

# Anti-Lymphangiogenic Terpenoids from the Heartwood of Taiwan Juniper, *Juniperus chinensis* var. *tsukusiensis*

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Table S1. Isolation and purification methods of known compounds isolated from *J. chinensis* var. *tsukusiensis*

Fraction	HPLC method	Obtained compounds (mg)
A	EtOAc:CH <sub>2</sub> Cl <sub>2</sub> : <i>n</i> -Hexane = 5:5:90	<b>34</b> (75.4 mg), <b>35</b> (26.4 mg), <b>36</b> (10.0 mg)
B	EtOAc:CH <sub>2</sub> Cl <sub>2</sub> : <i>n</i> -Hexane = 10:10:80	<b>8</b> (2.0 mg), <b>10</b> (8.6 mg)
C	EtOAc:CH <sub>2</sub> Cl <sub>2</sub> : <i>n</i> -Hexane = 5:5:90	<b>16</b> (5.6 mg)
D	EtOAc:CH <sub>2</sub> Cl <sub>2</sub> : <i>n</i> -Hexane = 10:10:80	<b>19</b> (2.3 mg), <b>30</b> (10.0 mg), <b>36</b> (10.0 mg)
E	EtOAc:CH <sub>2</sub> Cl <sub>2</sub> : <i>n</i> -Hexane = 10:10:80	<b>29</b> (3.5 mg)
F	EtOAc:CH <sub>2</sub> Cl <sub>2</sub> : <i>n</i> -Hexane = 10:5:85	<b>29</b> (3.0 mg)
G	EtOAc:CH <sub>2</sub> Cl <sub>2</sub> : <i>n</i> -Hexane = 10:5:85	<b>22</b> (450.2 mg)
H	EtOAc: <i>n</i> -Hexane = 20:80	<b>11</b> (142.0 mg), <b>14</b> (16.0 mg), <b>20</b> (24.3 mg), <b>21</b> (6.7 mg), <b>24</b> (5.6 mg), <b>28</b> (35.6 mg), <b>32</b> (16.7 mg), <b>37</b> (15.0 mg),
I	EtOAc:CH <sub>2</sub> Cl <sub>2</sub> : <i>n</i> -Hexane = 20:5:75	<b>6</b> (3.0 mg), <b>13</b> (12.0 mg), <b>38</b> (10.0 mg)
J	EtOAc:CH <sub>2</sub> Cl <sub>2</sub> : <i>n</i> -Hexane = 20:5:75	<b>26</b> (2.3 mg), <b>33</b> (363.0 mg)
K	EtOAc: <i>n</i> -Hexane = 30:70	<b>7</b> (12.0 mg)
L	EtOAc:CH <sub>2</sub> Cl <sub>2</sub> : <i>n</i> -Hexane = 20:20:60	<b>9</b> (3.6 mg), <b>15</b> (1.2 mg), <b>18</b> (3.8 mg), <b>23</b> (25.6 mg), <b>31</b> (13.7 mg),
M	EtOAc:CH <sub>2</sub> Cl <sub>2</sub> : <i>n</i> -Hexane = 40:20:40	<b>12</b> (6.5 mg), <b>17</b> (35.5 mg)
N	EtOAc:CH <sub>2</sub> Cl <sub>2</sub> : <i>n</i> -Hexane = 60:20:20	<b>25</b> (13.5 mg), <b>27</b> (3.2 mg)
O	EtOAc:CH <sub>2</sub> Cl <sub>2</sub> : <i>n</i> -Hexane = 60:20:20	

