Bioactive α-Pyrone Derivatives from the Endophytic Fungus Diaporthe sp. CB10100 as Inducible Nitric Oxide Synthase Inhibitors

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Inducible nitric oxide synthase (iNOS) produces NO from L-arginine and plays critical roles in inflammation and immune activation. Selective and potent iNOS inhibitors may be potentially used in many indications, such as rheumatoid arthritis, pain, and neurodegeneration. In the current study, five new compounds, including a dibenzo-α-pyrone derivative ellagic acid B (5), and four α-pyrone diaporpyrone A–D (9–12), together with three known compounds (6–8), were isolated from the endophytic fungus Diaporthe sp. CB10100. The structures of these new natural products were unambiguously elucidated using NMR, HRESIMS or electronic circular dichroism calculations. Ellagic acid B (5) features a tetracyclic 6/6/6/6 ring system with a fused 2H-chromene, which is different from ellagic acid (4) with a fused 2H-chromen-2-one. Both 2-hydroxy-alternariol (6) and alternariol (7) reduced the expression of iNOS at protein levels in a dose-dependent manner, using a lipopolysaccharide (LPS)-induced RAW264.7 cell models. Also, they decreased the protein expression levels of pro-inflammatory cytokines, such as tumor necrosis factor-α, interleukin-6 and monocyte chemotactic protein 1. Importantly, 6 and 7 significantly reduced the production of NO as low as 10 μM in LPS-induced RAW264.7 cells. Molecular docking of 6 and 7 to iNOS further suggests that both of them may interact with iNOS. Our study suggests that 6 and 7, as well as the alternariol scaffold may be further developed as potential iNOS inhibitors.

Keywords: endophytic fungus, Diaporthe sp., α-pyrone, anti-inflammation, inducible nitric oxide synthase, NO

INTRODUCTION

Inflammation plays important roles in the occurring and development of many diseases, including rheumatoid arthritis (RA) (Choy and Panayi, 2001), osteoarthritis (OA) (Ahmad et al., 2020), diabetes (Purkayastha and Cai, 2015), and cancers (Zhong et al., 2016). When inflammation occurs, excessive inflammatory mediators are produced by inducible nitric oxide synthase (iNOS) or cyclooxygenase-2 (COX-2), such as NO and prostaglandin E2 (PGE2) (Yu et al., 2019). Non-steroidal anti-inflammatory drugs (NSAIDs) and steroid hormone glucocorticoids are both used for the treatment of inflammation, despite serious side effects (Buchman, 2001; Salvo et al., 2011). iNOS...
is a mammalian protein composed of a C-terminal reductase and an N-terminal oxygenase domain, which produces micromolar NO by oxidizing \( L \)-arginine to \( L \)-citrulline in the presence of bacterial lipopolysaccharide (LPS) and/or proinflammatory cytokines. A considerable number of iNOS inhibitors, such as arginine derivatives, pyrimidines and aminopyrimidines, as well as aliphatic, aromatic, and cyclic amidines, have been developed (Cinelli et al., 2020). Although many of them showed promise in the treatment of arthritis or inflammatory and neuropathic pain in animal models, there are no iNOS inhibitors on the market. Therefore, there is strong need to discover new iNOS inhibitors as anti-inflammation agents.

\( \alpha \)-Pyrone (1, Figure 1A) is an aromatic unsaturated lactone and an important sub-structure of various natural products with interesting biological activities (Mcglacken and Fairlamb, 2005). For example, dothideopyrone F (2) inhibited NO production in LPS-induced BV2 cells and diminished the protein expression levels of iNOS and COX-2 (Kim et al., 2018). In addition, natural products with \( \alpha \)-pyrone sub-structure have antibacterial (Zhang et al., 2013), anti-influenza A virus (Hou et al., 2018), and anti-HIV (Appendino et al., 2007) activities. Some of them are signal molecules (Brachmann et al., 2013), while some are also cytotoxic agents against tumor cells (Rivera-Chávez et al., 2019; Zhu et al., 2019). In particular, dibenzo-\( \alpha \)-pyrone (3) belongs to a significant group of heptaketide coumarin metabolites that have a fused tricyclic core (Figure 1A), commonly identified from fungi, bacteria, and plants (Lai et al., 2016). Many of them also display diverse biological activities including cytotoxicity and anti-inflammatory activity (Mao et al., 2014). For example, ellagic acid (4), found in medicinal plants, vegetables and fruits, exhibits impressive anti-inflammatory and anti-diabetic activities (Rios et al., 2018). It is widely used in skin care products because of its anti-oxidation effect and skin protection effects.

