Addition/Correction

Synthesis and Biological Evaluation of Dimeric RGD Peptide–Paclitaxel Conjugate as a Model for Integrin-Targeted Drug Delivery.

Xiayuan Chen, Carmen Plasencia, Yingping Hou, and Nouri Neamati

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Additions and Corrections

2005, Volume 48

Xiayuan Chen,* Carmen Plasencia, Yingping Hou, Nouri Neamati: Synthesis and Biological Evaluation of Dimeric RGD Peptide−Paclitaxel Conjugate as a Model for Integrin-Targeted Drug Delivery.

Pages 1099 and 9A. In Figure 1 and the table of contents, one carbonyl group on the succinate linker was incorrectly omitted. The correct version of the structure is given below.

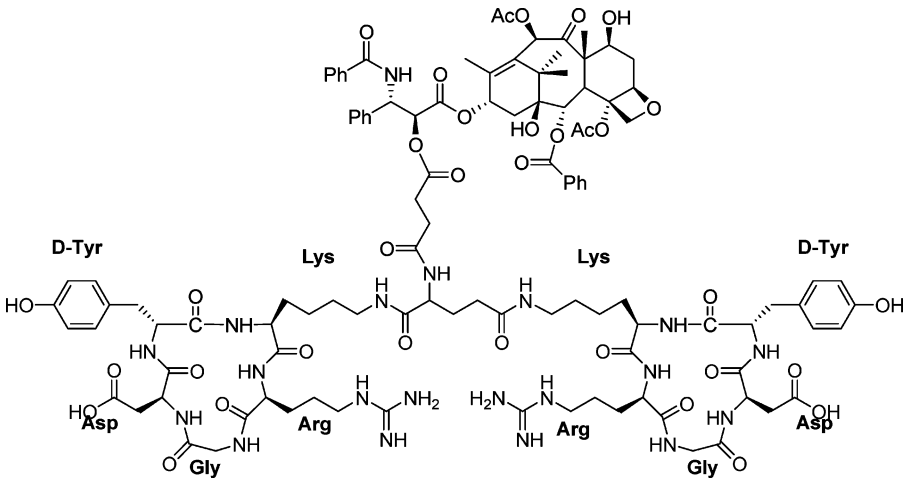

Figure 1. Schematic representation of the molecular structure of dimeric RGD peptide−paclitaxel conjugate. The succinate linkage is through the 2′-hydroxy group of paclitaxel and amino group of RGD glutamate residue.

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