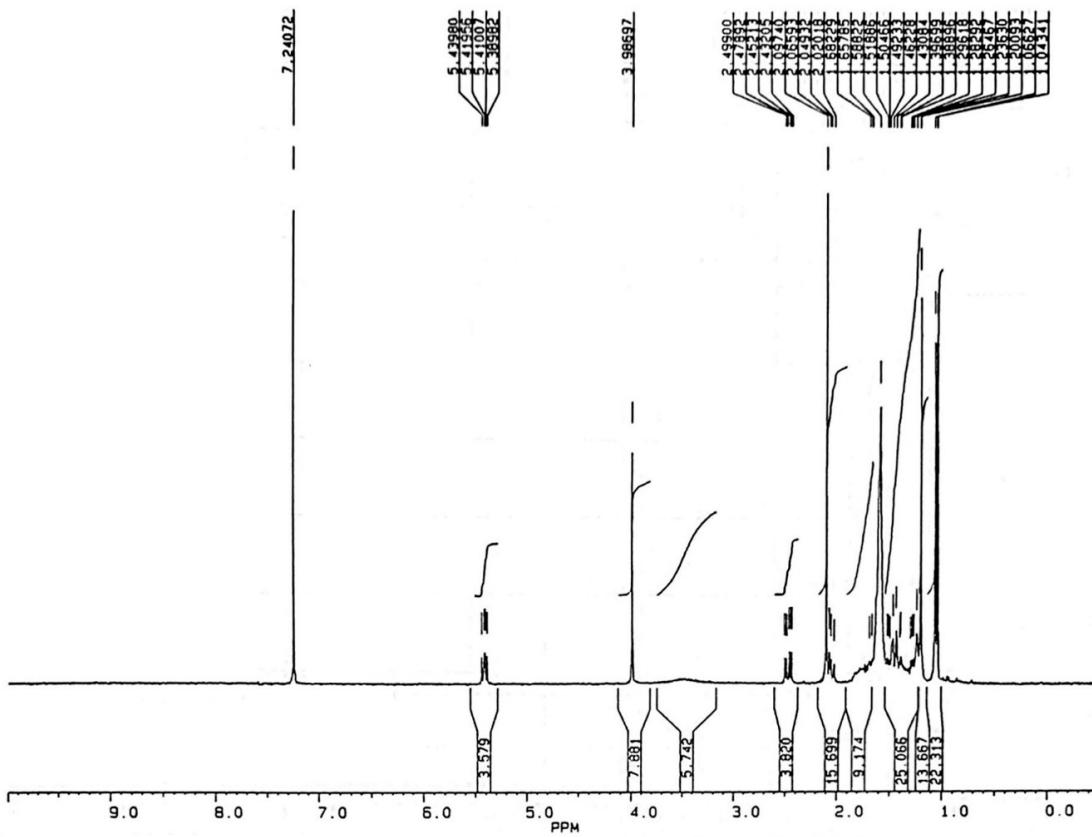


Figure S1.  $^1\text{H}$  NMR spectrum of 12-acetoxywiddrol (**1**) in  $\text{CDCl}_3$  at 300 MHz

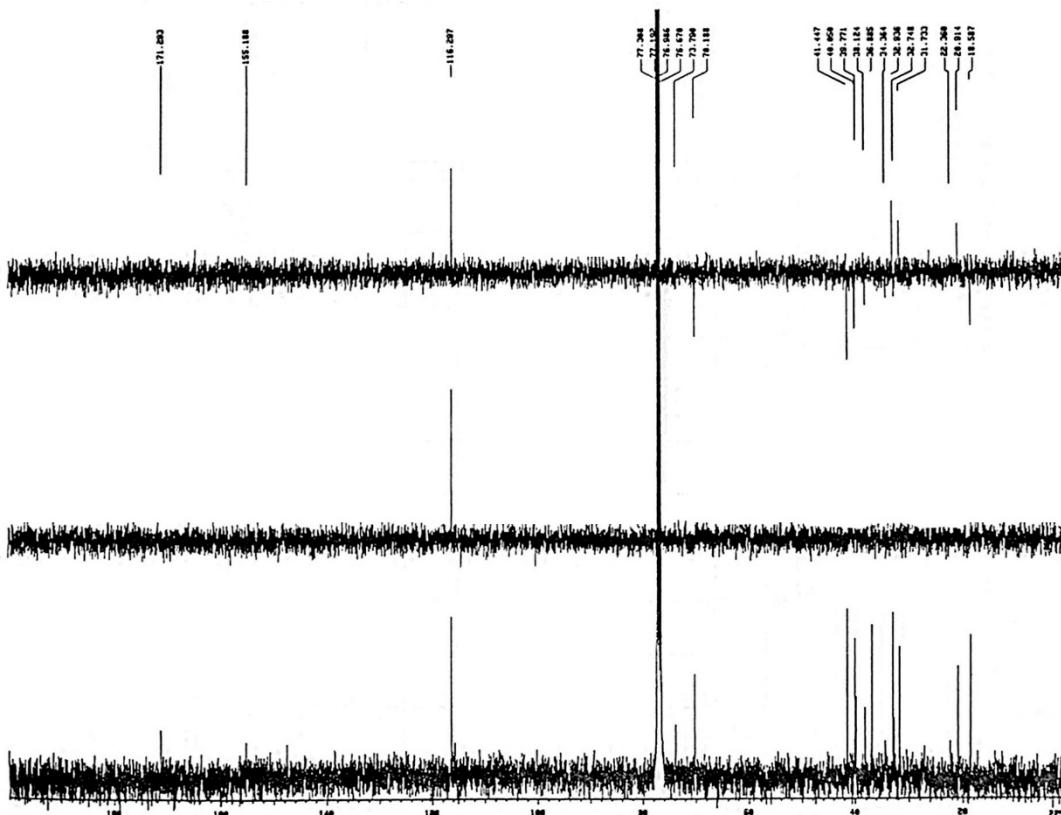


Figure S2. DEPT spectrum of 12-acetoxywiddrol (**1**)

