A New Dinor-cis-Labdane Diterpene and Flavonoids with Antimycobacterium Activity from Colebrookaea oppositifolia

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Abstract

The new 14,15-dinor-cis-labdane diterpene, named (+)-14,15-dinor-9a-hydroxy-cis-labd-11(E)-en-13-one (1), was isolated from the acetone extract of the aerial parts of Colebrookaea oppositifolia, along with the known compounds alunisint (2), moslovafone (3), flindulatin (4), 5,6,7-trimethoxy baicaulein (5), tanetin (6), scutellarein 4’-methyl ether (7), apigenin (8), caffeic acid (9), anisofolin A (10), apigetrin (11), and forsythoside A (12). Structures of the new and known compounds were established by detailed analysis of 1D and 2D nuclear magnetic resonance studies. The isolated compounds 1-12 were evaluated for their antimycobacterium activity against Mycobacterium tuberculosis H37Ra and Mycobacterium bovis in both dormant and active phases. Compounds 1, 7, and 8 exhibited inhibitory activity against M. tuberculosis with IC50 values in the range of 8.1-55.0 µM (MIC 11.5-17.1 µM) in the active phase and 7.4-43.5 µM (MIC 11.5-123.3 µM) in the dormant phase. Similarly 1, 7, and 8 exhibited inhibitory activity against M. bovis with IC50 values in the range of 4.1-98.5 µM (MIC 13.7-161.0 µM) in the active phase and 4.1-111.1 µM (MIC 13.0-166.4 µM) in the dormant phase.

Key words
Colebrookaea oppositifolia · Lamiaceae · 14,15-dinor-cis-labdane diterpene · antimycobacterium · flavonoids

Supporting information available online at http://www.thieme-connect.de/products

Colebrookaea (Lamiaceae) is a monotypic genus represented by Colebrookaea oppositifolia Sm. (syn. = Colebrookaea ternifolia Roxb.), commonly known as Panrasa, and is distributed in hilly parts of India and China [1,2]. The roots of this shrub are used for epilepsy and the leaves are applied for wound healing and bruises [1-3]. It is used for the treatment of fractures, traumatic injuries, and rheumatoid arthritis in China [2]. Some other traditional uses are: the decoction of its roots is given as an abortifacient; the juice of the leaves is used to stop bleeding and as an eye and ear drop; and the paste of the leaves is applied to toothaches and mouth and tongue sores [4]. Different extracts of this shrub are reported to exhibit antibacterial [5-7], antitymocobacterial [8], antioxidant [7], and antifertility [9] activities. Acteoside, a constituent from the aerial parts, exhibited an in vitro potent synergistic fungicidal effect in combination with amphotericin B [10].

Different parts of this plant have been studied phytochemically to isolate flavonoids [2,11-14], acteoside [10], sterols [15], and fatty compounds [11,15]. Despite the availability of treatment, tuberculosis (TB) continues to be a deadly disease [16-18]. We are continuously involved in the process of the isolation of novel phytochemicals with promising anti-TB activity [19-21]. During our program for the isolation of anti-TB compounds from plants found in Western Ghats of Maharashtra, India, a phytochemical analysis of the acetone extract of the aerial parts of C. oppositifolia was performed. Herein we report the isolation and structure elucidation of compounds 1-12 (Fig. 1) and their evaluation for antimycobacterium activity against two microbial strains, Mycobacterium tuberculosis H37Ra and Mycobacterium bovis in both active and dormant phases.

Compound 1 was obtained as yellow gum. Analysis of the 13C NMR and DEPT-135 spectra revealed 18 resonances along with a pseudomolecular peak [M + Na]+ at m/z 301.2135 in the high-resolution electrospray ionization mass spectrometry (HRESIMS; Fig. 1). Supporting Information) and allowed for the determination of the molecular formula as C18H30O2, corresponding to four indices of the hydrogen deficiency. The 1H NMR data (Table 1) showed the presence of four tertiary methyl singlets at δH 0.86, 0.90, 1.06, and 2.27, and one secondary methyl at δH 0.72 with coupling constant 6.8 Hz. Two methyne protons at δH 6.35 and 6.80 with coupling constant 15.9 Hz suggested the presence of a trans double bond. 13C NMR data (Table 1) showed the presence of a carbonyl carbon at δC 197.8 and two methine carbons at δC 130.1 and 151.4 accounted for two indices of hydrogen deficiencies, suggesting that 1 was a bicyclic diterpenoid. The NMR data of 1 was similar to previously reported dinor-labdane diterpene, 8-hydroxy-14,15-dinor-11-labden-13-one [22,23], except that a tertiary carbon at δC 73.5 at position C-8 was replaced by a methine carbon at δC 34.0, and a methine carbon at δC 67.0 at position C-9 was replaced by the oxygenated tertiary carbon at δC 79.6 in compound 1. These observations established the 14,15-dinor diterpene skeleton for 1. The structure was confirmed by 2D NMR studies as follows: The methyl protons at δH 0.86 and 0.90 showed the heteronuclear multiple bond correlation (HMBC; Fig. 6A, Supporting Information) with a methylene carbon at δC 41.6 and a quaternary carbon at δC 33.4. A methine proton at δH 1.31 and 1.50 showed the HMBC correlation with a methylene carbon at δC 30.6 and with a methyl carbon at δC 17.2. Methyl protons at δH 0.72 and 1.06 and a methine proton at δH 6.35 showed the HMBC correlation with an oxygenated tertiary carbon at δC 79.6. This suggested the placement of an oxygenated tertiary carbon at position C-9 and a methyl (δH 0.72) at position C-8 unequivocally. Other HMBC correlations from protons at δH 0.86 and 6.80 with the ketonic carbonyl carbon at δC 197.8 and a proton at δH 6.35 with a methyl carbon at δC 27.9 were also observed (Fig. 2). The correlation spectroscopy (COSY; Fig. 7A, Supporting Information) correlations were observed between δH 1.38 and 1.43, δH 1.43 and 1.17, δH 1.50 and 1.31, δH 1.31 and 1.52, δH 1.52 and 2.04, and δH 2.04 and 0.72, and suggested a H2-1-H2-2-H3-3 and H1-5-H6-6-H7-7-H8-8-H9-9 linkage, which confirmed the presence of a decalin ring skeleton in 1 (Fig. 2).

