E (−E1/2: hydroquinone dianion form) resulted in more difficult first- and second-electron reduction processes, as 6 had the most negative E1/2 values, followed by 4 and 5. The peak-to-peak separations (i.e., ΔE1/2 ranging from 60 to 80 mV) and anodic/cathodic peak current ratios (i.e., ipc/ipc = 0.90–1.12) for compounds 1–6 are presented in Table S3. In aqueous acetate buffer at pH = 5, the reduction involved a single step-two-electron-proton process (e.g., for 3: E1/2 = −0.417 V vs Ag/AgCl and ΔE1/2 = 88 mV; Figure S9).

Further characterization for the semiquinone radical anion and hydroquinone dianion forms of ent-shiraiachrome A (1) was performed using UV–vis spectrophotometry. Initially, the UV–vis spectrum for 1 in its starting quinone form, showing maximum absorption in the blue region (λmax = 430 nm) along with two weaker electronic absorptions at 540 and 580 nm (Figure 2). At −650 mV, where the semiquinone radical anion form of 1 would be the most dominant species, the UV–vis spectrum showed a more extended absorption profile in the visible and near-IR (vis–NIR) regions with the maxima in the red region (λmax = 640 nm). The dianion form of 1 was dominant at −850 mV, which showed two strong electronic absorptions at 500 and 530 nm. The reduced forms of ent-shiraiachrome A (1) absorb blue light with lower intensities as compared to the starting quinone form (Figure 2), and their photochemical behaviors (i.e., the nature of the excited electronic states and relaxation mechanisms) are remarkably different. Accordingly, unlike the photoactivated (i.e., excited electronic state) ent-shiraiachrome A (1), which reacts with dioxygen (1O2) through direct energy transfer to generate singlet dioxygen (1O2)6, the reduced form of 1 is expected to react with dioxygen in its ground electronic state through an electron transfer process, in the absence of proton transfer, producing superoxide (O2•−) or peroxide (O2•−2) species. Therefore, a significantly lower phototoxicity of the reduced form of 1, as compared to the starting quinone form, is expected. This further supports our supposition that fungi convert the photoactive forms of these metabolites into the reduced (i.e., nontoxic) forms as a self-protection mechanism. Similar observations were reported for the reduced derivatives of cercosporin17,21, suggesting that this could be a universal trait for these perylenequinones.

Chemical Reduction of Hypocrellins and Hypomycins under Aerobic Conditions. Since the cyclic voltammograms of hypocrellins (1–3) confirmed the electrochemical reversibility of their reduction processes, chemical reductions of these three compounds were performed using sodium dithionite (Na2S2O4, 1 M). As a start, these reactions were conducted at a small scale (~1 μg of each compound) under ambient conditions to maintain an adequate supply of O2. Treating the deep red solutions of 1–3 individually in CH3CN with excess amounts of Na2S2O4 aqueous solution resulted in an immediate color change from deep red to yellow (Figure S10A and B), representing the formation of the reduced hydroquinone forms of 1–3. Subsequently, the reversible reaction of the hydroquinone forms with O2 was observed via the development of the red color as air diffused into the solutions (Figure S10C). This indicated that compounds 1–3 were spontaneously reduced to the yellow hydroquinone forms in the presence of the appropriate reducing agent and were readily oxidized back to the red quinone forms through reacting with O2. The UPLC-HRESIMS data for the reaction mixtures of 1–3 confirmed the recovery of the original compounds in the quinone form, rather than the hydroquinone form (Figures S11–S13). In addition, a series of other side products were produced during these redox interconversions. The molecular formulas of these products, as deduced by HRESIMS, indicated the loss of one and/or two of the methyl
groups in the structures of 1–3 (Figures S11–S13). To further characterize these derivatives, scaled-up reduction reactions of 10 mg of ent-shiraiachrome A (1) with Na₂S₂O₄ were conducted in four replicates (total of 40 mg). Subsequently, the reaction mixtures were subjected to UPLC-HRESIMS, where the four reactions showed the same side products observed in the small-scale reduction reaction of 1 (Figure S14). The reaction mixtures were then combined and subjected to preparative HPLC to obtain compounds 8–10, in addition to the recovery of 1 in the quinone form (Scheme 2). The structures of 8–10 were characterized by analyzing HRESIMS data and 1D and 2D NMR spectra (Table S4 and Figures S15–S18). Accordingly, the methoxy groups attached to C-11 in 8, C-2 in 9, and both C-11 and C-2 in 10 were replaced with hydroxy moieties. Compounds 8, 9, and 10 were generated in approximately a 3:3:1 ratio, respectively (Scheme 2). The replacement of one or both methoxy groups at these positions (i.e., C-2 and C-11) by hydroxy groups is in excellent agreement with previous reports on simpler quinone systems that show that these positions were susceptible to nucleophilic substitution. Note that during this process the location of the oxidized half of the perylenequinone structures are shifted from the left side in 1 to the right side of the molecule in 8–10, so that the methoxy groups adjacent to the carbonyl sites.
are replaced by hydroxy moieties. This observation is also consistent with the fact that, during the aerobic reduction of these hypocrellins with a limited proton source (i.e., in the form of water), dioxygen molecules were continuously reduced and converted to hydroxide ions, thus acting as effective nucleophiles. The tautomeric structures of 8−11 were established by examining their HMBC spectra (Figure S18). Compounds 8−10 exhibited the characteristic electronic absorption profile of perylenequinones (Figure S19) and maintained the same axial chirality of ent-shiraiachrome A (1), as noted by their ECD spectra (Figure S21).

Similar experiments with the hypomycins (i.e., 4−6) resulted in no detectable substitution of the methoxy groups, one of which is attached to an sp3 carbon in hypomycins (i.e., C-2). Here, aerobic reduction with Na2S2O4 led to the full recovery of the original hypomycins in their quinone forms (Figures S22−S24). Moreover, as expected, no apparent color change was observed during the reduction process (Figure S25). Indeed, the electronic absorption profiles of hypomycins in the visible region are significantly different than hypocrellins, in that the former are yellow to orange-colored (Figure S26), as opposed to the deep red-color noted with hypocrellins (Figure S10).

