

Addition/Correction

**A Likely Biogenetic Gateway Linking 2-Aminoimidazolinone
Metabolites of Sponges to Proline: Spontaneous Oxidative
Conversion of the Pyrrole-Proline-Guanidine Pseudo-peptide to
Dispacamide A [J. Am. Chem. Soc. 2004, 126, 10252–10253].**

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J. Am. Chem. Soc., **2005**, 127 (29), 10454-10454 • DOI: 10.1021/ja053215n • Publication Date (Web): 29 June 2005

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A Likely Biogenetic Gateway Linking 2-Aminoimidazolinone Metabolites of Sponges to Proline: Spontaneous Oxidative Conversion of the Pyrrole-Proline-Guanidine Pseudo-peptide to Dispacamide A [*J. Am. Chem. Soc.* **2004**, *126*, 10252–10253]. Nathalie Traver and Ali Al-Mourabit*

Page 10252. In Scheme 1 and in the Supporting Information, dispacamide A (**2**) should be the (*Z*)-isomer and not the (*E*)-isomer, as drawn.

In the Supporting Information, **12**,AcO₂H should read as **12**,CF₃O₂H.

We thank Professor David A. Horne of Oregon State University for pointing out these errors.

Three syntheses of dispacamide have been reported but not referenced. Thus, the fourth sentence of the second paragraph should read as follows: Ornithine and proline have been respectively used in the synthesis of “oroidin-based” dibromophakellin by Büchi,^{8a} dispacamide A by Horne,^{8b} and dibromophakellstatin by Romo.⁹

Reference 8 should then read as follows:

(8) (a) Foley, L. H.; Büchi, G. *J. Am. Chem. Soc.* **1982**, *104*, 1776–1777. (b) The method has been developed by Horne for the synthesis of dispacamide A from the available 2-aminoimidazole derivative: Olofsen, A.; Yakushijin, K.; Horne, D. A. *J. Org. Chem.* **1998**, *63*, 1248–1253. For other syntheses, see: (c) Lindel, T.; Hoffmann, H. *Tetrahedron Lett.* **1997**, *38*, 8935–8938. (d) Fresneda, P. M.; Molina, P.; San, M. A. *Tetrahedron Lett.* **2001**, *42*, 851–854.

JA053215N

10.1021/ja053215n

Published on Web 06/29/2005