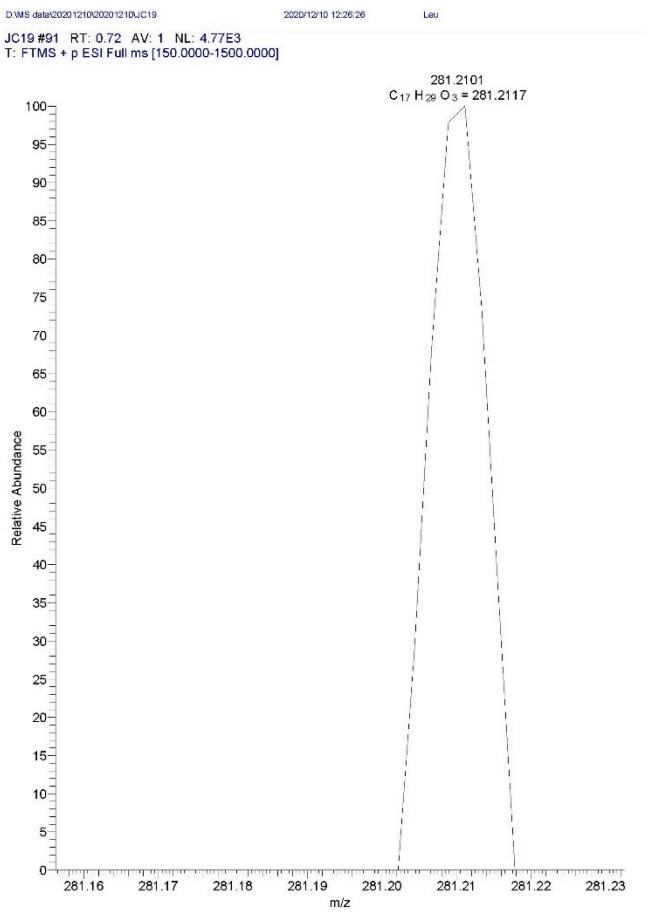


Figure S3. HRESIMS spectrum of 12-acetoxywiddrol (**1**)

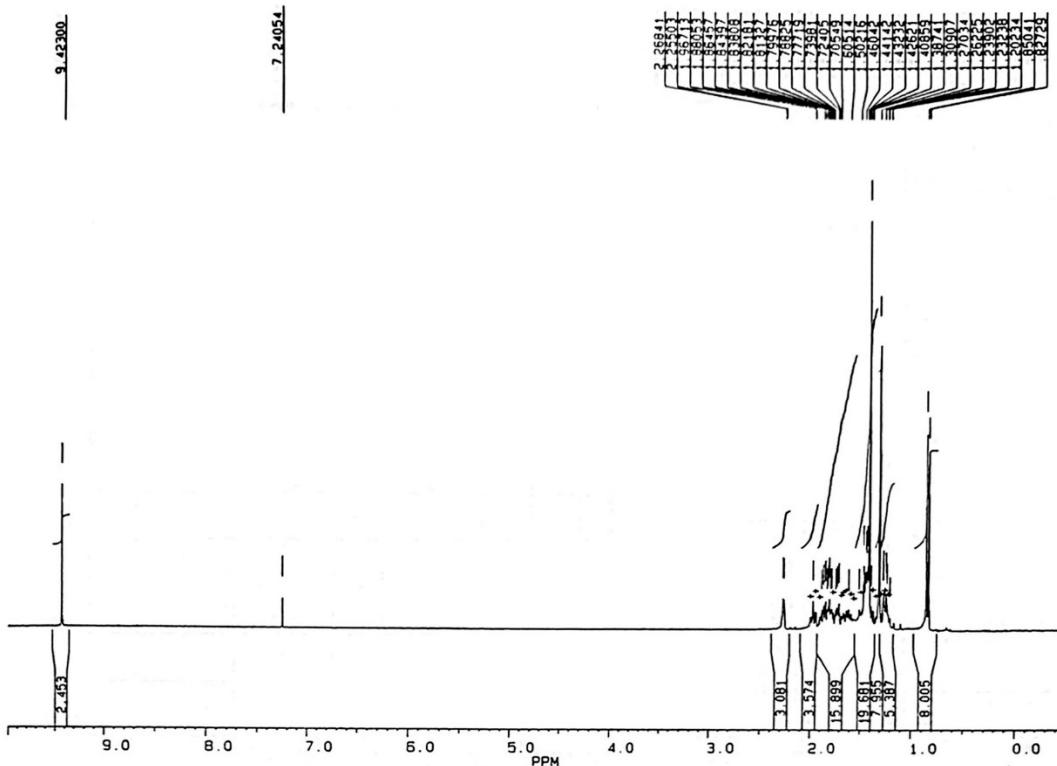


Figure S4.  $^1\text{H}$  NMR spectrum of cedrol-13-al (**2**) in  $\text{CDCl}_3$  at 300 MHz