Sinomenine (Supplementary Figure S1), a tetrahydrosoquinoline-type alkaloid isolated from the medicinal plant of Sinomenium acutum (Thunb.) (“青风藤” in Chinese), has strong anti-inflammatory effects and is used in China to treat RA for many years (Zhao et al., 2012; Wang et al., 2019). To our knowledge, there are few reports of endophytes and their natural products from \( S. \) acutum. Since endophytes from medical plants are attractive sources of bioactive natural products, we initiated an endeavor to isolate natural products from endophytes colonized in the stem and root of \( S. \) acutum, in our continuous search for new bioactive metabolites of microorganisms from un- or underexplored niches (Jiang et al., 2018; Jiang et al., 2021a). In this study, we report that these accumulating efforts have resulted in the isolation of multiple endophytes from \( S. \) acutum, and bioactivity-guided natural product isolation have yielded eight compounds, including a new dibenzo-\( \alpha \)-pyrone derivative, ellagic acid B (5) and four new \( \alpha \)-pyrone diaporpyrone A–D (9–12), together with three known compounds (6–8). Further biological evaluation of these compounds revealed that 2-hydroxy-alternariol (6) and alternariol (7) are potential iNOS inhibitors.

**MATERIALS AND METHODS**

**General Methods**

The instruments (including those applied for optical rotations, MS, ORD, NMR, and ECD) and routine reagents for chemical isolation and biological evaluation were the same as those reported previously (Jiang et al., 2021a). The details were listed in the Supporting Information.
Isolation of Endophytes and Fermentation

The separation methods of endophytic fungi are detailed in the Supporting Information. These endophytic fungi CB10098-CB10104 were isolated from the stems and roots of S. acutum, which were collected from Huaihua (Hunan, China) in October 2017. They were grown in PDA (0.3% potato extract, 2% glucose, 1.5% agar power) medium at 28°C for 5 day, and then inoculated to seed medium [glucose 2%, sucrose 1%, soybean powder 0.2%, peptone 1%, K2HPO4 0.03%, poly (ethylene glycol) 0.25%, NaNO3 0.3%, and (NH4)2SO4 0.3%]. The strain was deposited in the Xiangya International Academy of Translational Medicine, Central South University, Changsha, Hunan, China.

Large-Scale Fermentation of Diaporthe sp.

CB10100

For large-scale fermentation of Diaporthe sp. CB10100, 185 × 1 L Erlenmeyer flasks containing rice (100 g) and H2O (100 ml) were used. The fermented solid rice culture was processed to extract the respective fungus by ethyl acetate to obtain extracts C1

Table 1 | 1H NMR (500 MHz) and 13C NMR (125 MHz) data of 5 and 6 in DMSO-d6.

| Position | 5 δH, type | 5 δH, mult. (J in Hz) | 6 δH, type | 6 δH, mult. (J in Hz) |
|----------|------------|----------------------|------------|----------------------|
| 1        | 120.4, C   | 122.0, C             | 2          | 1.22, C              |
| 2        | 101.5, CH  | 6.69, s              | 3          | 141.7, C             |
| 3        | 149.4, C   | 147.7, C             | 4          | 142.1, C             |
| 4        | 102.8, C   | 144.9, C*            | 5          | 138.2, C             |
| 5        | 163.9, C   | 109.2, C             | 6          | 129.8, C             |
| 6a       | 94.1, C    | 138.7, C*            | 7          | 102.4, CH            |
| 7-0H     | 154.2, C   | 100.8, C             | 8          | 156.5, C             |
| 9        | 114.4, C   | 165.0, C             | 9a         | 9a                   |
| 10       |            |                      | 10         | 104.4, CH            |
| 11       | 63.3, CH3  | 7.26, br s           | 12         | 16.0, CH3            |

Table 1 | 1H NMR (500 MHz) and 13C NMR (125 MHz) data of 5 and 6 in DMSO-d6.

*Not detected (Chapla et al., 2014).

Isolation of Compounds 5–12

The EtOAc-soluble fraction (192.8 g) was subjected to silica gel chromatography using petroleum/EtOAc (v/v, 20:1 → 10:1 → 5:1 → 1:5 → 1:20), EtOAc, EtOAc/MeOH (v/v 1:0 → 1:5 → 0:1) to yield nine combined fractions (Fr. A to J). Fr. B (27.13 g) was subjected to silica gel column chromatography using a step gradient elution of petroleum/EtOAc (v/v, 20:1 → 10:1 → 5:1 → 1:1 → 1:10), EtOAc, EtOAc/MeOH (v/v 1:0 → 1:10:1 → 1:1 → 0:1) to yield ten combined fractions (Fr. 2A to 2H). Fr. 7 (58.6 g) was chromatographed on ODS column (MeOH:H2O v/v 3:7 → 4:6 → 6:4 → 7:3) to yield ten combined fractions (Fr. 2J-1 to 2J-10). Fr. 2J-2 (1.32 g) was chromatographed on ODS column (MeOH:H2O v/v 3:7 → 4:6 → 6:4 → 7:3), Sephadex LH-20 (MeOH) column and repeated purification by semi-preparative HPLC [containing 0.2% formic acid, MeCN/H2O, v/v, 5:95 → 25:75 (0–11 min); 25:75 (11–12 min); 25:75 → 5:95 (12–20 min); 5:95 (20–21 min)], which resulted in 9 (4.3 mg) and 10 (5.6 mg). In the same manner, 11 (2.7 mg) and 12 (1.0 mg) were obtained from Fr. 2 J-1.

