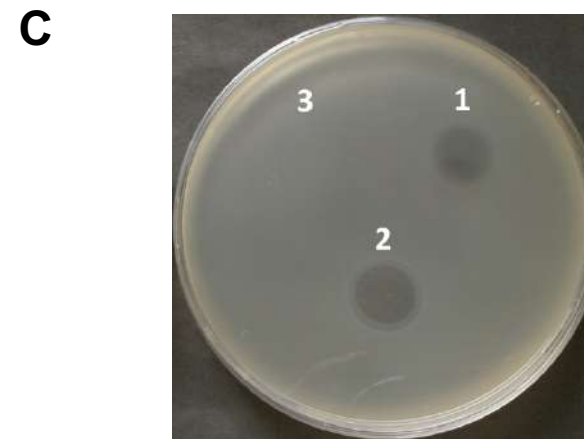
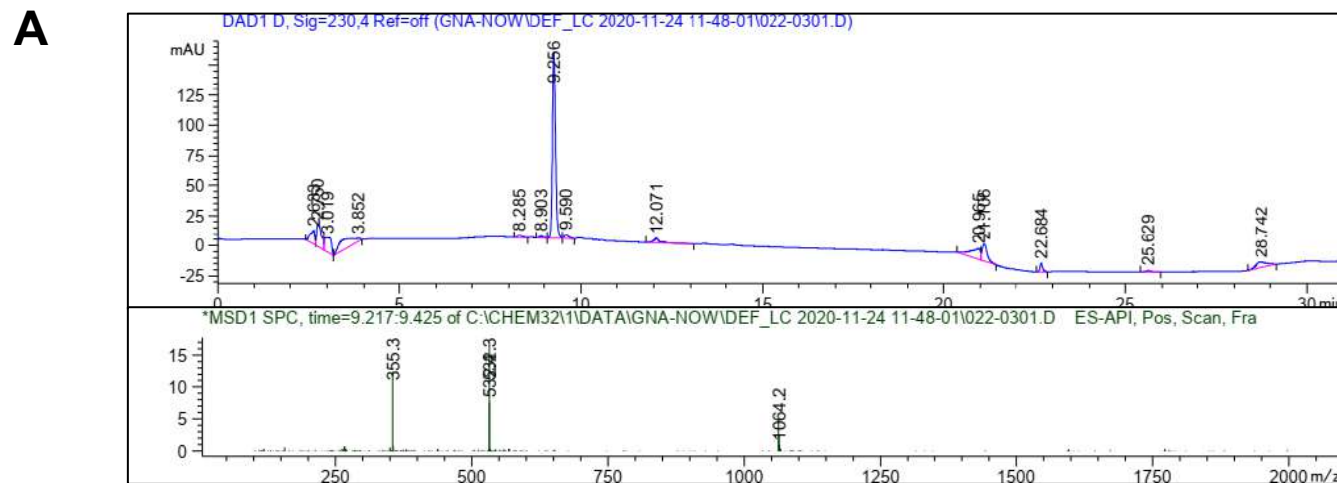


Fig. S6. Characterization of OatA activity against a NOSO-95179 analog. (A) LCMS analysis of NOSO-95179 (m/z: 1022) acetylated derivatives after 24 h of reaction with acetyl-CoA and odorhabdin acetyltransferase A (OatA): the form present was 100% mono-acetylated (m/z: 1064). (B) MS/MS fragmentation analysis of mono-acetylated NOSO-95179 (m/z: 1064): acetylation of the amine group on Dab(β OH)₂. (C) Antibacterial activity assay for NOSO-95179 after 24 h of reaction with acetyl-CoA (1), and OatA (2), or both (3) against *E. coli* ATCC2592. (D) Inhibition of synthesis of the GFP reporter protein in the *E. coli* cell-free transcription-translation system by NOSO-95179 after 24 h of reaction with acetyl-CoA, OatA, or both. Gentamicin (10 μ M) was used as a positive control. The data shown are the means of three experiments \pm SD.



B

Ion	m/z	Neutral loss from previous ion	Attribution	Assigned Structure
[M+2H] ²⁺	532			Lys-AcDab(β OH)-Ala-Gly-Orn-Pro-His-Lys-Dha
y ₈	935.5	129	Lys ₁	AcDab(β OH)-Ala-Gly-Orn-Pro-His-Lys-Dha
y ₇	777.4	158	AcDab(β OH) ₂	Ala-Gly-Orn-Pro-His-Lys-Dha
y ₆	706.4	71	Ala ₃	Gly-Orn-Pro-His-Lys-Dha
y ₅	649.4	57	Gly ₄	Orn-Pro-His-Lys-Dha
y ₄	535.3	114	Orn ₅	Pro-His-Lys-Dha
y ₃	438.3	97	Pro ₆	His-Lys-Dha

Abbreviation: **Ac**, acetyl group.