Chemical Reduction of Hypocrellins under Anaerobic Conditions. As the aerobic chemical reduction of hypocrellins (1−3) resulted in a series of demethylated analogues as minor products, we investigated this reaction under anaerobic conditions. ent-Shiraiachrome A (1, 10 mg) was treated with excess Na2S2O4 under argon in triplicate (total of 30 mg of 1). UPLC-HRESIMS data of the reaction mixtures confirmed the recovery of 1 in the quinone form as a major product (64%) (Figure S27). Surprisingly, eight other derivatives were also detected, and this was observed consistently across triplicate reactions (Figures 3A and S27). Based on matching the UPLC retention time and HRESIMS data, these products were identified as hypocrellin (2), hypocrellin B (3), hypomycin A (4), hypomycin C (5), and hypomycin E (6) (Figure S28). Three new products were also identified (i.e., 11−13) with molecular formulas of C29H24O9 for both 11 and 12 and C28H22O8 for 13 as deduced by HRESIMS data (Figure 3A). Accordingly, these compounds were missing either one (as in 11 and 12) or two methoxy groups (as in 13) relative to the starting compound (1). The generation of these natural (2−6) and un-natural (11−13) analogues through the anaerobic reduction of 1 suggests that the structural diversity of hypocrellins and hypomycins derives from their redox activities, which in some cases are followed by intramolecular or intermolecular nucleophilic reactions, vide infra.

To facilitate the isolation of the products, the anaerobic reaction was repeated using 50 mg of 1 followed by preparative HPLC to obtain 1−6 and 11−12 (Scheme 3). The structures were confirmed by 1H NMR and ECD spectra as compared to those reported previously (Figure S29).6,7 Aside from the recovered ent-shiraiachrome A (1), the highest yield was for hypomycin E (6, ~11%) followed by hypomycin C (5; ~5%). Hypomycin A (4) was isolated in low yield (~0.5%) from the
anaerobic reduction of 1 (Figure 3A). This was somewhat surprising, as hypomycin A (4) showed the highest abundance in the extract of strain MSX60519 as compared to 5 or 6. This led us to propose that hypomycin A (4) is a derivative of hypocrellin (2), rather than ent-shiraiachrome A (1), particularly since the axial chirality of 4 matches that of 2 (i.e., M; Chart 1) rather than that of 1 (i.e., P; Chart 1). In this context, we proposed that the reduction of 1 led to the generation of 2, which in turn underwent anaerobic reduction to provide hypomycin A (4).

To test this hypothesis, 4.6 mg of pure hypocrellin (2) were reduced using Na₂S₂O₄ under anaerobic conditions, and the UPLC-HRESIMS data of the reaction mixture showed the production of hypomycin A (4) in a relatively high yield (~65%), in addition to the recovery of 2 in the quinone form (~11%) (Figures 3B and S30). Interestingly, the yield of this intramolecular cyclization under anaerobic conditions to produce hypomycin A (4) from hypocrellin (2) was markedly higher as compared to the relatively low yield of hypomycin C (5) and E (6) from the anaerobic reduction of ent-shiraiachrome A (1) (i.e., 5% and 11%, respectively). This could be attributed to the stronger intramolecular hydrogen bonding present between HO-14 and the C-17 carbonyl in 2, as evidenced by the donor-acceptor distance of 2.70 Å in 2 compared to 2.79 Å in 1 (Figure 4). This would increase the electrophilicity, and hence the reactivity, of the C-17 carbonyl in 2 to produce relatively larger amounts of 4. These data correlated well with our previous findings, where the isolated amount of hypomycin A (4) was significantly higher than those of hypomycin C (5) and E (6) from fungal strain MSX60519.5

**Figure 4.** Intramolecular hydrogen-bonding between the C-14 hydroxy and the C-17 carbonyl groups in the 3D structures of ent-shiraiachrome A (1) and hypocrellin (2). The donor-acceptor distance of 2.70 Å in 2 and 2.79 Å in 1 indicates stronger hydrogen-bonding in hypocrellin (2). The 3D molecular structures and donor-acceptor distances of the intramolecular hydrogen bonding interactions were obtained through Spartan’10 (version 1.1.0) similarity analysis.

**Scheme 4.** Keto-Enol Tautomerism of ent-Shiraiachrome A (1) at C-17 Carbonyl Intermediates the Conversion Process of 1 into 2
Therefore, we postulate that the hypomycins (4–6) could derive from the hypocrellins, meaning that their production requires the reduction of the appropriate hypocrellins under anaerobic conditions. Indirectly, this indicates the ability of fungi to reduce various hypocrellins under oxygen-free conditions to generate a variety of hypomycins; a finding that is consistent with reports of oxidoreductases in diverse groups of fungi. More recently, the sporocarp environment in some fungi has been reported to be anoxic, and this includes a thick outer covering to protect the oxygen-sensitive nitrogenase enzyme, indicating the ability of fungi to create anaerobic environments despite the fact that they live in aerobic habitats. Other evidence from the literature that supports this conclusion is the recent discovery of hypomycin A enantiomer, (P)-hypomycin A, from the stromata extract of *Hypocrella bambusae*, where hypomycin A (i.e., the hypomycin enantiomer) was the major constituent in the extract. Accordingly, to satisfy the absolute configurations among related molecules, hypomycin A (4) with an absolute configuration of M(R), 1S, 2R, 14R, 15S, 17R would derive from hypocrellin (2) with an absolute configuration of M(R), 14R, 15S, while (P)-hypomycin A (enantiomer of 4) with an absolute configuration P(S), 1R, 2S, 14S, 15R, 17S would derive from hypocrellin A (enantiomer of 2) with an absolute configuration of P(S), 14S, 15R.

The conversion of ent-shiraiachrome A (1) to hypocrellin (2) was an interesting observation, as this required flipping the axial chirality from P(S) in 1 to M(R) in 2, along with changing the configuration at C-15 from 1S in 1 to 1S in 2 (Chart 1). From our 1H NMR measurements, we believe that the intramolecular hydrogen bonding between the C-17 carbonyl and C-14 hydroxyl maintains the structures of 1 and 2 in the keto form at C-17 (Figure 4) in solution, and the presence of this intramolecular hydrogen bonding interaction in the solid state was also previously confirmed in the X-ray crystal structure of 2. Accordingly, the 1SR configuration goes with the P(S) chirality, as seen in 1, while the 1SS configuration goes with the M(R) chirality, as seen in 2.