Ellagic acid B (5)

Dark-grey powder; UV (MeOH) λmax 200.9, 255.2, 287.2, 363.7 (see Supplementary Figure S4A); 1H, 13C and 2D NMR spectroscopic data, see Table 1 and Supplementary Figures S7–S13; HRESIMS m/z 287.0197 [M - H]+ (calcd for C14H7O7, 287.0197).

Diaporyrone A (9)

White powder; UV (MeOH) λmax 202.1, 220.9, 308.7 (see Supplementary Figure S4E); 1H, 13C and 2D NMR spectroscopic data, see Table 2 and Supplementary Figures S35–S48; HRESIMS m/z 195.0655 [M - H]+ (calcd for C10H13O4, 195.0663).

Diaporyrone B (10)

White powder; UV (MeOH) λmax 196.2, 301.5 (see Supplementary Figure S4F) [α]D 23.6 +1.92 (c 0.12, MeOH); ECD (MeOH) λ (ε) 303 (~1.22), 233 (2.02), 208 (~5.97), 1H, 13C and 2D NMR spectroscopic data, see Table 2; Supplementary Figures S49–S56; HRESIMS m/z 239.0922 [M - H]+ (calcd for C12H15O6, 239.0925).

Fungal Strain

The strain CB10100 was identified as Diaporthe sp. according to molecular identification and its ITS sequence has 99% sequence identity to that of Diaporthe sp. The sequence data have been deposited to GenBank with the accession number MW037206. The strain was deposited in the Xiangya International Academy of Translational Medicine, Central South University, Changsha, Hunan, China.
**Diaporpyrone C (11)**

White powder; UV (MeOH) \(\lambda_{max}\) 197.4, 223.3, 307.5 (see Supplementary Figure S4G) [\(\alpha\)]_D^23.6 170+14.72 (c 0.072, MeOH); \(^1\)H, \(^{13}\)C and 2D NMR spectroscopic data, see Table 2; Supplementary Figures S57–S64; HRESIMS \(m/z\) 211.0607 [M - H]\(^-\) (calcd for C\(_{10}\)H\(_{11}\)O\(_5\), 211.0612).

**Diaporpyrone D (12)**

White powder; UV (MeOH) \(\lambda_{max}\) 199.7, 220.9, 303.9 (see Supplementary Figure S4H); \(^1\)H, \(^{13}\)C and 2D NMR spectroscopic data, see Table 2; Supplementary Figures S65–S71; HRESIMS \(m/z\) 167.0340 [M - H]\(^-\) (calcd for C\(_9\)H\(_7\)O\(_4\), 167.0350).

**ECD Calculation Methods**

The ECD spectrum of diaporpyrone B (10) was calculated by using of the Gaussian 09 package (Frisch et al., 2010). Those configurations were optimized at the B3LYP/6-31G (d) level. The ECD spectrum were calculated by the TD-DFT method at the B3LYP/6-311+ +G (2 day, p) level with the CPCM model in methanol solution (Bruhn et al., 2013). The details were listed in the Supporting Information.

**Cell Culture and Cell Viability Assay**

Cell culture and cell viability assay (Liu et al., 2016) was utilized to investigate the maximum concentration of every compound that was not toxic to cells. The method for cell culture and cell viability assay was in the **Supporting Material**.

**Western Blotting for the Detection of iNOS and COX-2**

The iNOS and COX-2 protein levels affected by compounds 5–12 were detected using Western blotting according to the reported procedures, and dexamethasone was used as the positive control (Jiang et al., 2021a). The procedure is detailed in the **Supporting Material**.

**Detection of NO and Inflammatory Cytokines Production**

RAW264.7 cells were seeded in 12-well plates in a density of 2 × 10\(^5\) cells/mL and incubated for 24 h. Three different concentration compounds were added in the each well for 1 h before LPS stimulation. The culture medium was collected and stored at −80°C. The levels of nitric oxide in the culture medium were detected using the Nitric oxide detection kit (Invitrogen) and the levels of TNF-α, IL-6 and MCP-1 were analyzed using the ELISA kits (eBioscience).

**Molecular Docking Analysis**

Molecular docking studies were conducted by the software Molecular Operating Environment (MOE 2010.06; Chemical Computing Group, Canada) as the computational platform. The procedure is detailed in the **Supplementary Material**.

**Statistical Analysis**

The results are expressed as mean \(\pm\) SD of at least three independent experiments. Statistical significance of differences between groups was determined by ANOVA, and a level of *\(p < 0.05\) or **\(p < 0.01\) was considered statistically significant.

