

Supporting Information

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Anti-Inflammatory Components from the Fruits of *Amomum aromaticum*

Nguyen Hai Dang¹, Le Nguyen Thanh^{2*}, Do Hoang Giang³ and

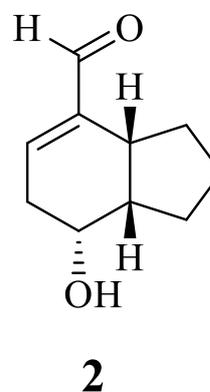
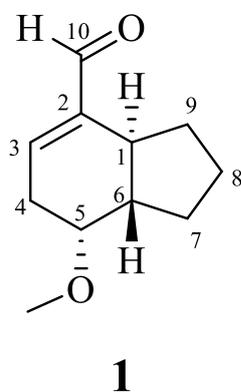
Nguyen Tien Dat³

¹University of Science and Technology of Hanoi, Vietnam Academy of Science and Technology (VAST), 18 Hoang Quoc Viet, Cau Giay, Hanoi, Vietnam

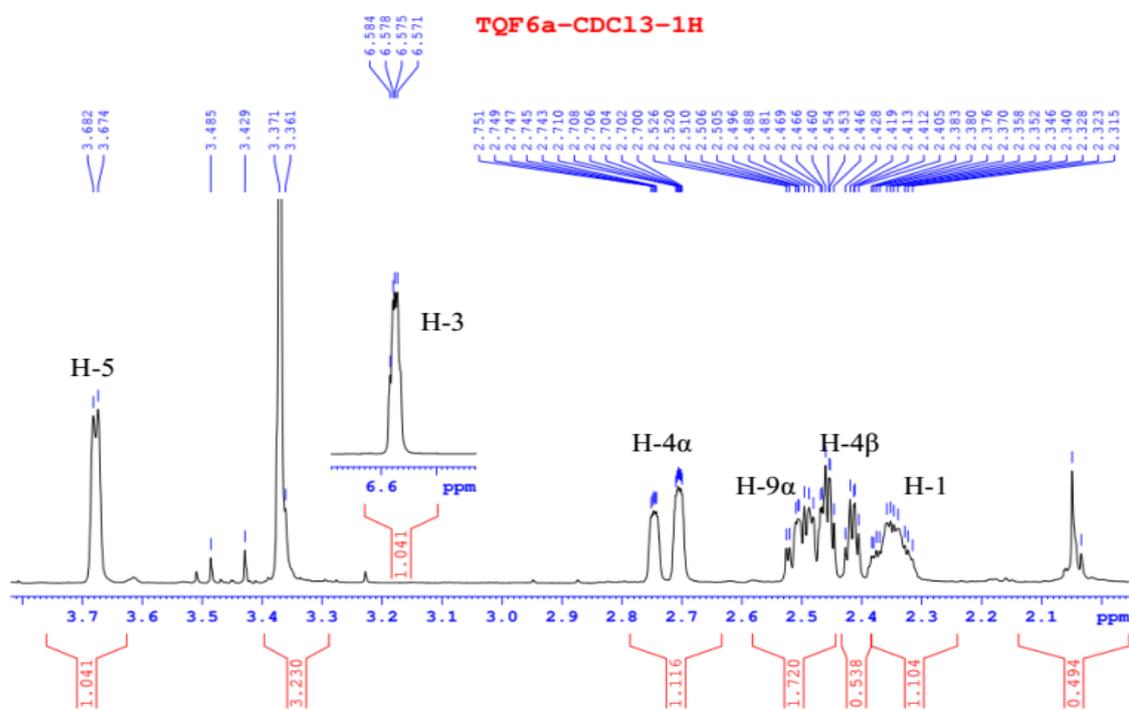
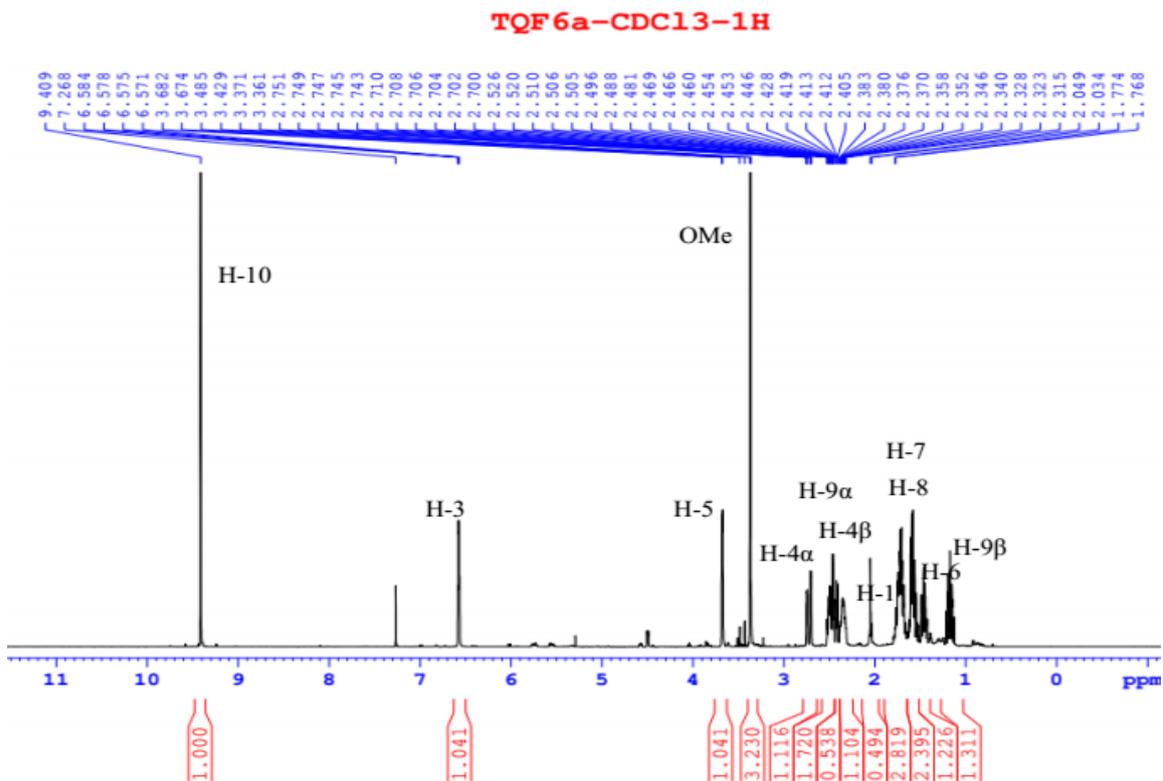
²Institute of Marine Biochemistry, VAST, 18 Hoang Quoc Viet, Cau Giay, Hanoi, Vietnam