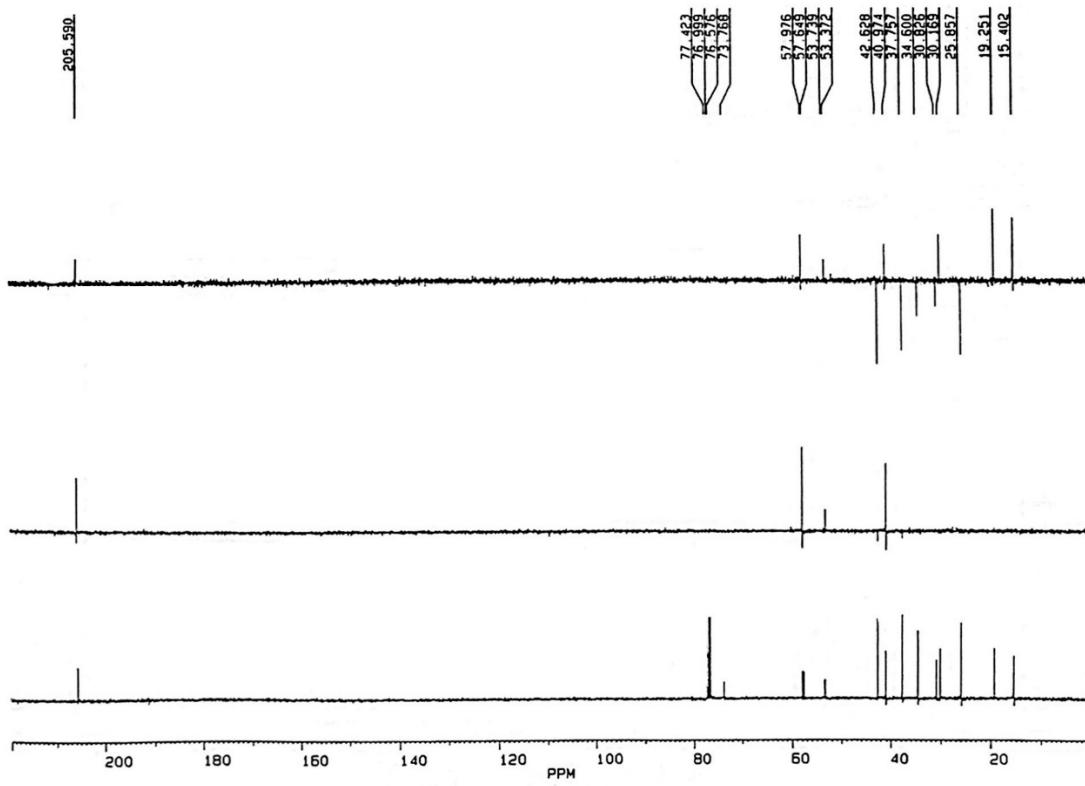


Figure S5. DEPT spectrum of cedrol-13-al (**2**)

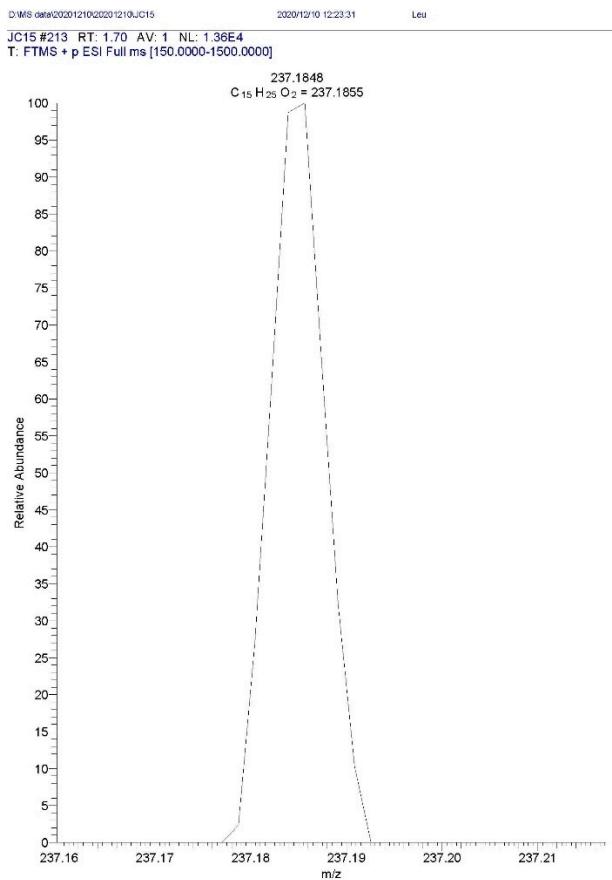


Figure S6. HRESIMS spectrum of cedrol-13-al (**2**)

