

EFFICIENT SYNTHESIS OF SPHINGOLIPID AND SPHINGOSINE-TYPE CHEMICAL MEDIATORS OF MICROBIAL ORIGIN

Daniel Lechnitz^a, Ella V. Ireland^b, Chia-Chi Peng,^a Nicole King^b,
Christine Beemelmans^a

^aChemical Biology of Microbe-Host Interactions, Hans-Knöll Institute (HKI),
07745 Jena, Germany

^bDepartment of Molecular and Cell Biology, UC Berkeley, Berkeley, CA 94720, USA

Sphingolipids and sphingosine-type compounds are widely distributed in nature; they convey a diverse set of signal transduction and stress response pathways and have profound physiological impacts.[1] Despite their recognized importance fully characterized examples of structure-activity relations are still rare. Recently, we became interested in the rosette-inducing sulfonosphingolipids, named RIFs, which induce cell differentiation in the predatory eukaryote choanoflagellate *Salpingoeca rosetta*. [2] We became particularly intrigued by the 6-hydroxy-(4*E*)-sphingenine-containing sulfonolipid named RIF-2 and its inhibitor IOR-1, both produced by the same prey bacterium. To generate the different possible stereoisomers of the (2*S*,3*S*)-6 hydroxy-(4*E*)-sphingenine-containing lipids and to determine the absolute configuration and further analyze the structure-activity relationship of these highly bioactive molecules,[3] we make use of a hydrozirconation addition reaction to Garner's aldehyde and derivatives. Here, we report about the synthesis of several unnatural (2*S*,3*S*)-6-hydroxy-(4*E*)-sphingenine-containing lipids including a fluorescence-labelled derivative for future biological studies.

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