**RESULTS AND DISCUSSION**

**Isolation of Endophytic Fungi From Sinomenium acutum**

Six endophytic fungi CB10098-CB10104 were isolated from the stems and roots of S. acutum, collected in Huaihua city in central China (Supplementary Figure S1). The fungus CB10100 was identified as **Diaporthe** species based on DNA sequencing of its internal transcribed spacer 4 (ITS4) and the phylogenetic analysis of its ITS4 with selected **Diaporthe** strains in GenBank (Supplementary Figure S2). **Diaporthe**, the sexual morph of **Phomopsis** and a known plant pathogen (Huang et al., 2013), lives in a diversity of plants (Guarnaccia and Crous, 2017). Some **Diaporthe** or **Phomopsis** species can also produce

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**TABLE 2** \(^1\)H NMR (\(500\) MHz) or (\(400\) MHz) and \(^{13}\)C NMR (\(125\) MHz) or (\(100\) MHz) data of 9–12 in CD\(_2\)OD.

| Position | \(\delta_c\), type | \(\delta_{HC}\), mult. (J in Hz) | \(\delta_c\), type | \(\delta_{HC}\), mult. (J in Hz) | \(\delta_c\), type | \(\delta_{HC}\), mult. (J in Hz) | \(\delta_c\), type | \(\delta_{HC}\), mult. (J in Hz) |
|----------|------------------|---------------------------------|------------------|---------------------------------|------------------|---------------------------------|------------------|---------------------------------|
| 2        | 165.2, C         |                                 | 216.9, C         |                                 | 165.2, C         |                                 | 165.2, C         |                                 |
| 3        | 113.7, CH        | 6.17, d (9.6)                  | 114.1, CH        | 6.23, d (9.6)                  | 113.7, CH        | 6.20, d (9.6)                  | 112.8, CH        | 6.18, d (9.5)                  |
| 4        | 149.6, CH        | 7.44, d (9.6)                  | 145.9, CH        | 7.62, d (9.6)                  | 149.7, CH        | 7.48, d (9.6)                  | 150.2, CH        | 7.45, d (9.5)                  |
| 5        | 116.9, C         |                                 | 120.6, C         |                                 | 116.8, C         |                                 | 113.2, C         |                                 |
| 6        | 160.4, C         |                                 | 160.0, C         |                                 | 160.6, C         |                                 | 161.7, C         |                                 |
| 1'       | 29.5, CH\(_2\)  | 2.41, t (7.2)                  | 26.6, CH\(_2\)  | 1.79, m                        | 37.7, CH\(_2\)  | 1.76, m 1.57, m               | 35.2, CH\(_2\)  | 1.98, m 1.86, m               |
| 2'       | 26.2, CH\(_2\)  | 1.79, m                        | 26.2, CH\(_2\)  | 1.36, m                        | 26.2, CH\(_2\)  | 1.36, m                        | 25.9, CH\(_2\)  | 2.52, m                        |
| 3'       | 34.4, CH\(_2\)  | 2.33, t (7.2)                  | 26.0, CH\(_2\)  | 1.63, m                        | 176.7, C         |                                 | 176.7, C         |                                 |
| 4'       | 176.8, C         |                                 | 35.2, CH\(_2\)  | 2.29, t (4.5)                  | 17.2, CH\(_3\)  | 2.27, s                        | 17.2, CH\(_3\)  | 2.32, s                        |
| 5'       | 172.1, C         |                                 | 17.8, C          |                                 | 17.6, CH\(_3\)  | 2.26, s                        |                   |                                 |

Not detected. 

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bioactive small molecules and protect host plants from fungi infection (Tan and Zou, 2001). For example, several bioactive metabolites were discovered, including anthaquinones (i.e., cytosykin A), terpenes (i.e., diaporthein A), alkaloids (i.e., diaprisoindoles A) (Chepkirui and Stadler, 2017) (Supplementary Figure S3). Therefore, a scale-up solid fermentation of Diaporthe sp. CB10100 (185 × 1 L flasks containing 100 g boiled rice) was next used to isolate natural products with anti-inflammatory activity.

Structure Elucidation
The obtained Diaporthe sp. CB10100 crude extract was further fractioned based on silico gel and Sephadex LH-20 chromatography and semi-preparative HPLC, which yielded compounds 5–12. Alternariol (7) and alternariol-9-monomethyl ether (8) are known compounds, and their structures were established based on the comparison of their 1D and 2D NMR spectra, HRMS, and UV spectra with the literature (Supplementary Figures S4C,D, S6C,D, S21–S34) (Hildebrand et al., 2015). Although 2-hydroxy- alternariol (6) was reported in 2007 and 2014 (Pfeiffer et al., 2007; Chapla et al., 2014), three carbons signals of 6 were not identified. Therefore, a complete set of NMR assignment of 6 is listed in Table 1, based on our full set of NMR spectra (Supplementary Figures S14–S20).