³Center for Research and Technology Transfer, VAST, 18 Hoang Quoc Viet, Cau Giay, Hanoi, Vietnam

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Table S1: NMR spectral data of compound **1** and **2**

C	1 (CDCl ₃)		2 (CDCl ₃)	
	δ_C^a	δ_H^b (m, <i>J</i> , Hz)	δ_C^a	δ_H^b (m, <i>J</i> , Hz)
1	35.4	2.35 (1H, m, H-1)	37.5	2.95 (1H, m)
2	145.2	-	144.8	-
3	148.5	6.57 (1H, dd, <i>J</i> = 4.0 Hz, 2.0 Hz)	146.7	6.63 (1H, ddd, 5.0, 3.0, 1.0)
4	32.6	2.72 (1H, ddd, <i>J</i> = 20.5, 4.0, 2.0 Hz, H-4 α) 2.43 (1H, ddd, <i>J</i> = 20.5, 4.0, 2.0 Hz, H-4 β)	31.4	2.38 (1H, m, H-4 α) 2.43 (1H, ddd, <i>J</i> = 20.5, 4.0, 2.0 Hz, H-4 β)
5	74.4	3.68 (1H, br d, <i>J</i> = 4.0 Hz)	68.6	4.02 (1H, ddd, 9.0, 5.0, 4.5)
6	47.8	1.45 (1H, m, H-6)	43.0	2.41 (1H, m)
7	24.2	1.58 (2H, m, H-7)	25.2	1.75 (1H, m, H-7 α) 1.54 (1H, m, H-7 β)
8	22.3	1.73 (2H, m, H-8)	24.9	1.73 (2H, m, H-8)
9	27.6	2.50 (1H, m, H-9 α) 1.17 (ddd, <i>J</i> = 12.5, 10.0, 9.5 Hz, H-9 β)	32.4	2.08 (1H, m, H-9 α) 1.39 (1H, m, H-9 β)
10	193.7	9.41 (1H, s)	193.9	9.38 (1H, s)
OMe	57.4	3.37 (3H, s)	-	-



