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**Brief Overview of Recently Reported Misassigned Natural Products and Their in Silico Revisions Enabled by DU8ML, a Machine Learning-Augmented DFT Computational NMR Method.**

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## Highlight

## Brief Overview of Recently Reported Misassigned Natural Products and Their *in Silico* Revisions Enabled by DU8ML, a Machine Learning-Augmented DFT Computational NMR Method.

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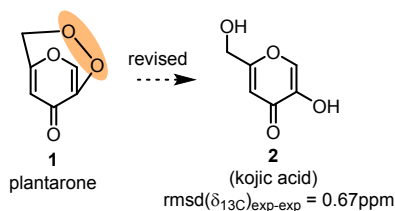
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This *Highlight* article describes a personal selection of recent misassigned structures of natural products and their revision with the aid of DU8ML, a machine learning-augmented DFT computational method for fast and accurate calculations of solution NMR chemical shifts and spin-spin coupling constants.

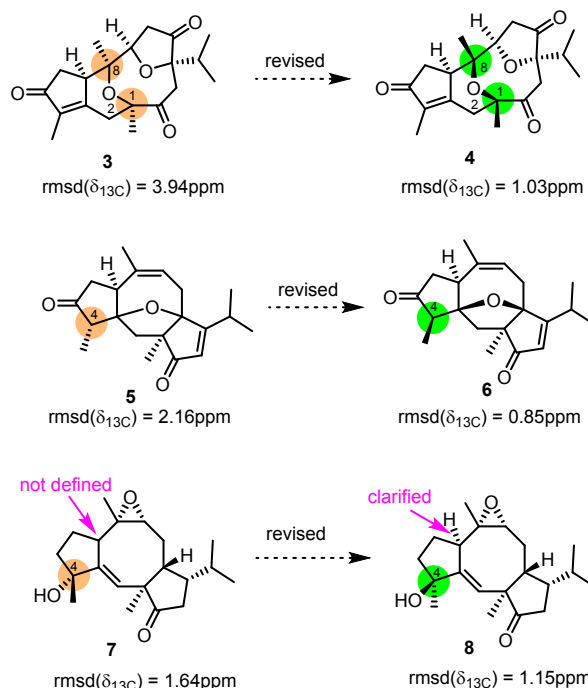
It is self-evident for the broader readership of this journal that natural product research is extraordinarily consequential. Among other things, it provides one of the most prolific pipelines for drug discovery. In this context it is critically important to determine the structures of NPs with utmost accuracy. Solution structure elucidation principally relies on NMR as the most informative method for non-crystallizable materials. There has been spectacular progress with new solution NMR techniques.<sup>1</sup> Yet, structure misassignments remain ubiquitous, perhaps excessively so. This brief overview of recent literature is intended to highlight such misassignments, offer revised structures, and further stimulate community discussions about pitfalls and common errors in structure elucidation. For this work we rely on our recently developed machine learning-augmented DFT computational NMR method, DU8ML.<sup>2</sup> Hopefully, such overviews will become a recurring feature in *Nat. Prod. Rep.*

At times, misassignments are grotesque and easy to spot. Recently reported new pyran derivative with antioxidant and anticancer properties, plantarone (**1**),<sup>3</sup> was flagged by the "Hot off the Press" reviewers Hill and Sutherland,<sup>4</sup> who called it "impossibly strained" and requiring revision. DU8ML analysis pointed to the known kojic acid (**2**), although even without computations the comparison of published *experimental* <sup>13</sup>C chemical shifts for plantarone and kojic acid, albeit in different solvents, produced  $\text{rmsd}_{[\text{exp-exp}]}$  of 0.67ppm.



In contrast, it is by far more difficult to detect structure misassignments in cases when imperceptible errors involve a small number of stereogenic centers. Such subtle cases of misassignment are most frequent. Computational methods – including DU8ML – could expeditiously identify problems and help guide the process of structure revision.