Ellagic acid B (5) was isolated as colorless powder. Its molecular formula is established as C_{14}H_{10}O_{5} based on the (-)-HRESIMS analysis at m/z 287.0197 [M - H]⁻ (calcd for C_{14}H_{10}O_{5}, 287.0197), suggesting 11 degrees of unsaturation. Its ¹³C NMR spectrum (Table 1) shows a total of 14 signals that could be classified as an ester carbonyl (δ_C 163.9), 12 sp²-hybridized carbons (including two methine and six quaternary carbons), and one oxygenated aliphatic carbon (δ_C 63.3), assisted by DEPT-135 and HSQC spectra (Supplementary Figures S10, S11). The presence of two aromatic singlet protons at δ_H 6.31 (s) and 6.69 (s) suggests the presence of a likely biphenyl structure. Based on these analyses and the similar UV spectra with 6–8, compound 5 may contain a new dibenzo-α-pyrone ring system with an additional fused α-pyrone moiety. This was corroborated by its HMBC spectrum, showing correlations from the aromatic proton H-2 (δ_H 6.69, s) to C-1, C-3, C-4, C-4b, and from H-7 (δ_H 6.31, s) to C-6, C-6a, C-6b, C-8, and C-9, as well as from one methylene (δ_H 5.29, s) to C-1, C-2, C-4b, C-9a (Figure 2). To our knowledge, this represents a new tetracyclic 6/6/6/6 ring system derived from dibenzo-α-pyrones.

Diaporpyrone A (9) was isolated as a colorless powder. Its molecular formula was determined as C_{14}H_{7}O_{7} on the basis of HRESIMS analysis, affording an [M - H]⁻ ion at m/z 195.0655 [M - H]⁻ (calculated for [M - H]⁻ ion at m/z 195.0663). The ¹H and ¹³C NMR data of 9 are similar to a previously isolated pyrone with a six-carbon aliphatic chain from an endophytic fungus Pericinia sp. F-31, with the replacement of the terminal methyl group by a carboxylic acid (δ_C 178.1) (Table 2; Supplementary Figure S36; Zhang et al., 2015). This was confirmed by the HMBC correlations from ¹H -CH₃ to C-5 (δ_C 116.9) and C-6 (δ_C 160.4), and from C-3 (δ_C 34.4) to C-4⁻ (δ_C 176.8) and C-2⁻ (δ_C 26.2). We named the new α-pyrene 9 as diaporpyrone A, since it is from the fungus Diaporthe sp. CB10100.

Diaporpyrone B (10) was isolated as colorless powder. Its molecular formula was established as C_{14}H_{8}O_{7} on the basis of the (-)-HRESIMS analysis at m/z 239.0922 [M - H]⁻ (calcd for C_{14}H_{8}O_{7}, 239.0925). The ¹³C NMR and DEPT spectra of 10 shows a total of 12 signals containing two carbonyl carbon (δ_C 171.8 and 164.8), two nonprotonated carbons (δ_C 160.0, 120.6), two olefinic methine carbon (δ_C 145.9, 114.1), an oxymethine carbons (δ_C 68.6), four methylene carbons (δ_C 37.7, 35.2, 26.2, and 26.0), and one methyl carbon (δ_C 17.0) (Table 2). The UV spectra of 10 resembled that of 9, and most of the ¹H and ¹³C NMR data of 10 are similar to those of 9. In the ¹H NMR, ¹³C NMR, and HSQC spectra of 10, additional signals attributed to

![Figure 2](image-url)
the presence of a hydroxy group on C-1′ (δC 68.6). This was confirmed by the HMBC correlations from C-1′ to C-4 (δC 145.9), C-5 (δC 120.6), C-6 (δC 160.0), and the correlations of H-1′/H-2′/H-3′/H-4′/H-5′ in the COSY spectrum of 10 (Figure 2A). Electronic circular dichroism (ECD) calculations were next employed to determine the absolute configuration of 10 by comparing the ECD spectra of (1’S)-10 and (1’R)-10 with the experimental result, which suggests that a (1’R)-10 configuration (Figure 2B; Bruhn et al., 2013). In addition, the specific rotation value [α]D25 +1.92 (c 0.12, MeOH) was in opposite with the published specific rotation value of similar α-pyrones dothideopyrone A ([α]D25 -77 (c 0.22, CHCl3)) (Chomcheon et al., 2009) and dothideopyrone F ([α]D25 -118.72 (c 0.05, MeOH)) (Kim et al., 2018), with an S-configuration at C-1′.

Diapopyrone C (11) was isolated as colorless powder, and its molecular formula was determined as C10H12O5 by the (-)-HRESIMS m/z 211.0607 [M - H]− (calcld for C10H12O5, 211.0612), differing from the molecular formula of 10 by two methylenes. The UV spectrum and 1D and 2D NMR data of 11 are similar to those of 10 (Supplementary Figures S4F,G; Table 2), suggesting that 10 and 11 are similar α-pyrones with a fatty acid side chain. The complete structure of 11 was fully assigned based on 1H COSY, HSQC, and HMBC data (Figure 2A; Table 2). Since diapopyrones B (10) and C (11) likely share the same biosynthetic route, diapopyrone C (11) may have the same absolute configuration at C-1′ position, which is further supported by its specific rotation value [α]D25 +14.72 (c 0.072, MeOH), similar to that of diapopyrone B (10).