In theory, conversion of 1 into 2 can be achieved through keto–enol tautomerism at the C-17 carbonyl (Scheme 4). The enol form of 1 could be favored if the hydrogen bonding between the C-17 carbonyl and C-14 hydroxy groups was disturbed, where the double bond in the enol form would be stabilized by engaging with the conjugated system of the pentacyclic core (Scheme 4). Therefore, the keto–enol tautomeration process at C-17 engenders the formation of the 1SR stereocenter along with P(S) chirality, as in ent-shiraiachrome A (1), or the formation of the 1SS stereocenter along with M(R) chirality, as in hypocrellin (2). To investigate this possibility, DMSO-d6 was used as a hydrogen bond acceptor to dissolve a sample of ent-shiraiachrome A (1). The 1H NMR spectrum of 1 in DMSO-d6 showed broadened signals, indicating the disruption of the intramolecular hydrogen bond and the presence of a sufficiently fast equilibrium between the keto- and the enol-forms, exceeding the NMR time scale and preventing the observation of separate resonances (Figure S31). After drying the sample and recollecting the 1H NMR data in CDCl3, two sets of signals appeared in the spectrum. One set matched that of ent-shiraiachrome A (1), while the other set matched the 1H NMR profile of hypocrellin (2) (Figure S32). This shows that hypocrellin (2) can derive from 1 simply by exposure to a solvent that is a strong hydrogen bond acceptor (e.g., DMSO), which facilitates the keto–enol tautomerism at the C-17 carbonyl, and thus, the reduction of 1 is not an essential step in the conversion of 1 into 2. Moreover, this suggests that a similar keto–enol tautomerism at C-17 occurs between hypocrellin A (the enantiomer of 2) and shiraiachrome A (the enantiomer of 1), as evidenced by their concurrent isolation from various fungi.

Despite the isolation of hypocrellin B (3) from various fungi as a naturally occurring perylenequinone, it is also known to be a dehydrogenation product of other hypocrellins; for example, hypocrellin B (3) can be produced readily by simple dehydration of either hypocrellin (2), hypocrellin A, shiraiachrome A, or ent-shiraiachrome A (1). The absence of ECD activity for 3 could result from its planar structure or, more likely, from a racemic mixture of the M(R) and P(S) enantiomers of 3. Following successful single crystal growth via ENaCl, we report the first X-ray crystallographic structure of 3, where a racemic mixture of the M(R) and P(S) enantiomers was observed in the solid state (Figures 5 and S33–S34). Accordingly, compound 3 exhibits a low barrier of atropisomerization, which can be observed at room temperature. The fact that hypocrellin B (3) exists as a racemic mixture of the two possible atropisomers further supports our theory that the intramolecular hydrogen bond interaction exhibits a chiral induction effect and is a major factor in determining the favored axial chirality of ent-shiraiachrome A (1) and hypocrellin (2), since losing that intramolecular force, as in 3, leads to a racemic mixture of the two atropisomers.

The conversion of hypocrellins (1 or 2) into hypomycins (4–6) requires the reduction of hypocrellins under oxygen-free conditions, since the production of hypomycins was not observed under any other conditions (i.e., exposure to DMSO, dehydration reactions, or reduction under aerobic conditions). Although the reduction reactions were conducted in oxygen-free settings, UPLC-MS data and product purifications were performed after drying the reaction mixture under an anaerobic environment but then exposing it to normal room conditions. Thus, a question that emerged was whether hypomycins formed before or after exposing the reaction mixture to dioxygen. To probe this, 1H NMR data for the reaction products of 1 and 2 were collected in deuterated CH3CN before air exposure (Figures S35 and S36). Interestingly, the main set of peaks observed in the reduction reaction of ent-
Scheme 5. Structural Relationships between the Natural (1−7) and Non-natural (8−13) Hypocrellin Derivatives

In our previous work, the absolute configuration of hypocrellin C (5) was assigned as $P(S)$, 1S, 2R, 14S, 15S, 17R, and for hypocrellin E (6) it was assigned as $P(S)$, 1S, 14S, 15S, 17R. These conclusions were based on their ECD spectra, analysis of NOESY data, and their structural similarity to hypomycin A (4) (Figure S37),6 which suggested that 5 and 4 differed in both configuration of the C-14 stereogenic center and axial chirality; compound 6 was assigned based on being a demethoxylated derivative of 5.6 However, the anaerobic reduction reactions of 1 and 2 revealed that hypomycin A (4) originates from hypocrellin (2), while hypocrellin C (5) and hypocrellin E (6) are derived from ent-shiraiachrome A (1). This new information caused us to revisit the structure elucidations of 4−6. The absolute configurations of hypomycin A (4) and hypocrellin (2) were in agreement (Chart 1), where both exhibit M(R) chirality and 14R configuration. However, the absolute configurations of hypocrellin C (5) and hypomycin E (6), which likely derive from ent-shiraiachrome A (1), and all share $P(S)$ axial chirality, were in discord at position 14, where 1 is 14R but 5 and 6 were reported as 14S (Figure S37). Since an X-ray crystallographic structure for 1, 5, and 6 has yet to be reported, a suitable single crystal of 1 was prepared via ENaCt and analyzed by X-ray diffraction.26 The resulting crystal structure allowed confirmation of the reported absolute configuration of $P(S)$, 14R, 15R by anomalous dispersion (Flack = 0.019(5)) (Figures S38−S40).6 Unfortunately attempts to grow single crystals of 5 and 6 suitable for X-ray diffraction analysis, using both ENaCt and classical methods, proved unsuccessful.38 Regardless, logic would follow that compounds 5 and 6 should have the 14R configuration, and their NOESY spectra were reanalyzed to evaluate this (Figure S41). As suggested by the NOESY correlations of 5 and 6 (Figure S41), the order of the six-membered ring and the seven-membered ring in 5 and 6 is flipped compared to 4 (Chart 1). The orientation of these two rings and their substituents were further confirmed by examining the observed exchangeable hydroxyl protons in the $^1$H NMR spectra and the presence of intramolecular hydrogen bonding in 4−6 (Figure S42). The $^1$H NMR data of hypomycin A (4) collected in CDCl$_3$ showed the two exchangeable hydrogens at the C-14 chemical shift (δ$_{14}$ 4.81) and the C-17 hydroxy groups (δ$_{17}$ 5.59),6 which suggests that these two protons are stabilized (i.e., slower chemical exchange rate) by forming intramolecular hydrogen bonds (Figure S43). The 3D structure of 4 confirmed the presence of hydrogen bonding interactions between the C-14 and C-17 hydrogen bonds (2.69 Å) (Figure S42). On the other hand, the C-17 hydroxy proton (δ$_{17}$ 4.96), but not the C-14 hydroxy proton, was observed in the $^1$H NMR data for hypomycin C (5) in CDCl$_3$ (Figure S44),6 which indicated that only the C-17 hydroxy group is stabilized through hydrogen bonding with the C-2 methoxy (Figure S44). Such hydrogen bonding between C-17 hydroxy and C-2 methoxy is not possible with the previously suggested structure, as the donor−acceptor distance would be 3.86 Å. Using DMSO-$d_6$ as a hydrogen bond acceptor allowed for

hypomycin A ($\text{Na}_2\text{SO}_4$: reducing agent; $\text{O}_2$: oxidizing agent; $\text{H}_2\text{O}$: proton source)

shiraiachrome A (1) matched those of hypomycin E (6) (Figure S35), while those matching hypomycin C (5) were at lower intensity. The anaerobic reduction reaction of hypocrellin (2) showed peaks that match those of hypomycin A (4) (Figure S36). Accordingly, we suggest that intramolecular cyclization of the hypocrellins to produce hypomycins occurs before exposure to any source of dioxygen.
observing the two exchangeable protons of the C-14 hydroxy ($\delta_2^2$ 4.25) and the C-17 hydroxy groups ($\delta_2^2$ 4.76) in 5 (Figure S43). The absence of a C-2 methoxy in 6 results in losing the hydrogen bonding with the C-17 hydroxy group (Figures S42 and S44). Therefore, neither of the exchangeable hydrogens at the C-14 and C-17 hydroxys were observed in the $^{1}$H NMR of 6 in CDCl$_3$.