Comparison of NMR values of 1 with those reported for 8-hydroxy-14,15-dinor-11-labden-13-one [22,23] revealed a deviation in the chemical shift at the carbon 5, i.e., upfield shift by 10 ppm at C-5 in 1. This upfield shift of C-5 ( ~ 10 ppm) can be explained by the cis-fused A/B ring junction [24,25]. The fused rings assume a nonsteroidal conformation (cis A/B ring junction),
as revealed by the strong nuclear overhauser effect spectroscopy (NOESY; Fig. 8, Supporting Information) correlation observed between H3-20 (δH 1.06) and H-5 (δH 1.50) (© Fig. 2). Other observed NOESY correlations were between δH 1.06 and 6.80 and between δH 1.06 and 2.04, and indicated that the presence of the side chain at position C-9 was β orientated and a methyl at position C-8 was α orientated. Thus, based on a combination of detailed analysis of the 2D NMR data and comparison of observed and literature NMR data with the reported compound [22, 23], 1 was identified as a new natural product, (+)-14,15-dinor-9α-hydroxy-cis-labd-11(E)-en-13-one, and belongs to the rare class of 14,15-dinor-diterpenes with a cis A/B ring junction. Compounds 2–12 (© Fig. 1) were identified by comparison with observed and literature NMR data, and supported by liquid chromatography electrospray ionization mass spectrometry (LCESIMS) data. Compounds 2–8 were identified as flavonoids alnustin (2) [26], mosloflavone (3) [27], flindulatin (4) [28], 5,6,7-trimethoxy baicalein (5) [29], tanetin (6) [30], scutellarein 4′-methyl ether (7) [31], and apigenin (8) [32]. Compound 9 was

**Fig. 1** Compounds 1–12 isolated from the acetone extract of the aerial parts of *C. oppositifolia*.
identified as a caffeic acid [33]. Compounds 10 and 11 were identified as flavonoid glycosides anisofolin A [34] and apigetrin [35], respectively. Compound 12 was identified as forsythoside A [36]. Compounds 2-4 and 6-12 are reported for the first time from the genus Colebrookea, except compound 5 was already reported from C. oppositifolia [12]. Anisofolin A (10), which was previously reported from Leucas mollissima Wall. ex Benth. (Lamiaceae) by Chinchansure and coworkers, with its antimycobacterium activity, is also reported in this study [21].

Material and Methods

General experimental procedures, chemical, and biochemicals: Melting points were measured on Buchi B-540 instrument. Optical rotations were measured with a JASCO P-1020 polarimeter. UV spectra were measured with a SpectraMax plus 384 Microplate Reader. The IR spectrum was measured with a Bruker ALPHA FT-IR Spectrometer. 1H and 13C NMR spectroscopic data were recorded on a Bruker Avance Ultra Shield NMR instrument (1H: 500 MHz, 13C: 125 MHz), see Table 1. HRESIMS data was recorded with a ThermaPULSAR spectrometer. 1H and 13C NMR data for compound 1 in CDCl3 (500 MHz for 1H and 125 MHz for 13C, 6 in ppm).

Table 1 1H and 13C NMR data for compound 1 in CDCl3 (500 MHz for 1H and 125 MHz for 13C, 6 in ppm).