Diapopyrone D (12) was isolated as colorless powder, and its molecular formula was determined as C10H14O4 by the (-)-HRESIMS m/z 167.0340 [M - H]− (calcld for C10H14O4, 167.0350), differing from the molecular formula of 9 by two methylenes. The UV maximum absorption and 1D NMR data were highly similar to those of 9 (Table 2; Supplementary Figures S4E,H), suggesting the presence of a similar carbon skeleton. The signals for 35.4 and 175.7 ppm are not present in the 13C NMR spectrum of 12, while they are present in both HSQC and HMBC spectra (Supplementary Figures S69, S70). The complete structure of 12 was fully assigned based on these data. The complete structure of 12 was fully assigned based on 1H–13C COSY, HSQC, and HMBC data (Figure 2A; Table 2).

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The biosynthesis of pyrone 9–12 may involve the condensation of three molecules of malonyl-CoA, followed by aldol-type cyclization and the subsequent lactonization to generate the key six-membered lactone intermediate by a PKS (Figure 3B). Next, the additional fatty acid side chains with various lengths may be biosynthesized by a PKS and finally attached to the lactone to generate 9–12. The detailed biosynthetic mechanism of 9–12 remains to be elucidated.

**Effects of Compounds 5–12 on LPS-Induced iNOS and COX-2 Expression in RAW264.7 Cells**

iNOS and COX-2 are key proteins in the inflammatory signaling pathway. When inflammation occurs, iNOS and COX-2 are usually highly expressed, resulting the formation of inflammatory mediators, including NO and PGE2 (Kyoung et al., 2019). Many phenolic compounds exhibit anti-inflammatory activities through inhibition of the iNOS (Ku et al., 2008; Laavola et al., 2015). No anti-inflammatory activity of compound 6 was reported (Pfeiffer et al., 2007; Chapla et al., 2014), while 7 was discovered to modify macrophage phenotype and to inhibit production of NO and inflammatory responses (Solhaug et al., 2015; Chen et al., 2020). In addition, dothideopyrone F (2) was able to diminish the expression of iNOS and COX-2 in lipopolysaccharide (LPS)-induced BV2 cells (Kim et al., 2018). Therefore, we hypothesize that the newly isolated natural products, e.g., 5–6 and 9–12, may have certain anti-inflammatory activity. In lipopolysaccharide (LPS)-stimulated RAW264.7 cell model, we next evaluated their inhibitory activity of iNOS and COX-2 protein expression, along with 7 and 8, using dexamethasone (DEX) as a positive control (Figures 4A–C).

Among the tested compounds, only 6 and 7 strongly inhibited iNOS protein expression, while they had no effects on COX-2 expression in a concentration of 30 μM (Figure 4A). To compare the anti-inflammatory activity of 6 and 7, the anti-inflammatory activity of 6 and 7 in concentrations ranging from 10 to 40 μM was determined in LPS-stimulated RAW264.7 cells. Both compounds inhibit the expression of iNOS in a dose-dependent manner (Figures 4D,E). For example, 6 was able to inhibit about 90% of iNOS expression when applied at 20 μM, while 7 showed slightly less inhibitory activity against iNOS expression. Therefore, the presence of multiple hydroxyl groups in the dibenzo-α-pyrene scaffold would be important for the observed inhibitory activity against iNOS expression, since the methylation of C-9 hydroxyl group abolishes the activity in 8, while the additional C-2 hydroxyl group increases the activity of 6. The fused α-pyrene moiety into dibenzo-α-pyrene scaffold abolishes the inhibitory activity against iNOS expression activity in 5.

No significant cytotoxicity of both compounds to RAW264.7 cells was observed under the test conditions, even up to 100 μM of 6 and 7 were used (Figures 4F,G). Solhaug et al. reported the cytotoxic activity of 7 in cells and we ascribe this difference as the
shorter treatment period (18 h) in our study, compared to the previously used 48 h (Solhaug et al., 2015). These results suggest that the inhibitory activities of 6 and 7 against iNOS expression in LPS-stimulated RAW264.7 cells did not involve general cytotoxicity.

2-Hydroxy-alternariol (6) and Alternariol (7) Decreased the Protein Levels of TNF-α, IL-6 and MCP-1 in LPS-Stimulated RAW264.7 Cells

Macrophages are central effectors in inflammation, and activated macrophages act through the release of cytokines, such as tumor necrosis factor α (TNF-α) and interleukin-6 (IL-6), as well as chemokines, such as monocyte chemoattractant protein-1 (MCP-1) (Fujisawa and Kobayashi, 2005). Since TNF-α, IL-6, and MCP-1 are major mediators of inflammation, we examined the effects of 6 and 7 on LPS-induced TNF-α, IL-6, and MCP-1 production in RAW264.7 cells, using DEX as a control (Figures 5A–F). After RAW264.7 cells were stimulated with LPS (100 ng/ml) for 18 h in the presence or absence of 6 and 7, the levels of TNF-α, IL-6, and MCP-1 in the culture media were measured by the ELISA. As shown in Figures 5A–C, compound 6 is able to inhibit the production of the tested cytokines and chemokines in a dose-dependent manner in RAW264.7 cells stimulated with LPS (p < 0.05). Significantly, the protein level of IL-6 is reduced to ~25% by 6 (40 μM), which is comparable to the treatment by DEX (0.5 μM). Similar inhibitory effects of 7 could be observed for TNF-α and IL-6, while intriguingly, higher concentrations of 7, e.g., 20 and 40 μM, have no obvious effects on MCP-1 production.