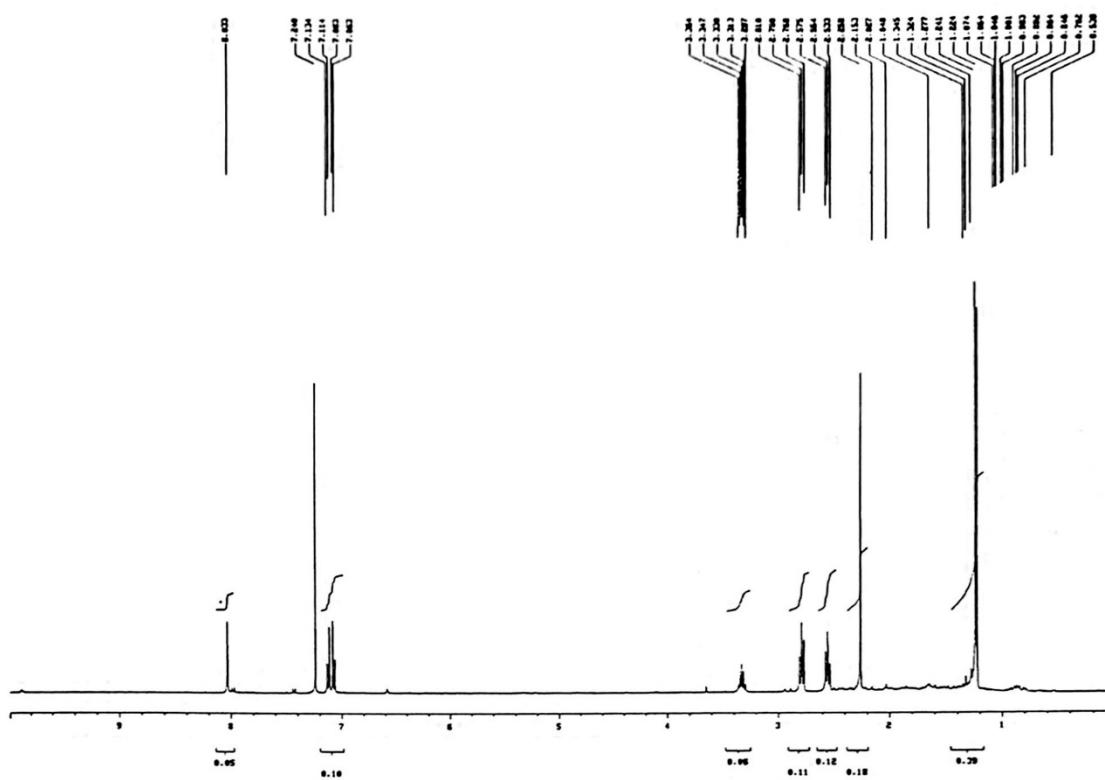


Figure S7.  $^1\text{H}$  NMR spectrum of  $\alpha$ -corocalen-15-oic acid (3) in  $\text{CDCl}_3$  at 400 MHz

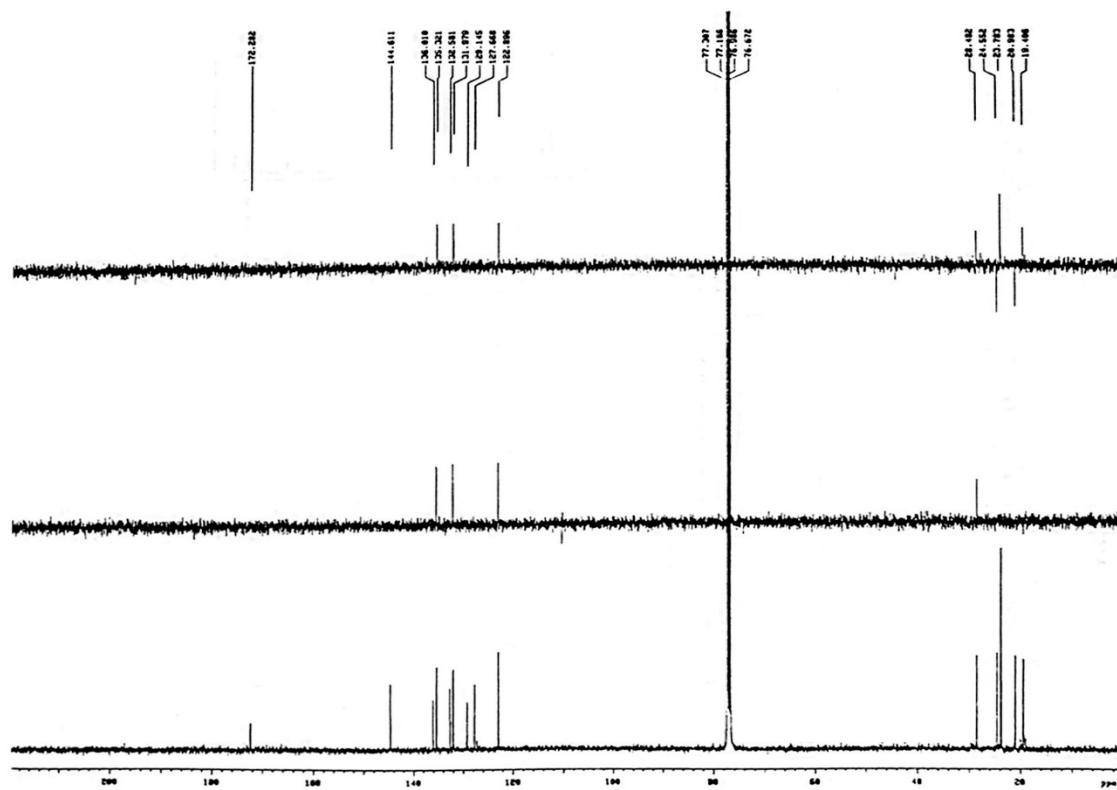


Figure S8. DEPT spectrum of  $\alpha$ -corocalen-15-oic acid (3)