Furthermore, the calculated ECD spectrum of the newly proposed structure of 5 showed higher similarity to the experimental data as compared to the previously proposed configuration (i.e., 84% vs 72%), and only the newly proposed [P(S), 1R, 2S, 14R, 15R, 17S]-configuration reproduced the negative Cotton effect at 500 nm (Figure S47). Previously, the appropriate Cotton effect at 500 nm was also observed for the calculated ECD spectrum of hypomycin A (4) only when the correct enantiomeric order was applied. Accordantly, the absolute configurations of 5 and 6 were revised to P(S), 1R, 2S, 14R, 15R, 17S and P(S), 1S, 14R, 15R, 17S, respectively (Chart 1).

Compounds 11 and 12 represent demethoxylated derivatives of 1, where, in each case, one of the four methoxy groups of 1 is missing as deduced by their HRESIMS data (Figure 3A). Analysis of 1D and 2D NMR spectra of 11 confirmed the loss of the methoxy group at C-2 compared to 1 (Table S7 and Figures S48–S50). The extra aromatic proton signal at $\delta_2^2$ 7.25 showed HMBC correlations with C-3, C-3a, C-1a, and C-13, which supported the position of this proton at C-2 in place of their demethylated derivatives (e.g., 3 and 4). The loss of the methoxy group at C-2 compared to 1 has the added benefit of generating wider chemical diversity. Finally, the reductions of hypocrellins under aerobic and anaerobic conditions helped to cement the structural relationships between these compounds, providing biogenetic insights and leading to a revision of the structures of hypomycin C (5) and hypomycin E (6) and the discovery of a series of new analogues (8–12). Moreover, X-ray crystal structures of ent-shiraiachrome A (1) and hypocrellin B (3) were reported for the first time.

### EXPERIMENTAL SECTION

#### General Experimental Procedures.

ECD was measured using an Olis DSM 17 ECD spectrophotometer (Olis, Inc.). UV–vis absorption spectroscopic data were collected using a Cary-60 spectrophotometer (Agilent). Electrochemical data were collected using a Bio-Logic SP-200 potentiostat. 1D and 2D NMR spectra were recorded in either CDCl$_3$, CD$_3$CN, or DMSO–d$_6$. An Agilent 700 MHz spectrometer equipped with a cryoprobe, a JEOL ECA-500 spectrometer, or a JEOL ECS-400 spectrometer equipped with a high sensitivity JEOL Royal probe and a 24-slot autosampler. The NMR shifts were referenced to CDCl$_3$ (δ$_3$ 7.26/77.2), DMSO–d$_6$ (δ$_6$ 2.50), or CD$_3$CN (δ$_3$ 1.94). UPLC-HRESIMS data were collected via an LTQ-Orbitrap XL mass spectrometer system (Thermo Finnigan, San Jose, CA, USA) connected to a Waters Acuity UPLC system. A BEH Shield RP18 column (Waters, 1.7 μm; 50 mm × 2.1 mm) heated to 40 °C was used. The mobile phase consisted of CH$_3$CN–H$_2$O (0.1% formic acid) in a gradient system of 15:85 to 100:0 over 10 min at a flow rate of 0.3 mL/min. MS data were collected from m/z 150 to 2000 in the positive mode. All analytical and preparative HPLC experiments were carried out using a Varian Prostar HPLC system equipped with ProStar 210 pumps and a Prostar 335 photodiode array detector (PDA). HPLC data were collected and analyzed using Galaxy Chromatography Workstation software (version 1.9.3.2, Varian Inc.). For preparative HPLC, a Synergi C12 column (Waters, 5 μm; 250 mm × 21.2 mm) was used. Flash chromatography was carried out using a Teledyne Isco CombiFlash RF 200 that was equipped with both UV and evaporative light-scattering detectors and using Silica Gold columns (from Teledyne Isco).

### CONCLUSION

This study expanded our knowledge on the biosynthesis and the redox behavior of hypocrellins and hypomycins, bringing to light four new aspects to our understanding of these perylenequinones. First, the biosynthetic precursor of hypocrellins was isolated and characterized, giving tangible evidence to the ultimate precursor to not only this class of compounds but also, by analogy, to the calphostins, phiechlorines, cercosporins, and elsinochromes. Second, our electrochemical measurements revealed that the hypocrellins and hypomycins undergo reversible reduction processes, with the former being more thermodynamically favored (i.e., milder redox potentials), which was further supported with our aerobic chemical reduction studies. Third, hypomycins originate from reduction of hypocrellins under anaerobic conditions, and the presence of significant amounts of hypomycins in fungal culture extracts suggests that hypocrellin-producing fungi reduce these photoactive compounds, potentially as a self-protection mechanism; this has the added benefit of generating wider chemical diversity. Finally, the reductions of hypocrellins under aerobic and anaerobic conditions helped to cement the structural relationships between these compounds, providing biogenetic insights.
All reagents and solvents were of commercially available grade, unless otherwise noted. Acetonitrile (MeCN) was used after passing through a 60-cm-long column of activated alumina (Innovative Technologies) under argon. Distilled water was further purified by a Nanopure Analytical Ultrapure Water System (Thermo Scientific) to obtain the specific resistance of 18.2 MΩ cm at 25 °C. Deoxygenation of solvents was achieved by either repeated freeze–pump–thaw cycles or bubbling with argon for 40–60 min. Sodium dithionite (for analysis EMSURE) was purchased from Millipore Sigma. 

**Media and Fermentations.** Fungus MSX60519 was identified as a *Shiria*-like fungus in the family Shiriaeeaceae, as described recently, and was maintained on potato dextrose agar (PDA; Difco) and transferred periodically to fresh PDA Petri plates. An agar plug from the leading edge of the PDA culture was transferred to a sterile tube with 10 mL of YESD (2% soy peptone, 2% dextrose, and 1% yeast extract). The YESD culture was grown for 7 days on an orbital shaker (100 rpm) at room temperature (∼23 °C) and then used to inoculate solid fermentation media.