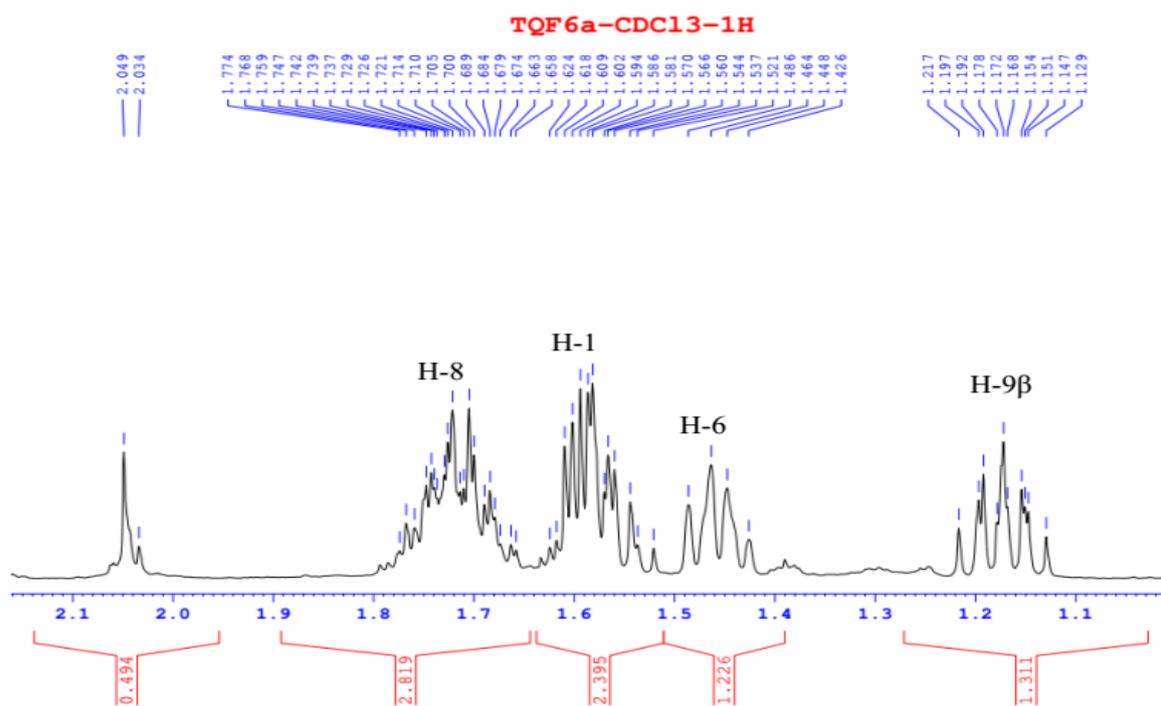


Figure S3: The extension (2) of ^1H -NMR spectrum of the compound (**1**)

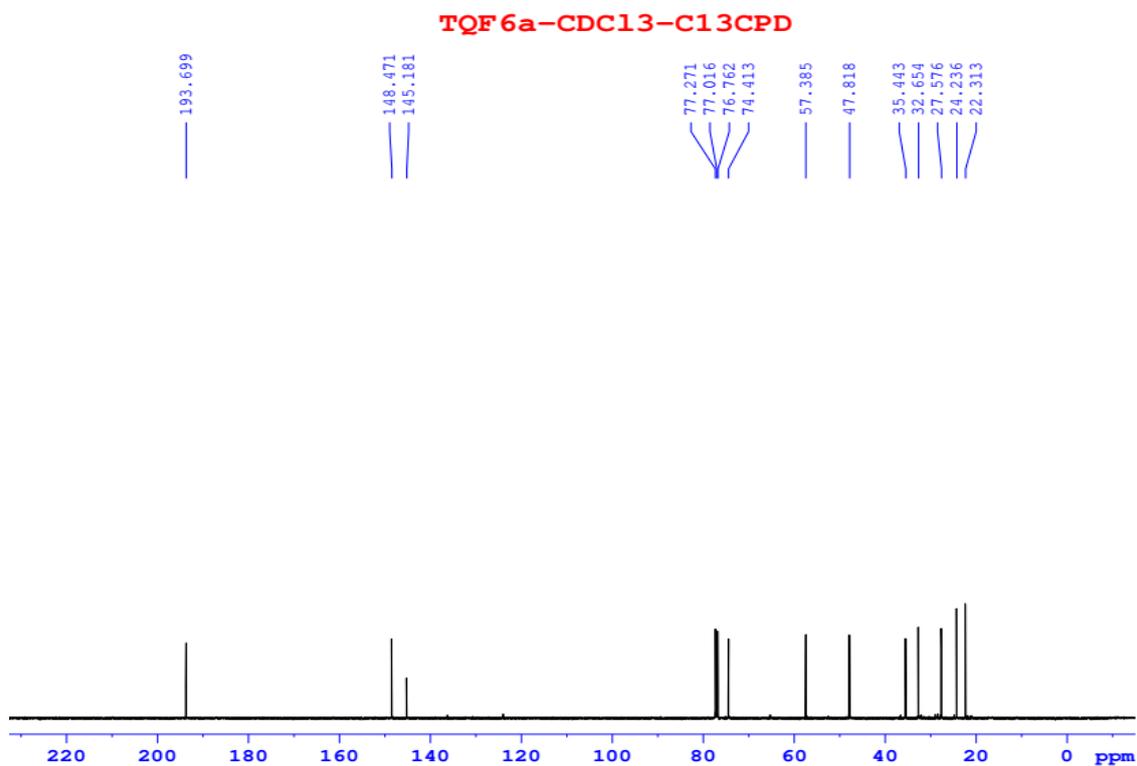


Figure S4: ^{13}C -NMR spectrum of the compound (**1**)