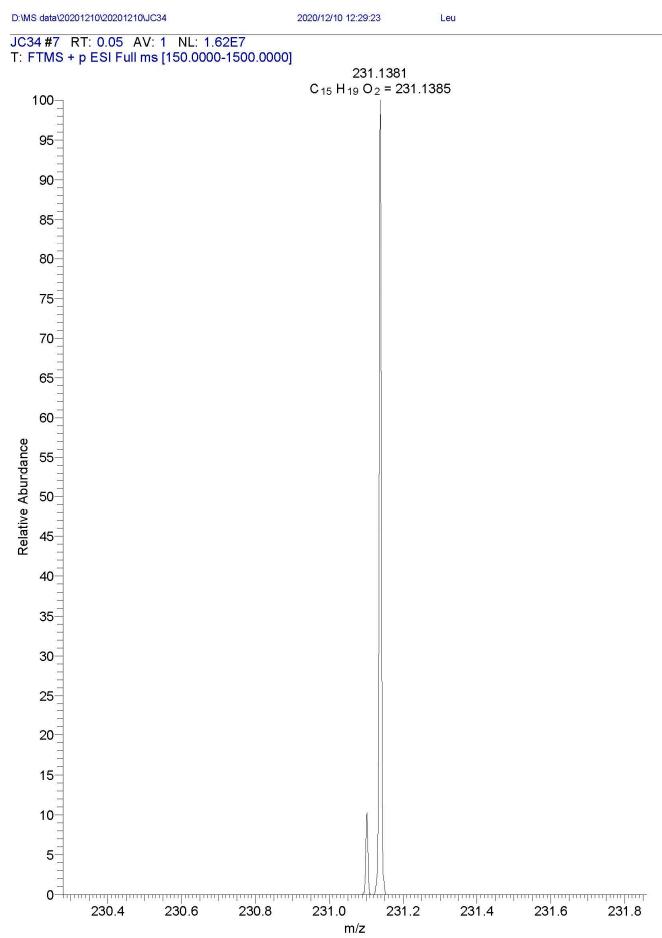
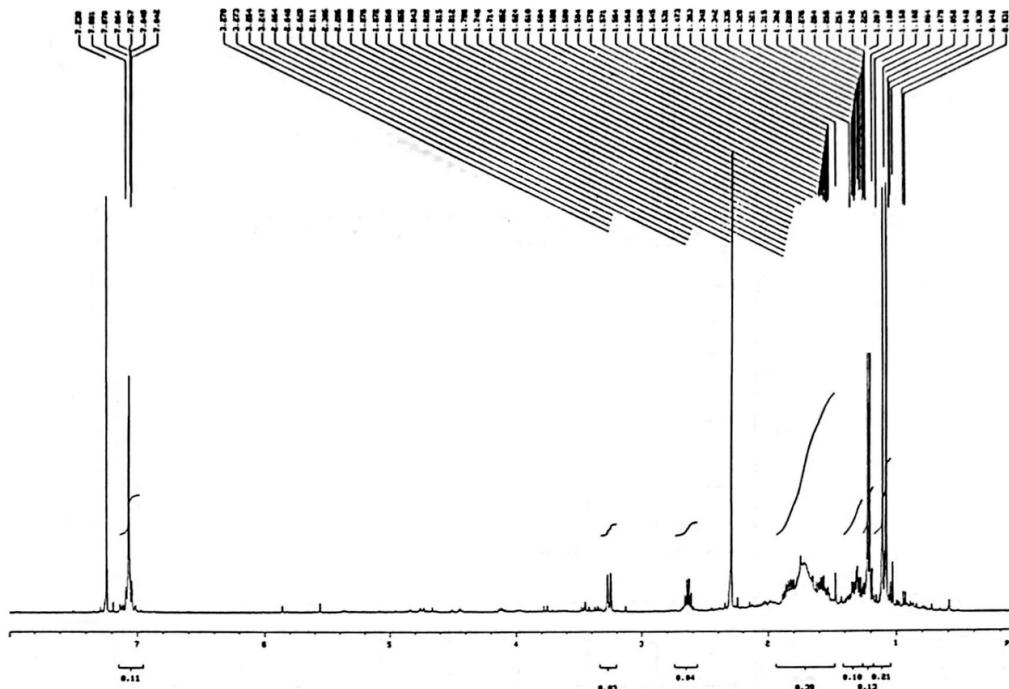


Figure S9. HRESIMS spectrum of  $\alpha$ -corocalen-15-oic acid (**3**)