Twelve cultures of strain MSX60519 were grown in rice medium in 250 mL Erlenmeyer flasks as previously described. To prepare rice medium, 10 g of rice were added to each flask with 20 mL of deionized water. After autoclaving, the flasks were inoculated with YESD seed cultures (described above) and incubated at room temperature (25 °C) and with gentle shaking (∼80 rpm) for 7 days. After extraction, the YESD culture was transferred to a sterile tube with 250 mL Erlenmeyer flasks, and then transferred into a separatory funnel. The bottom layer was drawn off, evaporated to dryness, and then reconstituted in 100 mL of 1:1 CH₃OH–CHCl₃. The extract was vacuum filtered, and 90 mL of CHCl₃ and 100 mL of DI H₂O were added to the filtrate. The mixture was stirred for 30 min and then transferred into a separatory funnel. The bottom layer was then evaporated under vacuum. The CH₃OH/CHCl₃ layer was then evaporated under vacuum. The organic extracts collected from the 12 cultures were combined to provide a total of 2.2 g. These were dissolved in CHCl₃, adsorbed onto Celite 545, and subdivided into five fractions via normal-phase flash chromatography using a gradient solvent system of hexanes–CHCl₃–CH₂OH at a 35 mL/min flow rate and 41 column volumes over 40 min. The first and the second flash chromatography fraction of each extract were subjected to preparative HPLC separation using a Phenomenex Synergi C₄ preparative column and a gradient system of 50:50 to 100% of CH₃CN and then treated with 3 mL of Na₂S₂O₄ solution (960 mM). The reaction mixtures were stirred for 3 h and then left open overnight. After 24 h, the four reaction mixtures were dried and subjected to UPLC-HRMS analysis as described above. To purify the resulting products, the four reactions were combined and subjected to a preparative reversed-phase HPLC separation using a Phenomenex Synergi C₁₂ preparative column and a gradient system of 50:50 to 60:40 CH₃CN–H₂O (0.1% formic acid) over 15 min at a flow rate of 21.24 mL/min to yield ent-shiraiachrome A (1, 580 mg), hypocrellin B (2, 23 mg), hypocrellin B (3, 1.4 mg), and hypymycin A (4, 4.6 mg). The third flash chromatography fraction was subjected to prep-HPLC using the same protocol to obtain hypymycin C (5, 1.3 mg), hypymycin E (6, 0.5 mg), and prehypocrellin (7, 1.5 mg).

**Prehypocrellin (7).** Dark red amorphous powder; UV (CH₃OH) λmax (log ε) 576 (3.74), 470 (4.02), 254 (4.31), 213 (4.46) nm; 1H NMR (500 MHz, CDCl₃) and 13C{1H} NMR (125 MHz, CDCl₃) (see Table S4); HRMS (ESI) m/z: [M + H]+ calcld for C₂₈H₂₃O₁₀ 519.1291; Found 519.1298. ECD (MeOH) nm (Δε): 707 (+4.5), 315 (−4.5), 455 (−4.3), 567 (−4.0).

**Cyclic Voltammetry Measurements.** A three-electrode setup was used for all voltammetry experiments with a 3-mm glassy carbon disk working electrode, a carbon rod counter electrode, and a leak-free Ag/AgCl reference electrode (3.4 M KCl) inside an OMNI-Lab inert atmosphere (<0.5 ppm of O₂ and H₂O) glovebox filled with nitrogen. The potentials were referenced to the Ag/AgCl electrode. All electrodes were cleaned with acetone and nanopure water before and after use. Acetonitrile was further deoxygenated by bubbling with argon for 45–60 min followed by storage over 3 Å molecular sieves for at least 72 h prior to use. The electrolyte tetra-n-butylammonium hexafluorophosphate (98%), [(nBu₄N)₂][PF₆], was purchased from Oakwood Chemicals and further purified by recrystallization from ethanol. OmniPur sodium acetate, trihydrate, and acetic acid (glacial; 100%) were purchased from Millipore Sigma.

**Spectroelectrochemistry.** A three-electrode setup was used for spectroelectrochemistry experiments with a platinum mesh working electrode, a platinum wire counter electrode, and a leak-free Ag/AgCl reference electrode (3.4 M KCl). The solutions were made inside the glovebox and transferred into a custom-made 1 mm Schlenk cuvette designed specifically for air-free spectroelectrochemistry experiments. The cuvette was then sealed and brought outside of the glovebox to obtain the UV–vis instrument. First, a blank was recorded which consisted of 100 mM of the supporting electrolyte, [(nBu₄N)₂][PF₆] in solution without the electrodes present. Then, a solution containing both electrolyte (100 mM) and compound 1 (248 μM) was added to the cuvette and the UV–vis spectrum was recorded. The electrodes were then inserted into the solution and connected to the potentiostat, and another spectrum was recorded, demonstrating a slight increase in the baseline due to the blockage of the working electrode. This solution was then used for a blank, and subsequent UV–vis difference measurements were recorded as a negative potential was applied to the working electrode. The starting potential was −0.3 V and was decreased by 0.05 V when the UV–vis spectra were unchanged. The final potential applied was −0.85 V.

**Small-Scale Reduction of Hypocrellins (1–3) and Hypymycins (4–6) with Sodium Dithionite.** Chemical reductions of compounds 1–6 using Na₂S₂O₄ as a reducing agent were first conducted at a pilot scale, where 1 mg of ent-shiraiachrome A (1), hypocrellin (2), and hypocrellin B (3) was placed in a scintillation vial and dissolved separately in 1 mL of acetonitrile. Each solution was then treated with excess Na₂S₂O₄ by adding 0.6 mL of aqueous Na₂S₂O₄ solution (480 mM) and mixing thoroughly. Similar procedures were followed for the chemical reductions of hypymycin A (4), C (5), and E (6), but using 0.5 mg of each compound. The six reaction vials were left open at room conditions overnight to provide a regular supply of dioxygen. After 24 h, UPLC-HRMS data were collected for the reaction mixtures of 1–6 as well as for the starting materials, i.e., pure samples of 1–6 dissolved in acetonitrile.

