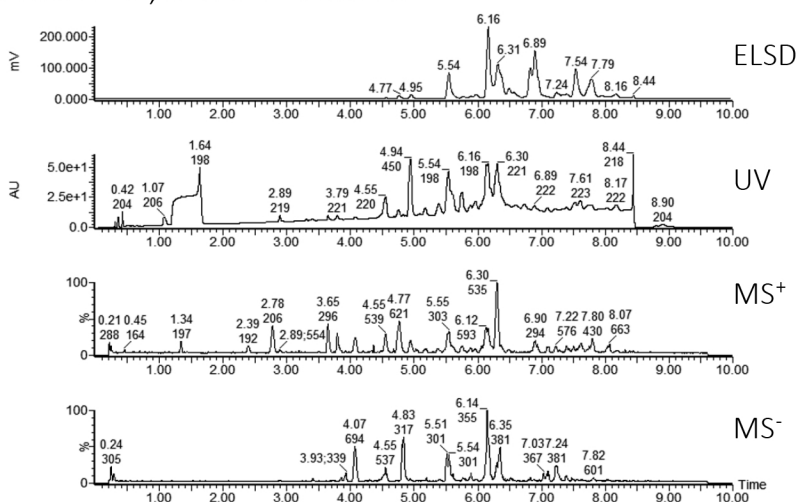
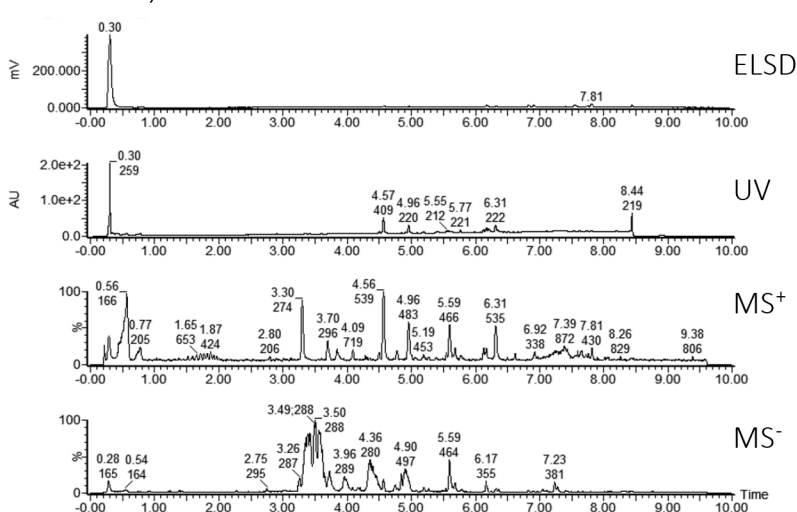