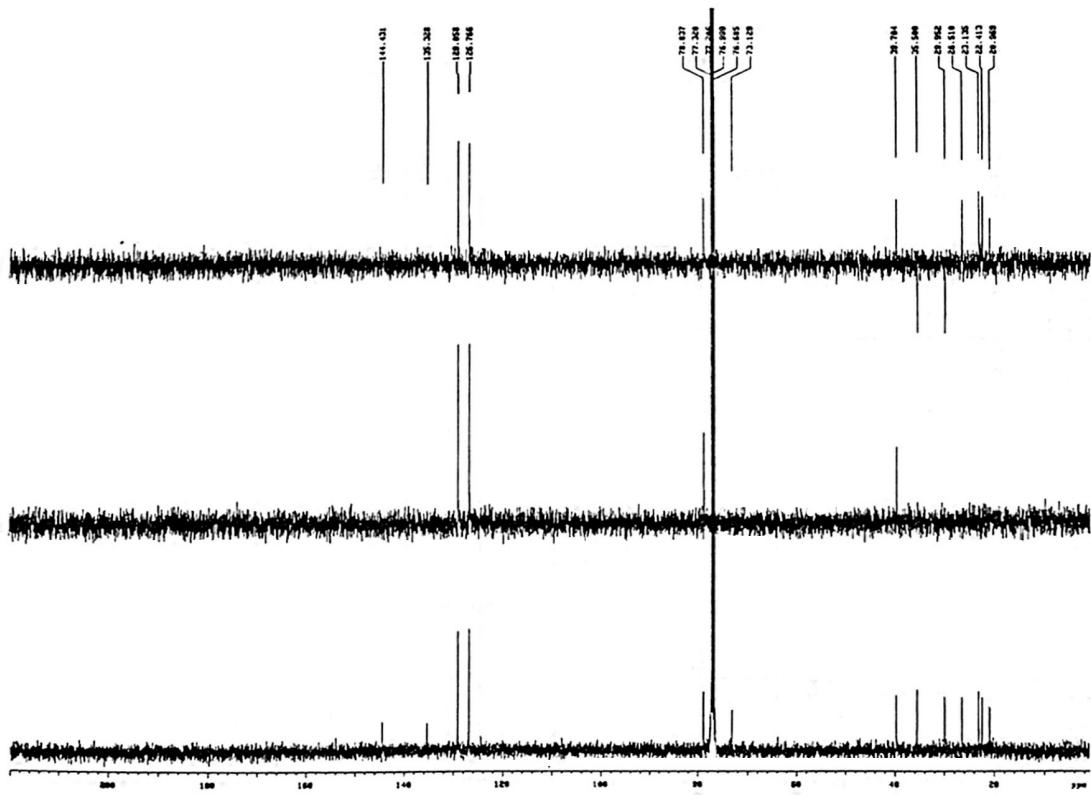


Figure S11. DEPT spectrum of 1,3,5-bisabolatrien-10-hydroperoxy-11-ol (4)

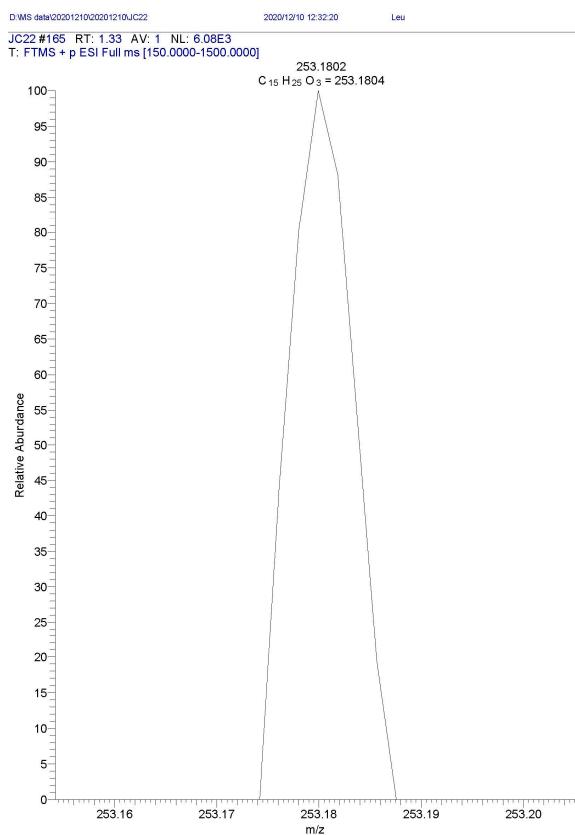


Figure S12. HRESIMS spectrum of 1,3,5-bisabolatrien-10-hydroperoxy-11-ol (4)

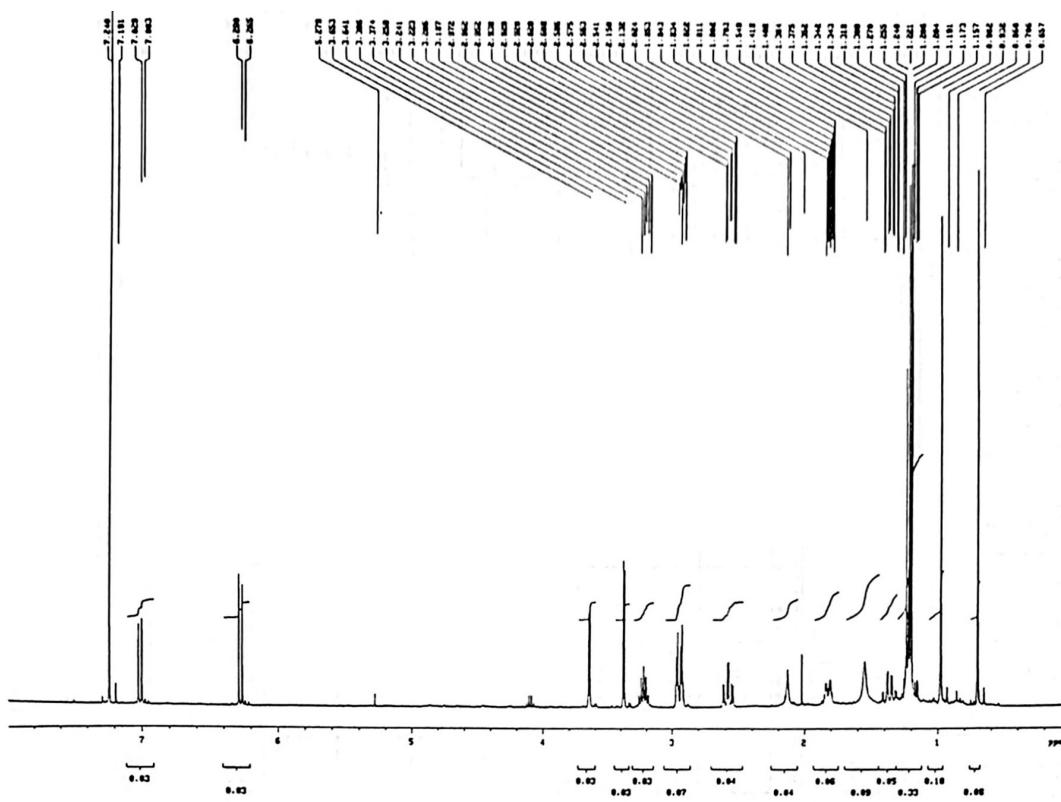


Figure S13. <sup>1</sup>H NMR spectrum of 1 $\beta$ ,2 $\beta$ -epoxy-9 $\alpha$ -hydroxy-8(14),11-totaradiene-3,13-dione (**5**) in  $\text{CDCl}_3$  at 400 MHz