Anti-Inflammatory Effects of 2-Hydroxy-Alternariol (6) and Alternariol (7) on LPS-Induced NO Production in LPS-Stimulated RAW264.7 Cells

NO, an important cellular messenger, may mediate various biological processes, including inflammation, induction of tumor cell death, and killing alien microorganisms (Jiang et al., 2021b). We next investigated the release of NO by the treatment of 6 and 7 in LPS-stimulated RAW264.7 cells (Figures 5G,H). The level of nitrite, a stable oxidized product of NO in the culture medium of RAW264.7 cells were measured using a nitric oxide detection kit. The treatment with 6 and 7 reduced LPS-induced NO production in a dose-dependent manner in the range of 10–40 μM. Compound 6 showed stronger inhibition of NO production than 7 in 10 μM, consistent with the more potent inhibition of iNOS by 6.

Molecular Docking Simulations of 6 and 7 With iNOS

Due to the reduction of NO level and the reduced expression of iNOS by the treatment of 6 and 7 in LPS-treated RAW264.7 cells, compounds 6 and 7 may interact with iNOS and block the biosynthesis of NO. To our knowledge, the dibenzo-α-pyrene
The scaffold has not been used as iNOS inhibitors. Therefore, the interactions of 6 and 7 with iNOS were investigated using molecular docking by Molecular Operating Environment (MOE) (Hu et al., 2017; Jiang et al., 2021b). Using the crystal structure of mouse inducible nitric oxide synthase (miNOS, 3NW2) as a template, compound 6 form four hydrogen-bonding interactions with the amino acid residue Trp457, Asp376, and Arg382 of iNOS, while 7 only has one hydrogen-bonding interaction and two π-π interactions with Met428 (Figures 5I,J). Taken together, 6 and 7 may interact with iNOS and could be further developed as specific inhibitors against iNOS.

CONCLUSION

In conclusion, five new compounds, including a new dibenzo-α-pyrene derivative, ellagic acid B (5) and four new α-pyrene,
diaporpyrone A–D (9–12), together with three known compounds (6–8), were isolated from the endophytic fungus *Diaporthe* sp. CB10100. Their structures were determined by the analyses of their NMR and HRESIMS spectra, while the absolute configuration of 10 and 11 was based on ECD spectra and their specific optical rotations. Ellagic acid B (5) features a tetracyclic 6/6/6/6 ring system with a fused 2H-chromene, while diaporpyrones consist of a pyrone and a short fatty acid side chain. The biosynthesis of 5 and 8 may involve a fungus PKS similar to SnPKS19 from *P. nodorum*, while the biosynthesis of 9–12 would likely be catalyzed by distinct fungi PKSs. All of the isolated compounds 5–12 were evaluated for their anti-inflammatory activities in LPS-stimulated RAW 264.7 macrophages. Among them, 6 and 7 display strong

![FIGURE 5](image-url)
inhibitory activities against iNOS and inhibit nitric oxide production. In addition, both compounds significantly reduced the production of pro-inflammatory mediators and cytokines, including NO, TNF-α, IL-6, and MCP-1. Molecular docking of 6 and 7 to iNOS further suggests that they are potential iNOS inhibitors. Considering the importance of iNOS in inflammation and signal transduction, the discovery of 6 and 7 the alternariol scaffold may be further developed as potential iNOS inhibitors.

DATA AVAILABILITY STATEMENT

The datasets presented in this study can be found in online repositories. The names of the repository/repositories and accession number(s) can be found in the article/Supplementary Material.

AUTHOR CONTRIBUTIONS

YH, YD, and HP conceived and designed the experiments. HP, WZ, BH, and CN was responsible for the isolation of compounds. YW performed the anti-inflammation of the compounds. YH, YD, and HP interpreted the data and wrote the paper. All authors read and approved the final manuscript.

