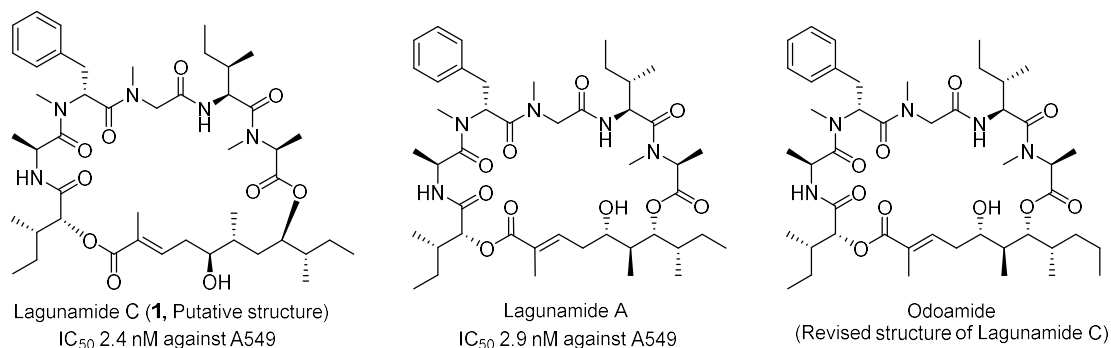


Synthesis and Structure Revision of Marine Cyanobacteria-Derived Natural Product Lagunamide C

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Lagunamide C, a 27-membered cyclodepsipeptide, was isolated from the marine cyanobacteria *Lyngbya majuscula* in 2011 by Tripathi et al.¹ and exhibits potent cytotoxicity against several cancer cells, comparable to that of the 26-membered analog lagunamide A.² The structural difference between the above compounds lies only in the presence or absence of a methylene carbon in the aliphatic acid moiety. However, it is known that the biological activity of cyclopeptides is strongly dependent on the conformation of their structure³. It should be interesting that the lagunamide families show comparable cytotoxicity regardless of the difference in the ring size of the cyclopeptide structure. Therefore, we planned the total synthesis of lagunamide C to elucidate its relationships between conformation and biological activity. In this presentation, we will report the total synthesis of the putative structure of lagunamide C and structure revision of lagunamide C to the related analog odoamide.



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