**Scaled-up Reduction of ent-Shiraiachrome A (1) under Aerobic Condition.** The scaled up aerobic reduction of ent-shiraiachrome A (1) was conducted in four replicates. For each reaction, 10 mg of 1 dissolved in 2.5 mL of CH₃CN and then treated with 3 mL of Na₂S₂O₄ solution (960 mM). The reaction mixtures were stirred for 3 h and then left open overnight. After 24 h, the four reaction mixtures were dried and subjected to UPLC-HRMS analysis as described above. To purify the resulting products, the four reactions were combined and subjected to a preparative reversed-phase HPLC separation using a Phenomenex Synergi C₁₂ preparative column and a gradient system of 50:50 to 60:40 CH₃CN–H₂O (0.1% formic acid) over 15 min at a flow rate of 21 mL/min to yield compounds 1 (23.6 mg), 8 (1.8 mg), 9 (2.5 mg), and 10 (0.6 mg).

11-O-Demethyl-ent-shiraiachrome A (8). Dark red amorphous powder; 1H NMR (400 MHz, CDCl₃) and 13C{1H} NMR (100 MHz, CDCl₃) (see Table S4); HRMS (ESI) m/z: [M + H]+ calcld for C₃₀H₂₇O₁₀ 533.1447; Found 533.1424. ECD (MeOH) nm (Δε): 270 (+4.5), 350 (−6.0), 455 (+12.4), 563 (−4.0).

2-O-Demethyl-ent-shiraiachrome A (9). Dark red amorphous powder; 1H NMR (500 MHz, CDCl₃) and 13C{1H} NMR (125 MHz, CDCl₃) (see Table S4); HRMS (ESI) m/z: [M + H]+ calcld for C₂₉H₂₅O₁₀ 547.1604; Found 547.1583. ECD (MeOH) nm (Δε): 280 (+5.5), 340 (−1.6), 410 (+5.3), 463 (+4.0), 573 (−0.3).

2,11-Di-O-Demethyl-ent-shiraiachrome A (10). Dark red amorphous powder; 1H NMR (700 MHz, CDCl₃) and 13C{1H} NMR (175 MHz, CDCl₃) (see Table S4); HRMS (ESI) m/z: [M + H]+ calcld for C₂₈H₂₄O₁₀ 519.1291; Found 519.1298. ECD (MeOH) nm (Δε): 238 (−5.6), 300 (+1.73), 348 (−5.0), 438 (+4.1), 558 (−1.3).

**Reduction of ent-Shiraiachrome A (1) and Hypocrellin (2) under Anaerobic Condition.** Anaerobic reduction of 1 was performed in triplicate, where 10 mg of 1 were used in each reaction (total of 30 mg). Under a dry oxygen-free argon atmosphere using a standard Schlenk technique, each sample of 1 was dissolved in 2.5 mL of CH₃CN and then treated with 3 mL of aqueous Na₂S₂O₄ solution (960 mM). The reactions were stirred under oxygen-free conditions.
for 30 min before drying in vacuo. After 24 h, samples for NMR analysis were prepared in an oxygen-free glovebox. Afterward, UPLC-HRESIMS data were collected for each reaction. Reference standards of 1–6 were also analyzed by UPLC-HRESIMS to identify the reaction products by matching their molecular ion peaks and retention times. For the purpose of purifying the observed products (i.e., compounds 2–6 and 11–12 in Scheme 3), the anaerobic reduction reaction of 1 was scaled up by reacting 50 mg of 1 following the same procedures. The reaction mixture was then subjected to a preparative reversed-phase HPLC separation using a Phenomenex Synergi C12 column and an isotropic system of 60:40 CH3CN-H2O (0.1% formic acid) for 30 min and at a flow rate of 21 mL/min to obtain compounds 1 (31.8 mg), 2 (2.1 mg), 3 (0.2 mg), 5 (2.3 mg), 6 (5.3 mg), 11 (1.6 mg), and 12 (0.5 mg). The anaerobic reduction of hypocrellin (2, 4.6 mg) was conducted following a nearly identical protocol to obtain compounds 2 (0.5 mg) and 4 (2.97 mg).

2-Demethoxy-ent-shiraiachrome A (11). Dark red amorphous powder; 1H NMR (400 MHz, CDCl3) and 13C{1H} NMR (100 MHz, CDCl3) (see Table S7); HRMS (ESI) m/z: [M + H]+ calcd for C26H32O13 678.1592, found 678.1592. 11-Demethoxy-ent-shiraiachrome A (12). Dark red amorphous powder; 1H NMR (700 MHz, CDCl3) and 13C{1H} NMR (175 MHz, CDCl3) (see Table S7); HRMS (ESI) m/z: [M + H]+ calcd for C26H29O12 650.1536, found 650.1534.

Encapsulated Nanodroplet Crystallization (ENaCt) of Hypocrellin B (3). The crystallization of hypocrellin B (3) was examined via the published ENaCt approach.43 Stock solutions of hypocrellin B (3) were prepared in MeOH, aliquoted into screw top glass vials and allowed to evaporate. Samples of hypocrellin B (3) were then dissolved in a range of solvents. Droplets of hypocrellin B (3) containing solutions (50 nL) were dispensed via an STP Labtech Mosquito liquid handling robot into 96 well glass plates (SWISSCI LCP Modular, 100 µl spacer) containing either an appropriate solvent, or solutions (50 nL) containing 10% DMF. The anaerobic reduction of hypocrellin B (3) was conducted following a nearly identical protocol to obtain single crystals suitable for X-ray diffraction analysis (Figure S38). Wells were opened, and crystals were analyzed using techniques and instruments identical to the procedure described above for 3.

Shiraiachrome A (1). Crystallographic data of Shiraiachrome A (1) were deposited at the Cambridge Crystallographic Data Centre (deposition numbers: CCDC 2085795). Crystallographic data of this paper. These data can be obtained emailing data_request@ccdc.cam.ac.uk, or by contacting The Cambridge Crystallographic Data Centre, 12 Union Road, Cambridge CB2 1EZ, UK; fax: +44 1223 336033.

Computational Methods. Molecular Merck force field (MMFF) and density functional theory (TD-DFT/DFT) calculations were carried out with Spartan’10 (Wave function Inc., Irvine, CA, USA) and GaussView 06 software, respectively. The conformers were optimized using DFT calculations at the B3LYP/cc-pVTZ level in CH3OH for ECD prediction of 3. The ECD spectra were generated using SpecDis 1.71 software.44

ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge at https://pubs.acs.org/doi/10.1021/acs.joc.1c02639.

Accession Codes

CCDC 2085795–2085796 contain the supplementary crystallographic data for this paper. These data can be obtained free of charge via www.ccdc.cam.ac.uk/data_request/cif or by emailing data_request@ccdc.cam.ac.uk, or by contacting The Cambridge Crystallographic Data Centre, 12 Union Road, Cambridge CB2 1EZ, UK; fax: +44 1223 336033.