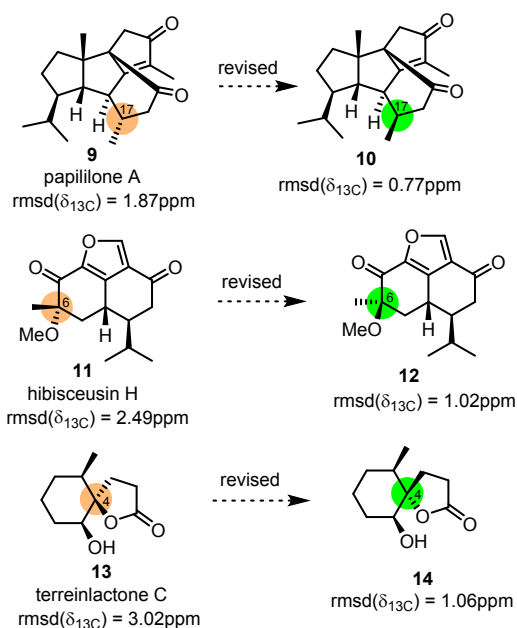
For example, of ten new fusicoccane diterpenes recently isolated from the roots of *H. forsskaolii*,<sup>5</sup> at least three required revisions of stereoconfiguration for one or two stereogenic centers: compound **3** needed revision to diastereomer **4**; compound **5** was revised to 4-epimer **6**; and compound **7** was revised to 4-epimer **8** with stereoconfiguration at C6 – which was not defined in the original publication – clarified as shown. Notice that in all revisions *relative* stereochemistry is implied.



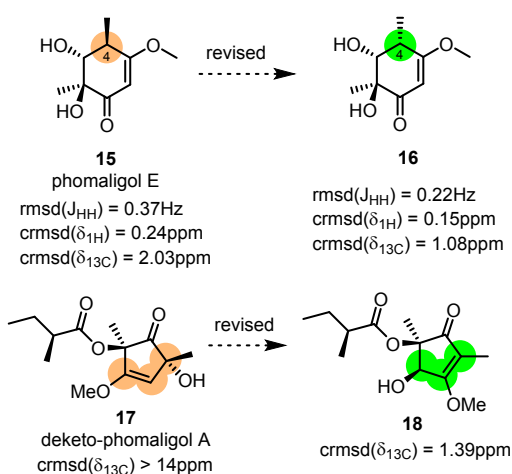
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In another recent study, secondary metabolites from *Papiliomyces* sp were investigated and the first tetracyclic diterpenoid with an unprecedented 5/5/5/6 ring system, papililone A (**9**), was reported.<sup>6</sup> While the authors augmented its structure elucidation by both HOSE and Neural Net methods for <sup>13</sup>C chemical shift prediction, our DU8ML analysis still necessitated revision to 17-epimer **10**.

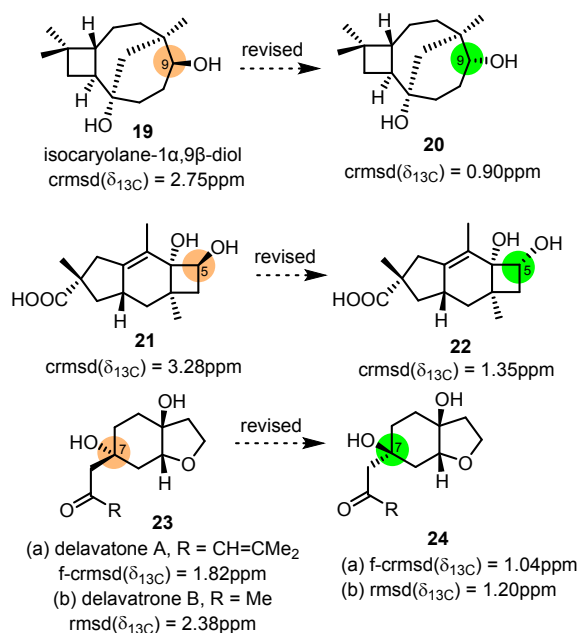
Eight new cadinene sesquiterpenoids, hibisceusins A-H, were isolated from infected stems of the semi-mangrove plant, *Hibiscus tiliaceus*.<sup>7</sup> Structures of five of them were determined by X-ray crystallographic analysis. Yet according to DU8ML, hibisceusin H (**11**), characterized only by solution NMR, required revision to its 6-epimer **12**. The new spirocyclic lactone, terreinlactone C (**13**),<sup>8</sup> isolated from the fungus *Aspergillus terreus*, also needed revision to 4-epimer **14**.



Culture broth from marine-derived fungus, *Aspergillus flocculosus*, produced two new compounds, phomaligol E (**15**), and deketo-phomaligol A (**17**).<sup>9</sup> Our analysis showed that phomaligol E (**15**), needs revision to 4-epimer (**16**), while deketo-phomaligol A (**17**) required a more extensive revision into  $\alpha,\beta$ -unsaturated cyclopentanone **18**.