TQF6A-CDC13-HSQC

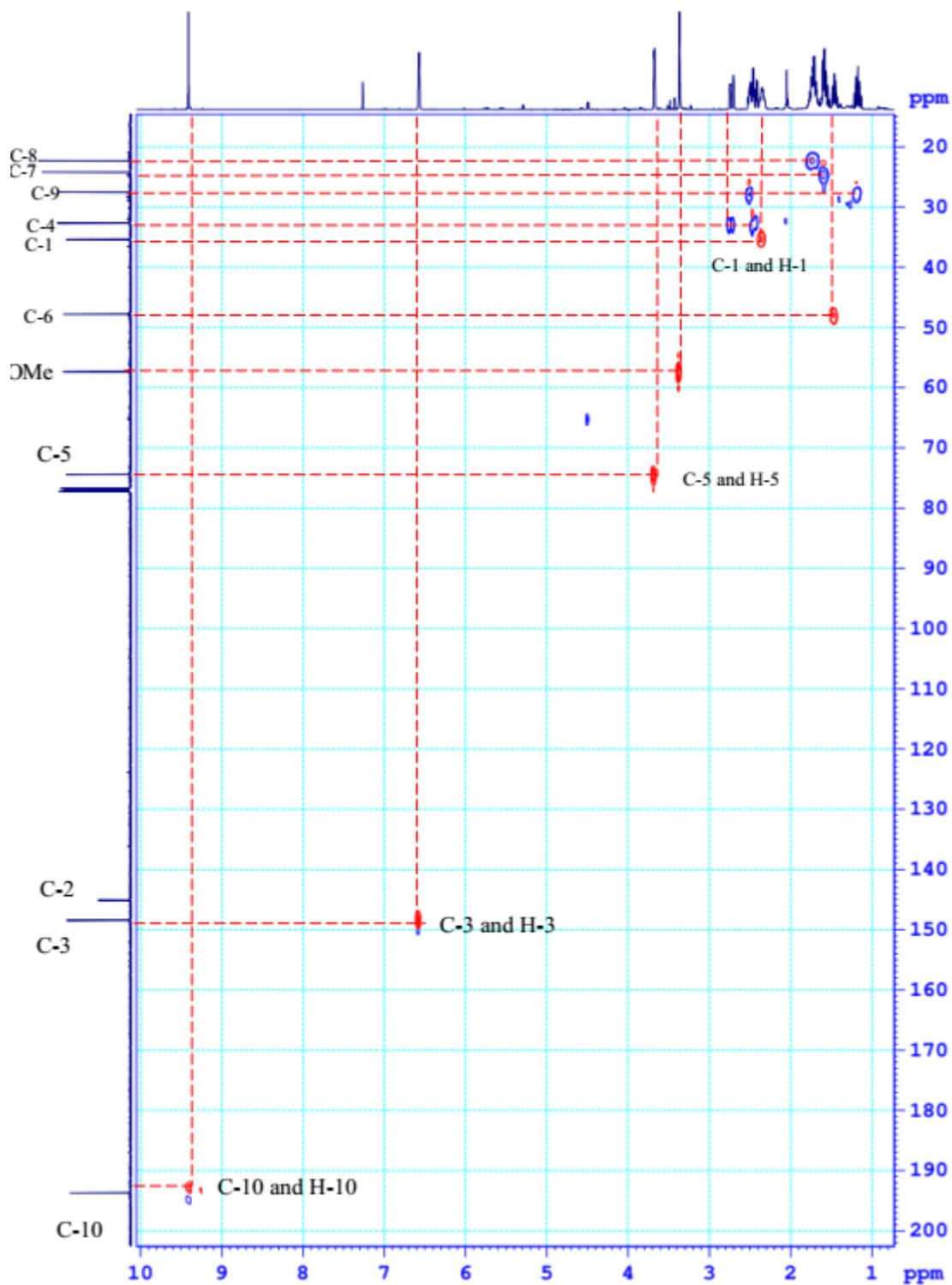


Figure S5: HSQC spectrum of the compound (1)

