



Corrigendum to “Approaches to the total synthesis of phomactins”

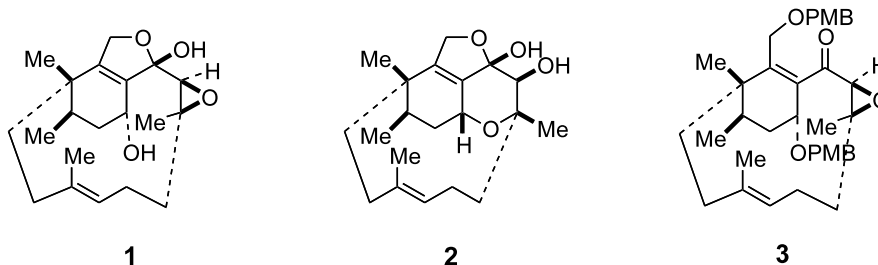
[Tetrahedron Lett. 44 (2003) 2713][☆]

Andrew S. Balnaves, Graham M^cGowan, Peter D. P. Shapland and Eric J. Thomas*

The Department of Chemistry, The University of Manchester, Manchester M13 9PL, UK

Footnote 5 of the above *Tetrahedron Letter* states that “The spontaneous rearrangement of Sch. 49028 into phomactin A was observed by Pattenden et al. during their synthesis, see ref. 4.¹” However, it should be made absolutely clear that Goldring and Pattenden did not state in their paper¹ that Sch. 49028 spontaneously rearranged into phomactin A. These authors did not isolate Sch. 49028 (still attributed the originally assigned² hydroxy-epoxide structure **1** in the above *Tetrahedron Letter*) and could not therefore study its rearrangement.

In the final step of their synthesis of phomactin A **2**,¹ Goldring and Pattenden deprotected the bis-*p*-methoxybenzyl ether **3** and isolated phomactin A **2** directly. They did not isolate the hydroxyepoxide **1** (Sch. 49028?) although it may be an intermediate in the conversion of **3** into **2**. Moreover, there is some doubt as to whether the compound identified as Sch. 49028 was correctly identified originally² as the hydroxyepoxide **1** or whether it should have been identified as phomactin A **2** since the NMR spectrum reported for Sch. 49028² in deuterated chloroform was found by both Pattenden¹ and Halcomb³ to correspond to that of synthetic phomactin A **2**.



References

1. Goldring, W. P. D.; Pattenden, G. *Chem. Commun.* **2002**, 1736.
2. Chu, M.; Truumees, I.; Gunnarsson, I.; Bishop, W. R.; Kreutner, W.; Horan, A. C.; Patel, M. G.; Gullo, V. P.; Puar, M. S. *J. Antibiot.* **1993**, 46, 554.
3. Mohr, P. J.; Halcomb, R. L. *J. Am. Chem. Soc.* **2003**, 125, 1712.

[☆] PII of original article: S0040-4039(03)00341-1

* Corresponding author. Tel.: 0044 (0)161 275 4614; fax: 0044 (0)161 275 4939; e-mail: e.j.thomas@man.ac.uk