AUTHOR INFORMATION

Corresponding Authors

Nicholas H. Oberlies — Department of Chemistry and Biochemistry, University of North Carolina at Greensboro, Greensboro, North Carolina 27402, United States; orcid.org/0000-0002-0354-8464; Email: nicholas_oberlies@uncg.edu

Shabnam Hematian — Department of Chemistry and Biochemistry, University of North Carolina at Greensboro, Greensboro, North Carolina 27402, United States; orcid.org/0000-0002-0788-7615; Email: s_hemati@uncg.edu

Authors

Zeinab Y. Al Subeh — Department of Chemistry and Biochemistry, University of North Carolina at Greensboro, Greensboro, North Carolina 27402, United States; orcid.org/0000-0001-9051-4969
Notes
The authors declare the following competing financial interest(s): Nicholas Oberlies is on the Scientific Advisory Board of Mycosynthetix, Inc.

ACKNOWLEDGMENTS
This research was supported in part by the National Institutes of Health (US)/National Cancer Institute via Grant P01 CA125066 and by the Medical Research Council (UK) via Grant MR/T000740/1. X-ray crystallography facilities at NCL were supported by the Engineering and Physical Sciences Research Council (UK) via Grant EP/F03637X/1. S.H. and A.L.W. are thankful to the University of North Carolina at Greensboro for the startup funds and the New Faculty Research Award.

REFERENCES
(1) Jiang, Y.; Leung, A. W.; Wang, X.; Zhang, H.; Xu, C. Effect of photodynamic therapy with hypocrellin B on apoptosis, adhesion, and migration of cancer cells. Int. J. Radiat. Biol. 2014, 90, 575.
(2) Qi, S.; Guo, L.; Yan, S.; Lee, R. J.; Yu, S.; Chen, S. Hypocrellin A-based photodynamic action induces apoptosis in A549 cells through ROS-mediated mitochondrial signaling pathway. Acta Pharm. Sin. B 2019, 9, 279.
(3) Yang, Y.; Wang, C.; Zhuge, Y.; Zhang, J.; Xu, K.; Zhang, Q.; Zhang, H.; Chen, H.; Chu, M.; Jia, C. Photodynamic antifungal activity of hypocrellin A against Candida albicans. Front. Microbiol. 2019, 10, 1810.
(4) Su, Y.; Sun, J.; Rao, S.; Cai, Y.; Yang, Y. Photodynamic antimicrobial activity of hypocrellin A. J. Photochem. Photobiol. B, Biol. 2011, 103, 29.
(5) Mazzini, S.; Merlini, L.; Mondelli, R.; Scaglioni, L. Conformation and tautomeration of hypocrellins. Revised structure of shiraiachrome A. J. Chem. Soc., Perkin Trans. 2001, 2, 409.
(6) Al Subeh, Z. Y.; Raja, H. A.; Monro, S.; Flores-Bocanegra, L.; El-Elimat, T.; Pearce, C. J.; McFarland, S. A.; Oberlies, N. H. Enhanced production and anticancer properties of photoactivated perylenequinones. J. Nat. Prod. 2020, 83, 2490.
(7) Al Subeh, Z. Y.; Raja, H. A.; Monro, S.; Flores-Bocanegra, L.; El-Elimat, T.; Pearce, C. J.; McFarland, S. A.; Oberlies, N. H. Correction to enhanced production and anticancer properties of photoactivated perylenequinones. J. Nat. Prod. 2020, 83, 3765.
(8) Liu, W. Z.; Shen, Y. X.; Liu, X. F.; Chen, Y. T.; Xie, J. L. A new perylenequinone from Hypomyces sp. Chin. Chem. Lett. 2001, 12, 431.
(9) Shen, Y. X.; Liu, W. Z.; Rong, X. G.; Sun, Y. H. Studies on the chemical constituents of a fungus producing perylenequinones. Acta Pharm. Sin. 2003, 38, 834.
(10) Newman, A. G.; Townsend, C. A. Molecular characterization of the cescorpin biosynthetic pathway in the fungal plant pathogen Cercospora nicotianae. J. Am. Chem. Soc. 2016, 138, 4219.
(11) Chen, H. Q.; Lee, M. H.; Chung, K. R. Functional characterization of three genes encoding putative oxidoreductases required for cescorpin toxin biosynthesis in the fungi Cercospora nicotianae. Microbiolology 2007, 153, 2781.
(12) Deng, H.; Gao, R.; Liao, X.; Cai, Y. Genome editing in Shiria bambusicola using CRISPR-Cas9 system. J. Biotechnol. 2017, 259, 228.
(13) Chooi, Y. H.; Zhang, G.; Hu, J.; Muria-Gonzalez, M. J.; Tran, P. N.; Pettitt, A.; Maier, A. G.; Barrow, R. A.; Solomon, P. S. Functional genomics-guided discovery of a light-activated phototoxin in the wheat pathogen Parastagonospora nodorum via pathway activation. Environ. Microbiol. 2017, 19, 1975.
(14) Leung, A. W.; Ip, M.; Xu, C. S.; Wang, X. N.; Yung, P. T.; Hua, H. Y. Sonodynamic bactericidal efficacy of hypocrellin A and B against mexiticillin-resistant Staphylococcus aureus. Hong Kong Med. J. 2017, 23, 36.
(15) Li, Y.-T.; Yang, C.; Wu, Y.; Lv, J.-J.; Feng, X.; Tian, X.; Zhou, Z.; Pan, X.; Liu, S.; Tian, L.-W. Axial chiral binaphthoxquinone and perylenequinones from the stromata of Hypocrella bambusae are SARS-CoV-2 entry inhibitors. J. Nat. Prod. 2021, 84, 436.
(16) Sollod, C. C.; Jenns, A. E.; Daub, M. E. Cell surface redox potential as a mechanism of defense against photosensitizers in fungi. Appl. Environ. Microbiol. 1992, 58, 444.
(17) Daub, M. E.; Leisman, G. B.; Clark, R. A.; Bowden, E. F. Reductive detoxification as a mechanism of fungal resistance to singlet oxygen-generating photosensitizers. Proc. Natl. Acad. Sci. U. S. A. 1992, 89, 9588.
(18) Daub, M. E.; Herrero, S.; Chung, K.-R. Photoactivated perylenequinone toxins in fungal pathogenesis of plants. FEMS Microbiol. Lett. 2005, 252, 197.
(19) Daub, M. E.; Ehrenshaft, M. The photovactiated toxin cercosporin as a tool in fungal photobiology. Physiol. Plant. 1993, 89, 227.
(20) Daub, M. E.; Li, M.; Bilski, P.; Chignell, C. F. Dihydrocercosporin singlet oxygen production and subcellular localization: a possible defense against cercosporin phototoxicity in Cercospora. Photochem. Photobiol. 2000, 71, 135.
(21) Leisman, G. B.; Daub, M. E. Singlet oxygen yields, optical properties, and phototoxicity of reduced derivatives of the photosensitizer cercosporin. Photochem. Photobiol. 1992, 55, 373.
(22) Amrine, C. S. M.; Raja, H. A.; Darveaux, B. A.; Pearce, C. J.; Oberlies, N. H. Media studies to enhance the production of verticillins facilitated by in situ chemical analysis. J. Ind. Microbiol. Biotechnol. 2018, 45, 1053.
(23) Al Subeh, Z. Y.; Raja, H. A.; Obike, J. C.; Pearce, C. J.; Croatt, M. P.; Oberlies, N. H. Media and strain studies for the scaled production of cis-enoate resorcylic acid lactones as feedstocks for semisynthesis. J. Antibiot. 2021, 74, 496.
(24) O’Brien, E. M.; Morgan, B. J.; Mulrooney, C. A.; Carroll, P. J.; Kozlowski, M. C. Perylenequinone natural products: total synthesis of hypocrellin A. J. Org. Chem. 2010, 75, 57.
(25) Mulrooney, C. A.; Morgan, B. J.; Li, X.; Kozlowski, M. C. Perylenequinone natural products: enantioselective synthesis of the oxidized pentacyclic core. J. Org. Chem. 2010, 75, 16.
(26) O’Brien, E. M.; Li, J.; Carroll, P. J.; Kozlowski, M. C. Synthesis of the cores of hypocrellin and shiraiachrome: diastereoselective 1,8-diketone aldol cyclization. J. Org. Chem. 2010, 75, 69.
(27) Hu, J.; Sarrami, F.; Li, H.; Zhang, G.; Stubbs, K. A.; Lacey, E.; Stewart, S. G.; Karton, A.; Piggott, Andrew M.; Chooi, Y.-H.; Brien, E. M.; Li, J.; Carroll, P. J.; Kozlowski, M. C. Synthesis of the cores of hypocrellin and shiraiachrome: diastereoselective 1,8-diketone aldol cyclization. J. Org. Chem. 2010, 75, 69.
Heterologous biosynthesis of elsinochrome A sheds light on the formation of the photosensitive perylenequinone system. Chem. Sci. 2019, 10, 1457.