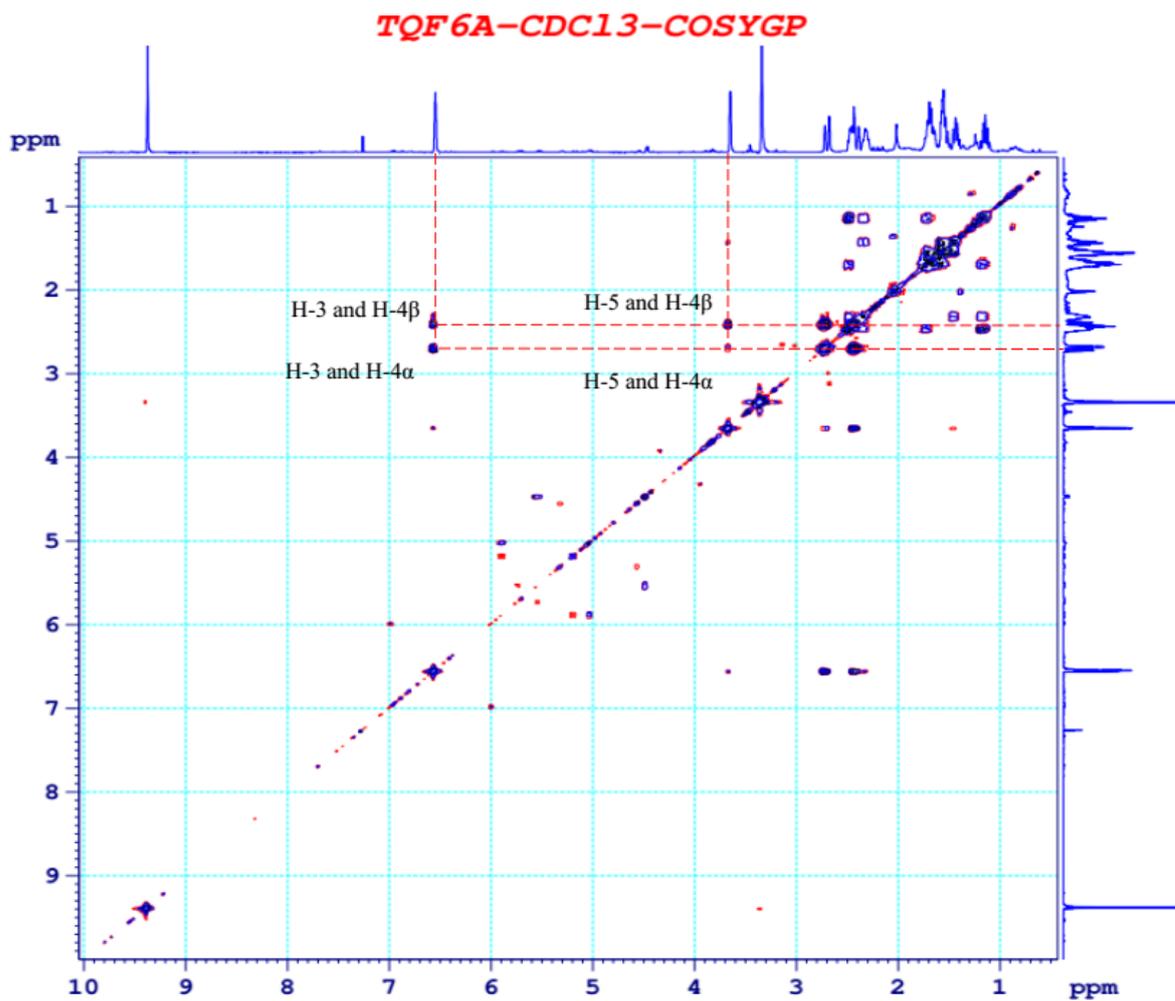


Figure S6: COSY spectrum of the compound (1)

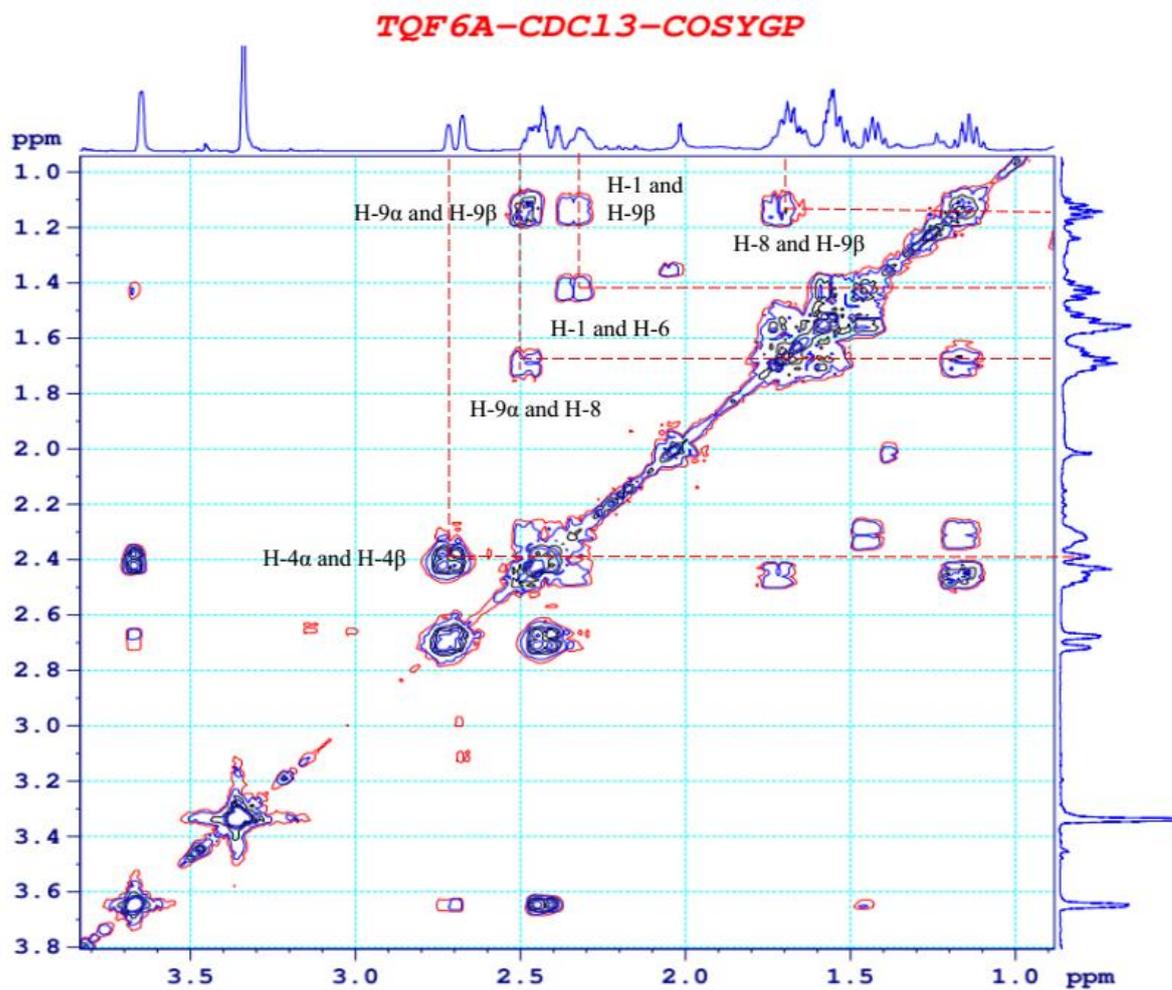
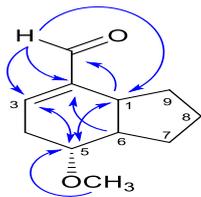