Carbon-13 NMR data for isocaryolane-1 $\alpha$ ,9 $\beta$ -diol (**19**), extracted from aerial parts of the plant *Perovskia artemisioides*<sup>10</sup> produced poor match with the computed chemical shifts. It was revised to 9-epimer **20**. A sterpurane derivative **21** isolated from a coculture of *P. orientoasiaticus* and *X. flaviporus* required similar single-stereogenic center revision to 5-epimer **22**.<sup>11</sup>



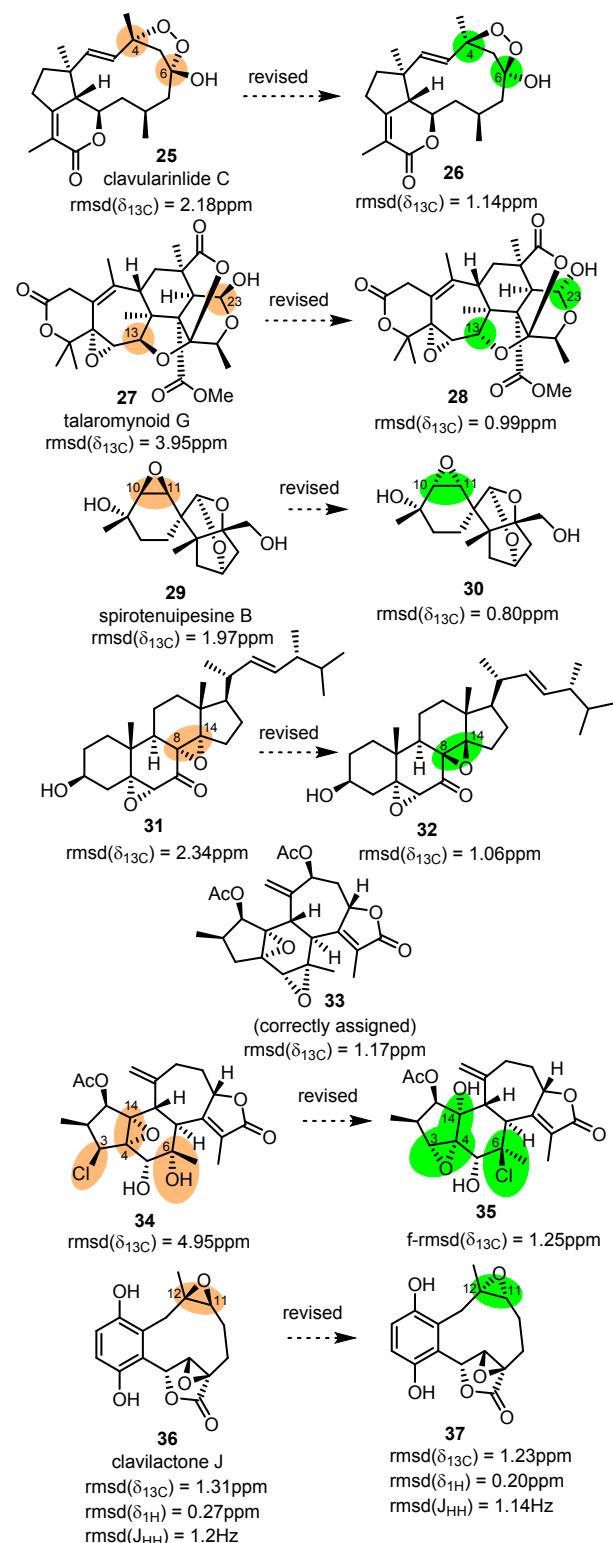
Delavatone A (**23a**, R = CH=CMe<sub>2</sub>) and its close relative, delavatrone B (**23b**, R = Me) were isolated from the ethanol extract of whole plant of *Incarvillea delavayi* and exhibited "remarkable suppression on NO production in LPS-induced BV-2 cells."<sup>12</sup> Their structures were elucidated by NMR "with the help of chemical shift calculations coupled with DP4+ probability analysis". Yet, DU8ML necessitated the revision of stereoconfiguration at C7 to structures **24** with the calculated chemical shifts matching the experimental data very well.

Clavularinlide C (**25**)<sup>13</sup> isolated from soft coral *Clavularia inflata* needed revision to diastereomer **26**. Nine new highly oxygenated

3,5-dimethylorsellinic acid-derived meroterpenoids, talaromynoids A-I, were isolated from the marine-derived fungus *Talaromyces purpureogenus* SCSIO 41517.<sup>14</sup> All of them possessed unprecedented polycyclic ring systems, with talaromynoid A characterized with X-ray crystallographic analysis. However, talaromynoid G (**27**), needed revision to diastereomer **28**.