(28) Kishi, T.; Tahara, S.; Taniguchi, N.; Tsuda, M.; Tanaka, C.; Takahashi, S. New perylenequinones from Shiraia bambusicola. Planta Med. 1991, 57, 376.

(29) Wu, H.; Lao, X.-F.; Wang, Q.-W.; Lu, R.-R.; Shen, C.; Zhang, F.; Liu, M.; Jia, L. The shiraiachromes: novel fungal perylenequinone pigments from Shiraia bambusicola. J. Nat. Prod. 1989, 52, 948.

(30) Bogeski, I.; Gulaboski, R.; Kappl, R.; Mirceski, V.; Stefova, M.; Petreska, J.; Hoth, M. Calcium binding and transport by coenzyme Q. J. Am. Chem. Soc. 2011, 133, 9293.

(31) Zhou, Z.; Takaya, N.; Nakamura, A.; Yamaguchi, M.; Takeo, K.; Shoun, H. Ammonia fermentation, a novel anoxic metabolism of nitrate by fungi. J. Biol. Chem. 2002, 277, 1892.

(32) Krueger, M. C.; Bergmann, M.; Schlosser, D. Widespread ability of fungi to drive quinone redox cycling for biodegradation. FEMS Microbiol. Lett. 2016, 363, fnw105.

(33) Pedrini, N.; Ortiz-Urquiza, A.; Huarte-Bonnet, C.; Fan, Y.; Juárez, M. P.; Keyhani, N. O. Tenebrionid secretions and a fungal benzoquinone oxidoreductase form competing components of an arms race between a host and pathogen. Proc. Natl. Acad. Sci. U. S. A. 2015, 112, No. E3651.

(34) Romans, H.; Schlosser, D.; Wick, L. Y. Untapped potential: exploiting fungi in bioremediation of hazardous chemicals. Nat. Rev. Microbiol. 2011, 9, 177.

(35) Koch, R. A.; Yoon, G. M.; Aryan, U. K.; Lail, K.; Amirebrahimi, M.; LaButti, K.; Lipzen, A.; Riley, R.; Barry, K.; Henrissat, B.; Grigoriev, I. V.; Herr, J. R.; Aime, M. C. Symbiotic nitrogen fixation in the reproductive structures of a basidiomycete fungus. Curr. Biol. 2021, 31, 3905.

(36) Wei-Shin, C.; Yuan-Teng, C.; Xiang-Yi, W.; Friedrichs, E.; Puff, H.; Breitmaier, E. Die Struktur des Hypocrellins und seines Photooxidationsproduktes Peroxyhypocrellin. Liebigs Ann. Chem. 1981, 1981, 1880.

(37) Fang, L. Z.; Qing, C.; Shao, H. J.; Yang, Y. D.; Dong, Z. J.; Wang, F.; Zhao, W.; Yang, W. Q.; Liu, J. K. Hypocrellin D, a cytotoxic fungal pigment from fruiting bodies of the ascomycete Shiraia bambusicola. J. Antibiot. 2006, 59, 351.

(38) Tyler, A. R.; Ragbir, R.; McMonagle, C. J.; Waddell, P. G.; Heaps, S. E.; Steed, J. W.; Thaw, P.; Hall, M. J.; Probert, M. R. Encapsulated nanodroplet crystallization of organic-soluble small molecules. Chem. 2020, 6, 1755.

(39) Mayhew, S. G. The redox potential of dithionite and SO-2 from equilibrium reactions with flavodoxins, methyl viologen and hydrogen plus hydrogenase. Eur. J. Biochem. 1978, 85, 535.

(40) Blankenhorn, G.; Moore, E. G. Sulfoxylate ion (HSO2-), the hydride donor in dithionite-dependent reduction of NAD+ analogs. J. Am. Chem. Soc. 1980, 102, 1092.

(41) Bruhn, T.; Schaumlöffel, A.; Hemberger, Y.; Bringmann, G. SpecDis: quantifying the comparison of calculated and experimental electronic circular dichroism spectra. Chirality 2013, 25, 243.

NOTE ADDED AFTER ASAP PUBLICATION
This paper published ASAP on January 25, 2022 with an error in the text. The error was corrected and the revised manuscript reposted on February 15, 2022.