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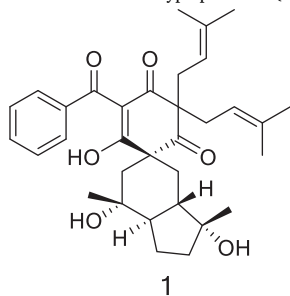
Corrigendum

Corrigendum to “Hyperpatulols A-I, spirocyclic acylphloroglucinol derivatives with anti-migration activities from the flowers of *hypericum patulum*” [Bioorg. Chem. 87 (2019) 409–416]

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1. The structure of hyperpatulol A (**1**) please delete



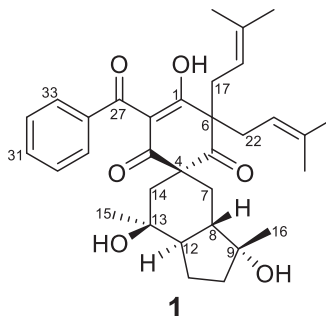
2. In the section “Structure elucidation” (Pg. 412, Line 24) please delete

Comparing their ^{13}C NMR data disclosed the differences in signals of C-1 and C-3 between **1** and hyperbeanol C. The shielded chemical shifts of C-1 (δ_{C} 193.3 in **1**; δ_{C} 196.9 in hyperbeanol C) and C-3 (δ_{C} 193.3 in **1**; δ_{C} 195.6 in hyperbeanol C) suggested that the enol group at C-1 and the carbonyl group at C-3 in hyperbeanol C were exchanged in **1** [20–23] (Figure 2). This deduction was further confirmed by the correlations from H-7a (δ_{H} 1.82) and H-14a (δ_{H} 2.35) to C-3 (δ_{C} 193.3), C-4 (δ_{C} 62.5), and C-5 (δ_{C} 213.9), and from H₂-17 (δ_{H} 2.83) and H-22a (δ_{H} 2.68) to C-1 (δ_{C} 193.3), C-5 (δ_{C} 213.9), and C-6 (δ_{C} 61.4) in its HMBC spectrum.

3. In the section “Structure elucidation” (Pg. 412, Line 38) please delete

Subsequently, the obvious ROE correlations of H-8/H-14a, Me-15/H-14a, and H-8/Me-15 revealed that H-8 and H-14a were β -oriented, indicating the *trans*-fused conformation of the cyclohexane and cyclopentane moieties. The β -orientation for Me-15 and Me-16 was confirmed by the observed ROE correlations of H-8/Me-16, H-8/Me-15, H-14a/Me-16, and H-14a/Me-15. In addition, the ROE interactions of H-29 with H-14a and Me-15 revealed that the benzoyl group was at the upper side of the cyclohexane moiety. Therefore, the structure of **1** was defined.

Substitute with



Substitute with

Comparing their ^{13}C NMR data disclosed the differences in signals of Me-15 between **1** and hyperbeanol C. The deshielded chemical shift of Me-15 (δ_{C} 29.1 in **1**; δ_{C} 21.6 in hyperbeanol C) suggested that they could be C-13 epimers (Figure 2) [19].

Substitute with

Subsequently, the obvious ROESY correlations of HO-13 (δ_{H} 5.30)/H-8, H-8/H-14a and H-8/Me-16 revealed that HO-13, H-8, Me-16 and H-14a were β -oriented, indicating the *trans*-fused conformation of the cyclohexane and cyclopentane moieties. The ROESY correlation of Me-15/H-14b indicated the α -orientation for Me-15. The α -orientation for H-12 was same as hyperbeanol C determined by the ^{13}C NMR signal of C-12 (δ_{C} 51.1) [19, 20]. Therefore, the structure of **1** was defined.

[20] Y.R. Li, W.J. Xu, S.S. Wei, W.J. Li, J. Luo, L. Y. Kong, *Phytochemistry*. 159 (2019) 56–64, <https://doi.org/10.1016/j.phytochem.2018.12.005>.

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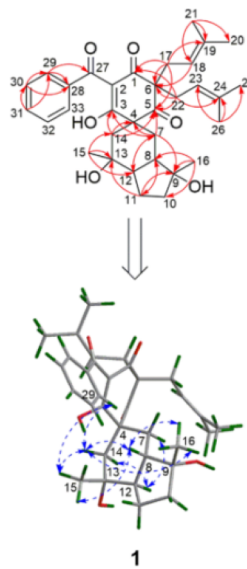
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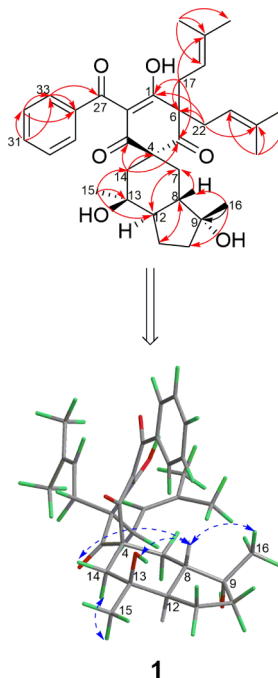
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4. In the section "Structure elucidation" (Pg. 413, Fig. 2) please delete



Substitute with



The authors would like to apologize for these errors and any consequent inconvenience caused.