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SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fchem.2021.679592/full#supplementary-material

REFERENCES

Ahmad, N., Ansari, M. Y., and Haqiqi, T. M. (2020). Role of iNOS in Osteoarthritis: Pathological and Therapeutic Aspects. *J. Cell. Physiol.* 235, 6366–6376. doi:10.1002/jcp.29607

Appendino, G., Ottino, M., Marquez, N., Bianchi, F., Giana, A., Ballero, M., et al. (2007). Arzanol, an Anti-inflammatory and Anti-HIV-1 Phloroglucinol α-Pyrene from *Helichrysum italicum* ssp. microphyllum. *J. Nat. Prod.* 70, 608–612. doi:10.1021/np060581r

Brachmann, A. O., Brumeyer, S., Kresovic, D., Hitkova, I., Kopp, Y., Manske, C., et al. (2013). SpecDis: Circular Dichroism Spectra. *Chirality* 25, 243–249. doi:10.1002/chir.22138

Buchman, A. L. (2001). Side Effects of Corticosteroid Therapy. *J. Clin. Gastroenterol.* 33, 289–294. doi:10.1097/00004836-200110000-00006

Chapala, V., Zeraik, M., Ximenes, V., Zanardi, L., Lopes, M., Cavalheiro, A., et al. (2014). Bioactive Secondary Metabolites from *Phomopsis* sp., an Endophytic Fungus from *Senna Spectabilis*. *Molecules* 19, 6597–6608. doi:10.3390/molecules19055697

Chen, S., Deng, Y., Yan, C., Wu, Z., Guo, H., Liu, L., et al. (2020). Secondary Metabolites with Nitric Oxide Inhibition from Marine-Derived Fungus *Alternaria* sp. 5102. *Mar. Drugs* 18, 426. doi:10.3390/md18080426

Chepkirui, C., and Stadler, M. (2017). The Genus *Phomopsis*: A Rich Source of Diverse and Bioactive Metabolites. *Mycol. Prog.* 16, 477–494. doi:10.1007/s11557-012-1887-y

Chomcheon, P., Wiwakrutta, S., Sribolmas, N., Ngamrojanavanich, N., Mahidol, C., Ruchirawat, S., et al. (2009). Metabolites from the Endophytic Mitosporic *Dothideomycete* sp. LRUB20. *Phytochemistry* 70, 121–127. doi:10.1016/j.phytochem.2008.10.007

Chooi, Y.-H., Muria-Gonzalez, M. J., Mead, O. L., and Solomon, P. S. (2015). SnFKS19 Encodes the Polyketide Synthase for Alternariol Mycotoxin Biosynthesis in the Wheat Pathogen *Parastagonospora nodorum*. *Appl. Environ. Microbiol.* 81, 5309–5317. doi:10.1128/AEM.00278-15

Choy, E. H. S., and Panayi, G. S. (2001). Cytokine Pathways and Joint Inflammation in Rheumatoid Arthritis. *N. Engl. J. Med.* 344, 907–916. doi:10.1056/NEJM2001032234441207

Cinelli, M. A., Do, H. T., Miley, G. P., and Silverman, R. B. (2020). Inducible Nitric Oxide Synthase: Regulation, Structure, and Inhibition. *Med. Res. Rev.* 40, 158–189. doi:10.1002/med.21599

Frisch, M. J., Trucks, G. W., Schlegel, H. B., Robb, M. A., Cheeseman, J. R., Scalmani, G., et al. (2010). Gaussian 09, Revision C.01. Gaussian, Inc.

Gyorkos, T. W., and Sermeus, W. (2004). Cytokines and Immunopathology of Infectious Disease. *Clin. Rev. Allergy Immunol.* 26, 129–143. doi:10.1023/B:CLRE.0000015822.33526.fc

Hildebrand, A. A., Kohn, B. N., Pfeiffer, E., Metzler, M., and Bunzel, M. (2015). Conjugation of the Mycotoxins Alternariol and Alternariol Monomethyl Ether in Tobacco Suspension Cells. *J. Agric. Food Chem.* 63, 4728–4736. doi:10.1021/acs.jafc.5b00806

Hou, L., Huang, H., Li, H., Wang, S., Ju, J., and Li, W. (2018). Overexpression of a Type III PKS Gene Affording Novel Violapyrones with Enhanced Anti-inflammation A Virus Activity. *Microb. Cell Fact.* 17, 1–11. doi:10.1186/s1294-018-0958-9

Hu, Z., Wu, Y., Xie, S., Sun, W., Guo, Y., Li, X.-N., et al. (2017). Phomosterone A and B, Two Functionalized Ergostane-Type Steroids from the Endophytic Fungus *Phomopsis* sp. TJ507A. *Org. Lett.* 19, 258–261. doi:10.1021/acs.orglett.6b03557

Huang, F., Hou, X., Dewdney, M. M., Fu, Y., Chen, G., Hyde, K. D., et al. (2013). *Diaporthe* Species Occurring on Citrus in China. *Fungal Divers.* 61, 237–250. doi:10.1007/s11235-013-0245-6

Jiang, L., Pu, H., Qin, X., Liu, J., Wen, Z., Huang, Y., et al. (2021a). Syn-2, 3-diols and anti-inflammatory Indole Derivatives from *Streptomyces* sp. CB09001. *Nat. Prod. Res.* 35, 144–151. doi:10.1080/14787449.2019.1611812