Figure S7: The extended ^1H - ^1H COSY spectrum of the compound (**1**)



TQF 6A-CDC13-HMBC

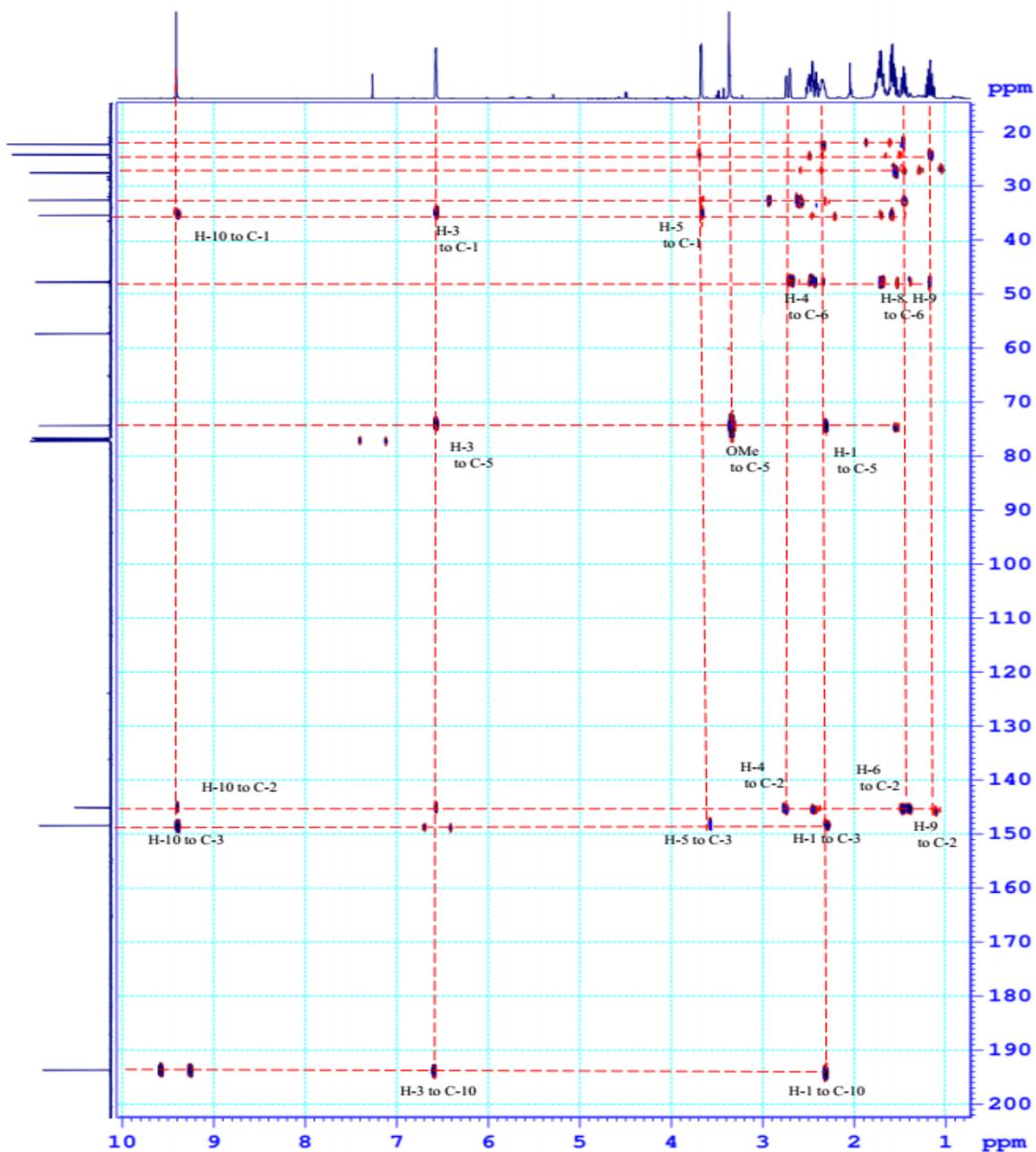


Figure S8: HMBC spectrum of the compound (1)