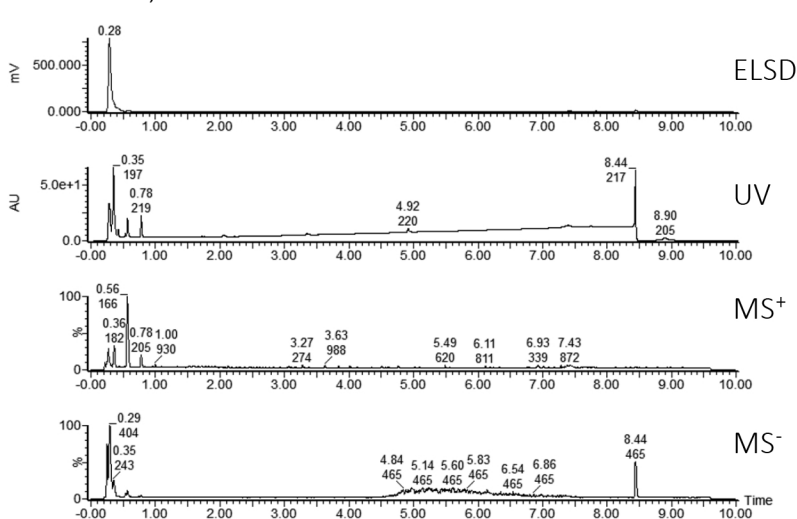
Ophiura sarsii, chloroform extract



Ophiura sarsii, ethanol extract

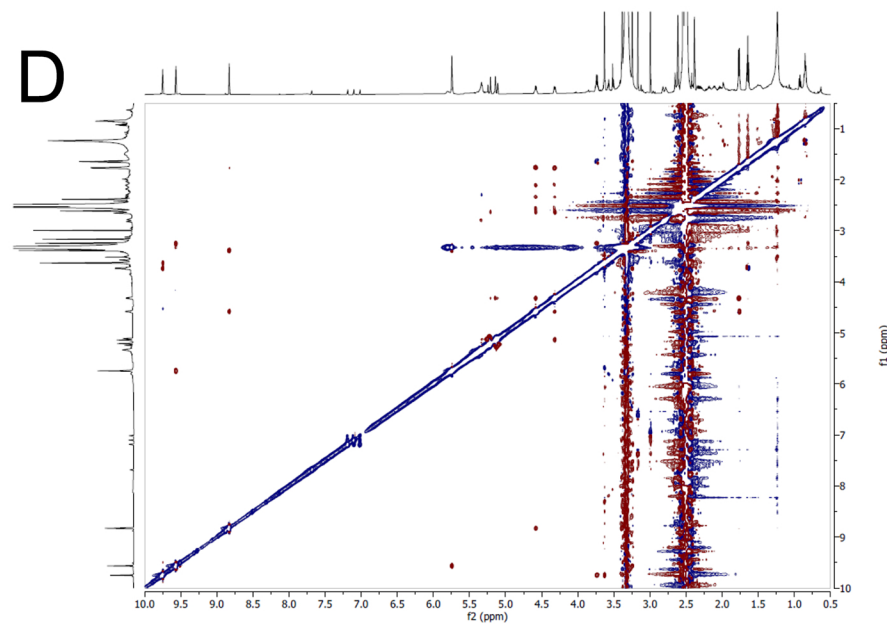
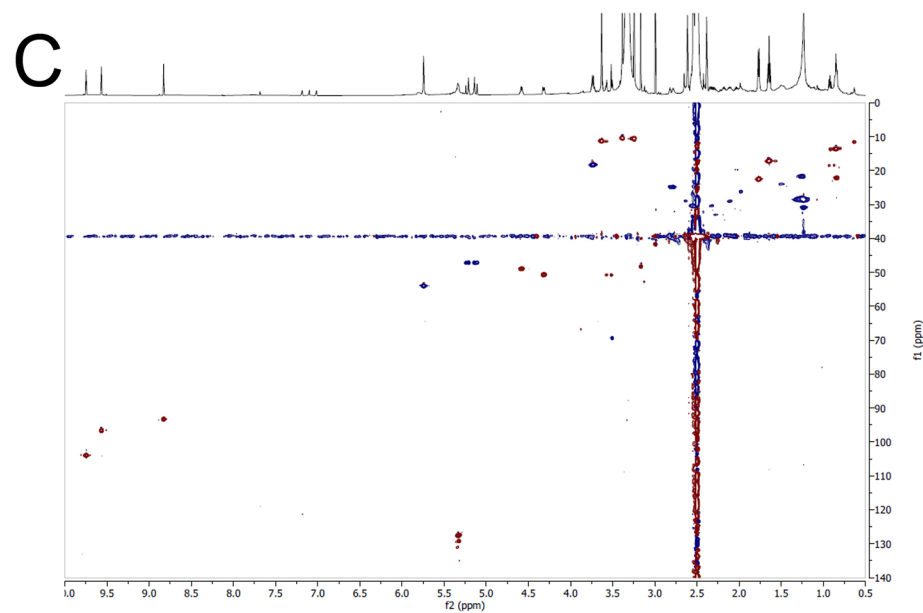
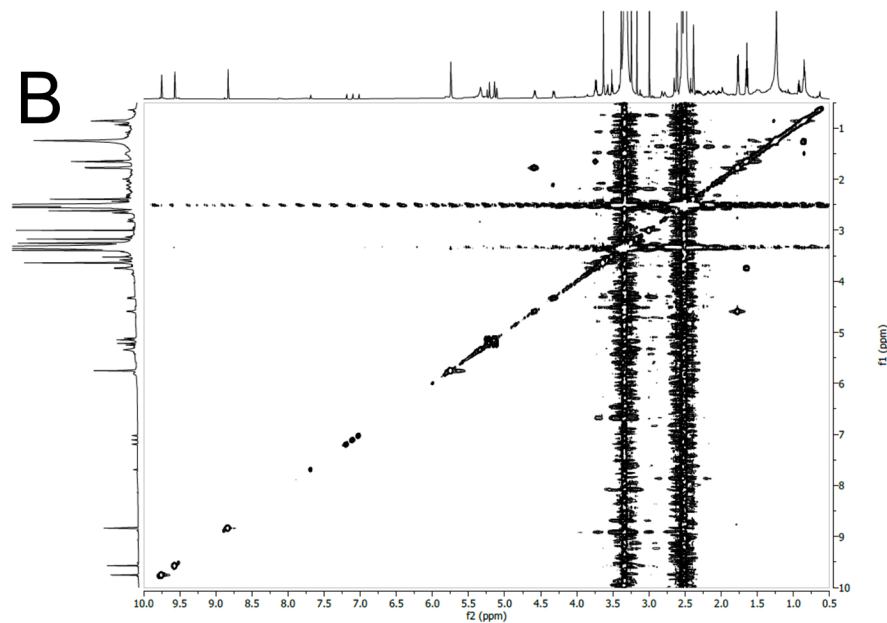
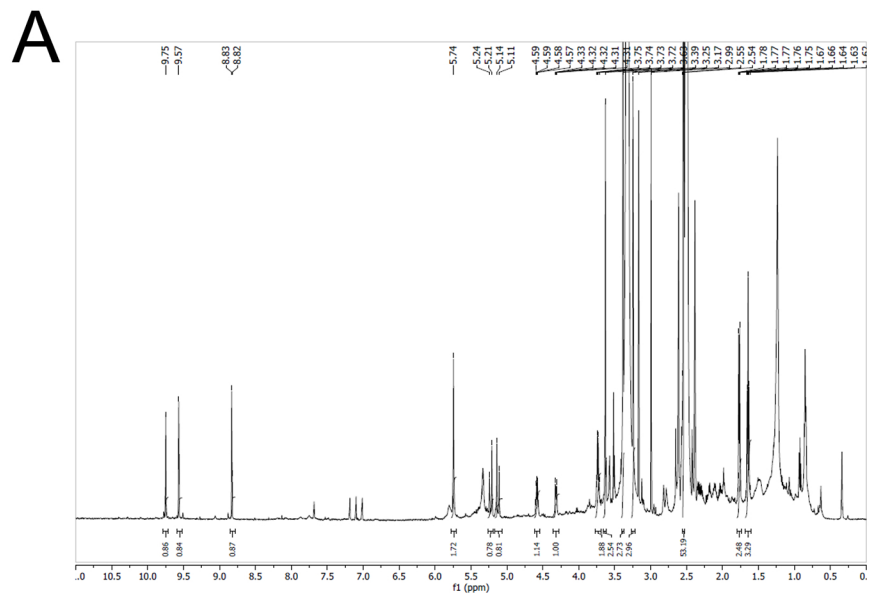