Recent synthesis of spirotenuipesine A, a promoter of neurotrophic factor secretion from glial cells<sup>15</sup> prompted us to look at the original isolation work for spirotenuipesine A and its epoxide, spirotenuipesine B, which was assigned structure **29**.<sup>16</sup> DU8ML analysis necessitated revision of spirotenuipesine B to a stereoisomeric epoxide **30**.<sup>17</sup> As we reported in the past, complex epoxides often present significant challenge to accurate determination of their stereochemistry.<sup>18</sup> Another recent example of this is bis-epoxy ergostadienediol.<sup>19</sup> The authors reassigned an incorrect pentahydroxy ergostane

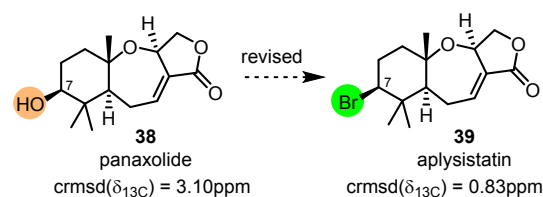
structure to bis-epoxide **31**, because it matched the experimental chemical shifts for this bis-epoxide reported a decade ago.<sup>20</sup> The problem is that this revision was incomplete and, according to DU8ML, required additional correction of stereoconfiguration for the second oxirane moiety, i.e. to diastereomer **32**.



Two new anti-HIV diterpenoids **33** and **34** were isolated from *C. megalocarpus*, with chloride **34** described as a potential artifact of extraction.<sup>21</sup> According to our analysis, diepoxide **33** is assigned correctly. However, chloride **34** needed significant revisions to **35**.

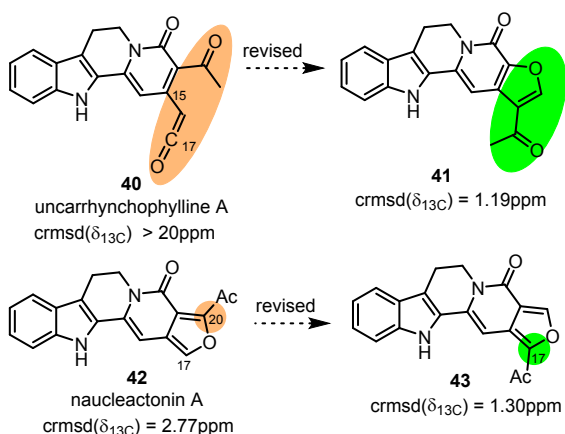
The case of meroterpenoid clavilactone J (**36**), recently isolated from the basidiomycete *Clitocybe clavipes*,<sup>22</sup> further underscores significant difficulty that conformationally flexible NPs possessing the oxirane moiety pose for structure elucidation. The authors calculated DP4+ all-data probabilities for the candidate structures, which pointed with high confidence (99.87%) to *11S,12R* diastereomer **36**, although the H data-based probability disagreed with C data. DU8ML analysis showed very small differences between the two diastereomers. However, all criteria that we used, i.e.  $^{13}C$  and  $^1H$  NMR chemical shifts, and proton spin-spin coupling constants matched the *11R,12S* diastereomer **37** slightly better. We therefore revise clavilactone J to structure **37**.

Another challenge for the *in silico* structure elucidations present the cases where computer-driven revisions are not in keeping with the published interpretation of mass-spectrometry data. One such example is a new sesquiterpene lactone panaxolide (**38**) which was recently isolated from the leaves of Vietnamese ginseng found in the Ngoc Linh mountain range.<sup>23</sup> While most of the calculated chemical shifts closely tracked the experimental values, the C7 carbon bearing the hydroxy group was singularly off by >10ppm, contributing to poor overall rmsd of 3.1ppm. Replacing the 7-OH group by bromine improved rmsd to 0.83ppm, indicating that panaxolide should be revised to **39**, which implies misinterpretation of the MS data. Luckily, bromide **39** is a known compound, aplysiastatin, first isolated in 1977 from a sea hare, with structure established by X-ray crystallography,<sup>24</sup> imparting confidence that DU8ML has arrived at the correct revision. We are not certain how a brominated marine NP ended up in Vietnamese ginseng found at high elevation.