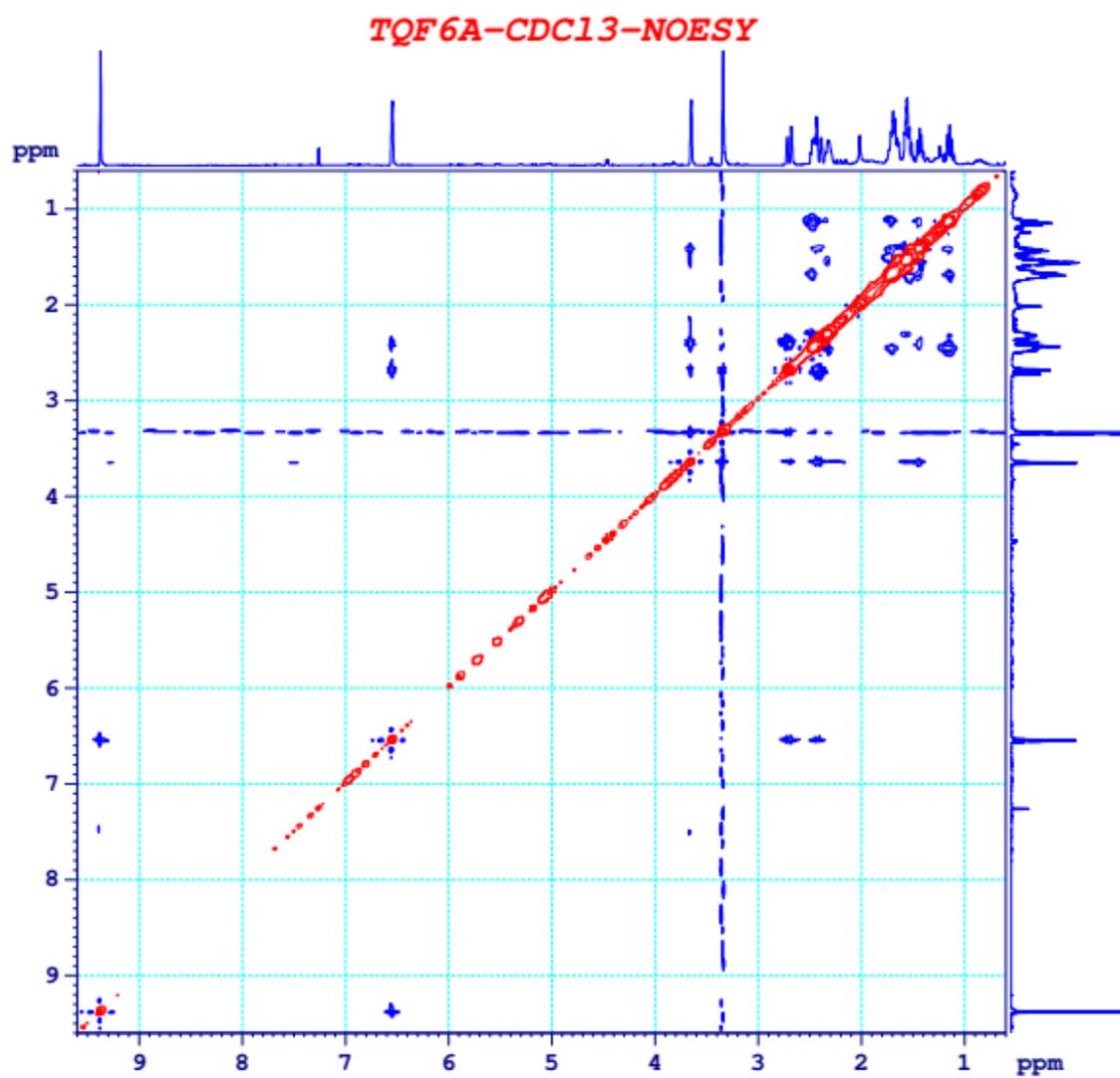


Figure S9: NOESY spectrum of the compound (1)