<table>
<thead>
<tr>
<th>Position</th>
<th>δH</th>
<th>δC</th>
<th>HMBC</th>
</tr>
</thead>
<tbody>
<tr>
<td>1a</td>
<td>1.38 (m)</td>
<td>21.4</td>
<td>–</td>
</tr>
<tr>
<td>1b</td>
<td>1.59 (m)</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>2a</td>
<td>1.43 (m)</td>
<td>18.4</td>
<td>–</td>
</tr>
<tr>
<td>2b</td>
<td>1.52 (m)</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>3a</td>
<td>1.17 (m)</td>
<td>41.6</td>
<td>C-18, C-19</td>
</tr>
<tr>
<td>3b</td>
<td>1.36 (m)</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>4</td>
<td>–</td>
<td>33.4</td>
<td>–</td>
</tr>
<tr>
<td>5</td>
<td>1.50 (m)</td>
<td>45.4</td>
<td>C-6, C-20</td>
</tr>
<tr>
<td>6a</td>
<td>1.31 (m)</td>
<td>30.6</td>
<td>C-8</td>
</tr>
<tr>
<td>6b</td>
<td>1.65 (m)</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>7a</td>
<td>1.12 (m)</td>
<td>32.8</td>
<td>C-20</td>
</tr>
<tr>
<td>7b</td>
<td>1.52 (m)</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>8</td>
<td>2.04 (m)</td>
<td>34.0</td>
<td>C-6</td>
</tr>
<tr>
<td>9</td>
<td>–</td>
<td>79.6</td>
<td>–</td>
</tr>
<tr>
<td>10</td>
<td>–</td>
<td>42.4</td>
<td>–</td>
</tr>
<tr>
<td>11</td>
<td>6.80 (d, J = 15.9 Hz)</td>
<td>151.4</td>
<td>C-13</td>
</tr>
<tr>
<td>12</td>
<td>6.35 (d, J = 15.9 Hz)</td>
<td>130.1</td>
<td>C-9, C-13, C-16</td>
</tr>
<tr>
<td>13</td>
<td>–</td>
<td>197.8</td>
<td>–</td>
</tr>
<tr>
<td>14</td>
<td>2.27</td>
<td>27.9</td>
<td>–</td>
</tr>
<tr>
<td>15</td>
<td>0.72 (d, J = 6.8 Hz)</td>
<td>16.2</td>
<td>C-9</td>
</tr>
<tr>
<td>16</td>
<td>0.90 (s)</td>
<td>33.7</td>
<td>C-3, C-4</td>
</tr>
<tr>
<td>17</td>
<td>0.86 (s)</td>
<td>22.0</td>
<td>C-3, C-4</td>
</tr>
<tr>
<td>18</td>
<td>1.06 (s)</td>
<td>17.2</td>
<td>C-5, C-7, C-9</td>
</tr>
</tbody>
</table>

Antimycobacterial assay: M. tuberculosis H37Rv (ATCC No. 23000) and M. bovis (ATCC 35734) strains were tested for their susceptibility to compounds 1-12 in active and dormant phases by using the XRMA protocol. All the experiments were performed in triplicate, and IC50 and MIC values were calculated from their dose-response curves. The MIC was defined as the lowest concentration of the anti-tubercular agents that prevented visible growth with respect to the growth control [37-39]. Rifampicin was used as a positive control.

Compounds 1, 7, and 8 exhibited inhibitory activity against M. tuberculosis with IC50 values in the range of 8.1-55.0 µM (MIC 14.4-119.7 µM) in the active phase and 7.4-43.5 µM (MIC 11.5-123.3 µM) in the dormant phase (Table 2). Similarly, 1, 7, and 8 exhibited inhibitory activity against M. bovis with IC50 values in the range of 4.1-98.5 µM (MIC 13.7-161.0 µM) in the active phase and 4.1-111.1 µM (MIC 13.0-166.4 µM) in the dormant phase (Table 2).

Supporting information

HRESIMS and 1D and 2D NMR data of compound 1, dose dependence curves for compound 1, the isolation of compounds 1–12, and NMR and other characterization data for compounds 2–12 are available as Supporting Information.

Acknowledgements

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Conflict of Interest

The authors declare no conflict of interest.

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Table 2 Antimycobacterial activity of compounds 1, 7, and 8 against M. tuberculosis H37Ra and M. bovis.

<table>
<thead>
<tr>
<th>Compound</th>
<th>Mycobacterium tuberculosis H37Ra</th>
<th>Mycobacterium bovis</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Active I C50 (MICT)</td>
<td>Dormant I C50 (MICT)</td>
</tr>
<tr>
<td>1</td>
<td>55.0 ± 0.3</td>
<td>(119.7 ± 0.4)</td>
</tr>
<tr>
<td>7</td>
<td>23.7 ± 0.2</td>
<td>(80.3 ± 0.3)</td>
</tr>
<tr>
<td>8</td>
<td>8.1 ± 0.1</td>
<td>(14.4 ± 0.2)</td>
</tr>
<tr>
<td>Rifampicin</td>
<td>0.0021 ± 0.0004</td>
<td>(0.019 ± 0.003)</td>
</tr>
</tbody>
</table>

* µM = micro molar

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genin 7-O-glucoside from chamomile (Chamomilla recutita [L.] Rau-

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