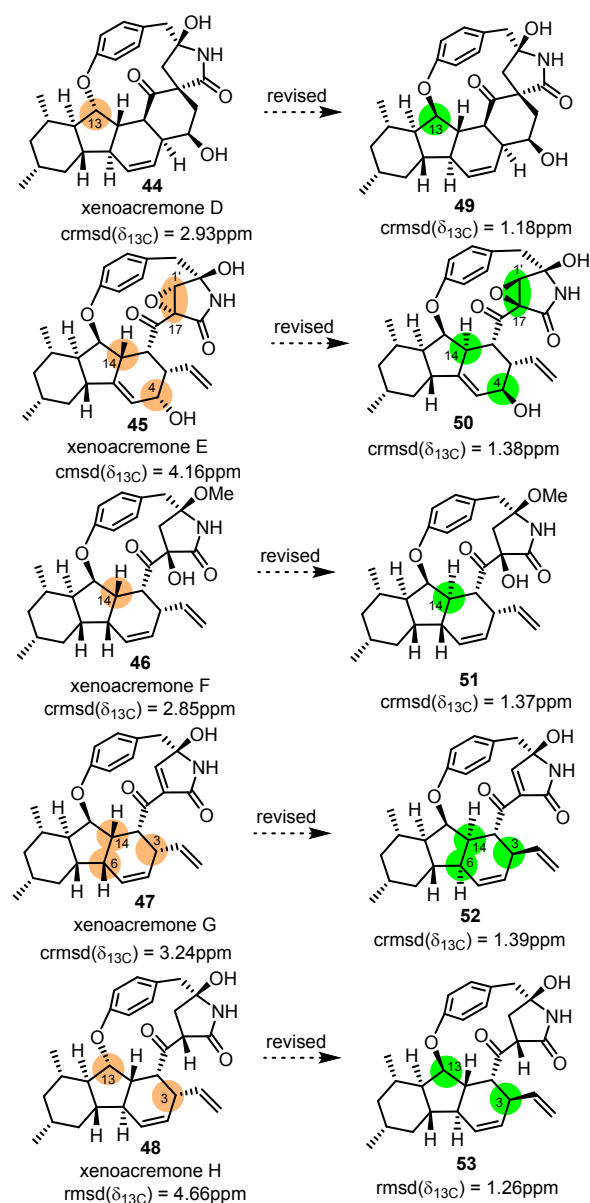


Highly reactive moieties offer another red flag of structure misassignment. An unusual alkaloid, uncarrhynophylline A (**40**), purportedly containing a ketene moiety was isolated from stems of *Uncaria rhynchophylla*.<sup>25</sup> DU8ML analysis showed irreconcilable problems with this structure, which we revised to acetylfuran **41**.

Revised uncarrhynophylline A (**41**) is a new unreported structure. However, an alkaloid with a high degree of similarity, naucleactonin A (**42**),<sup>26</sup> was previously described. Instructively, according to DU8ML, naucleactonin A also required revision to **43**, i.e. relocation of the acetyl group from C20 to C17.



Finally, a group of bioactive tyrosine-decahydrofluorene analogues, xenoacremones D-H (**44-48**), were recently isolated from the plant-derived fungus *Xenoacremonium sinensis*.<sup>27</sup> While the structure connectivities in these complex NPs were determined correctly, stereoconfiguration of one or more stereogenic centers in each of these NPs required revision.



The case of xenoacremones is particularly challenging because of structure distortions related to the *p*-methylenephenoxy bridge linking positions 13 and 2'. The cyclic C2'-hemiaminal moiety further aggravated the situation. DU8ML-guided analysis helped us revise xenoacremone D (**44**) to 13-epimer **49**; xenoacremone E (**45**) – to diastereomer **50**; xenoacremone F (**46**) – to 14-epimer **51**; xenoacremone G (**47**) – to diastereomer **52**; and xenoacremone H (**48**) – to diastereomer **53**. It appears that stereoconfiguration of centers C7, C9, C11, C12, and C13 is "conserved," unlike the other stereogenic centers, e.g., C3, C6, C14, and C15.

## Conclusions

DU8ML offers ample cost-benefit outcomes, allowing for fast and accurate computations of NMR parameters, and thus enabling *high-throughput* validation or revision of published structures on a massive scale. However, no method of structure elucidation is perfect. While most of the computational NMR methods these days are reasonably reliable in flagging potential misassignments, the proposed revisions are what they are – the most likely structures. This overview is an invitation to a discussion of how to make structure elucidation by NMR more reliable and learn to recognize most challenging aspects of assigning connectivity and stereochemistry in newly isolated natural products.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

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**Biographies**

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