Ophiura sarsii, water extract



Klimenko *et al.*, Supplementary Figure 1.

UHPLC-PDA-ELSD-MS analysis of the initial chloroform, ethanol, and water extracts of *O. sarsii*. The ELSD trace shows relative scarcity of compounds in the water extract and abundance of compounds extracted with chloroform and ethanol.



Klimenko *et al.*, Supplementary Figure 2. Details of the NMR identification of fraction 14 as compound 1, (3S,4S)-14-Ethyl-9-(hydroxymethyl)-4,8,13,18-tetramethyl-20-oxo-3-phorbinepropanoic acid.

Supplementary Table 1: NMR chemical shifts of compound **1** in DMSO-*d*₆ and the same compound reported by Tamiaki, H., Shinkai, A. and Kataoka, Y. (2009). Synthesis of galactosylated zinc bacteriochlorophyll-d analogs and their self-aggregation in an aqueous methanol solution. *Journal of Photochemistry and Photobiology a-Chemistry* **207**, 115-125.

No	Compound 1 in DMSO- <i>d</i> ₆ at 600 MHz		Compound 1 in CD ₃ OD-CDCl ₃ reported by Tamiaki et al. 2009*
	δ _H (Multiplicity, J, nH)	δ _C	
2a	3.39 (s, 3H)	10.3	3.57 (s, 3H)
3a	5.74 (s, 2H)	53.9	5.80 (s, 2H)
5	9.57 (s, 1H)	96.7	9.42 (s, 1H)
7a	3.25 (s, 3H)	10.5	3.36 (s, 3H)
8a	3.74 (q, 7.6 Hz, 2H)	18.3	3.63 (q, 8.0 Hz, 2H)
8b	1.64 (t, 7.6 Hz, 3H)	17.1	1.64 (t, 8.0 Hz, 3H)
10	9.75 (s, 1H)	103.9	9.42 (s, 1H)
12a	3.63 (s, 3H)	11.2	3.21 (s, 3H)
13b'	5.23 (d, 19.4 Hz, 1H)	47.1	5.19 (d, 20.0 Hz, 1H)
13b''	5.12 (d, 19.4 Hz, 1H)		5.02 (d, 20.0 Hz, 1H)
17	4.32 (dt, 10.0, 2.9 Hz, 1H)	50.6	4.18 (m, 1H)
17a'	2.64 (m, 1H)	29.0	2.31-2.23, 2.00-1.97
17a''	2.11 (m, 1H)		2.31-2.23, 2.00-1.97
17b'	2.55 (m, 1H)	30.4	2.31-2.23, 2.00-1.97
17b''	2.32 (m, 1H)		2.31-2.23, 2.00-1.97
18	4.58 (qd, 7.4, 2.9 Hz, 1H)	48.9	4.42 (q, 8.0 Hz, 1H)
18a	1.77 (d, 7.4 Hz, 3H)	22.5	1.74 (d, 8.0 Hz, 3H)
20	8.83 (s, 1H)	93.2	8.53