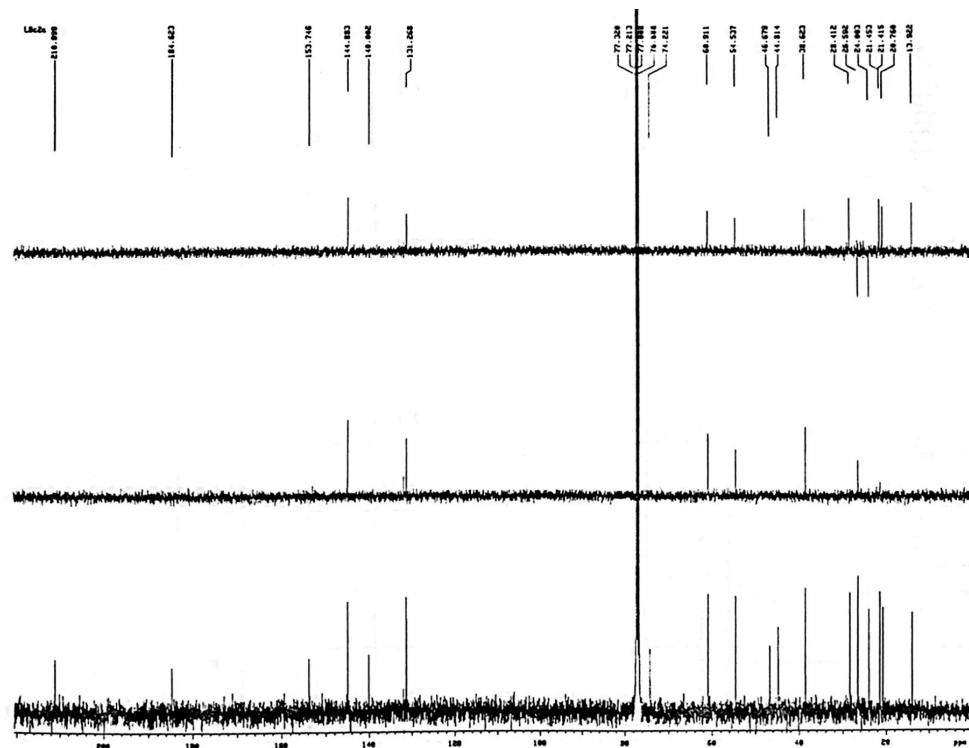


Figure S14. DEPT spectrum of 1 $\beta$ ,2 $\beta$ -epoxy-9 $\alpha$ -hydroxy-8(14),11-totaradiene-3,13-dione (**5**) in  $\text{CDCl}_3$  at 400 MHz

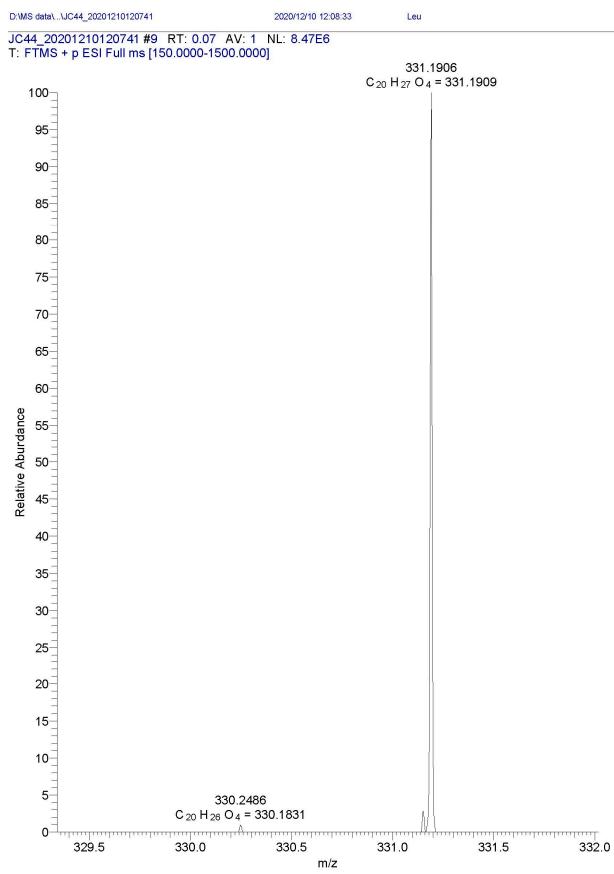


Figure S15. HRESIMS spectrum of  $1\beta,2\beta$ -epoxy- $9\alpha$ -hydroxy- $8(14),11$ -totaradiene- $3,13$ -dione (**5**) in  $CDCl_3$  at 400 MHz