TQF6A-CDC13-NOESY

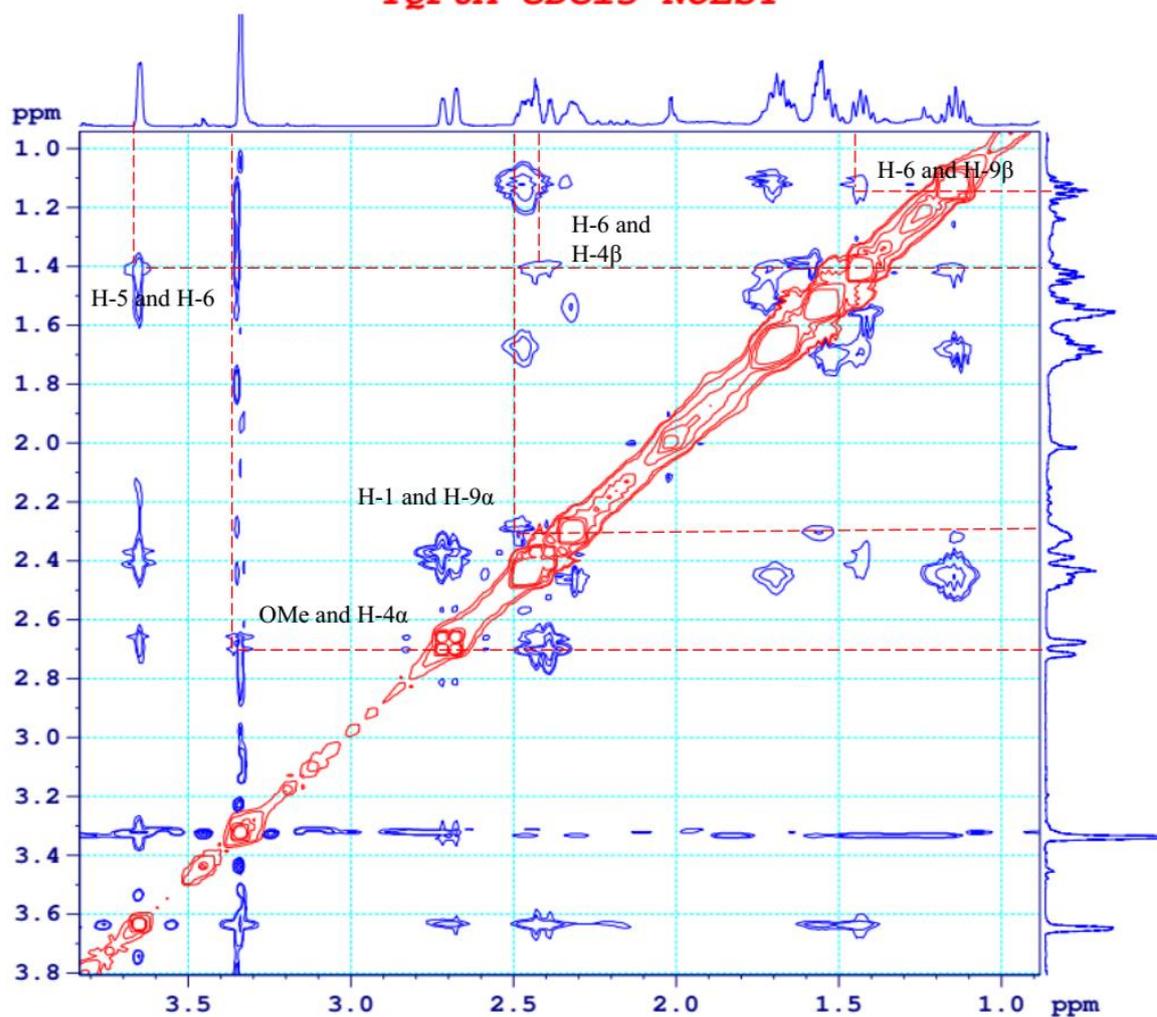


Figure S10: The extended NOESY spectrum of the compound (1)

TQF_170228164229 #955 RT: 3.25 AV: 1 NL: 6.11E6
T: FTMS + c ESI Full ms [50.00-500.00]

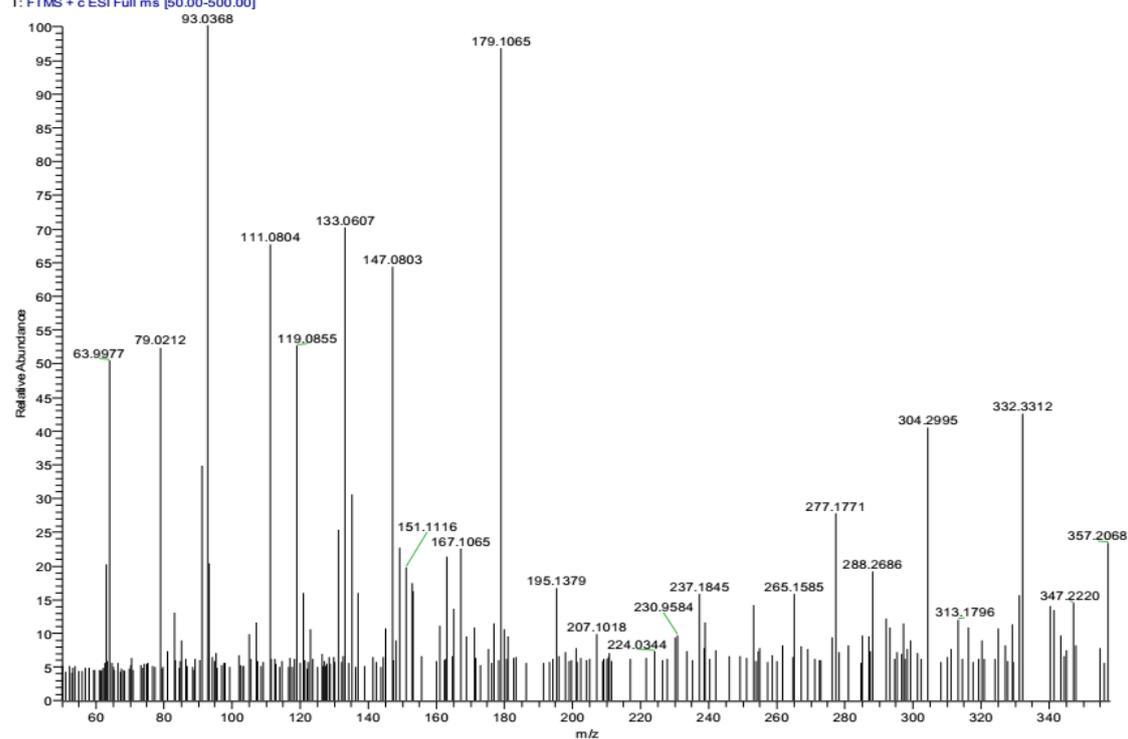


Figure S11: HRESIMS spectrum of the compound **1**

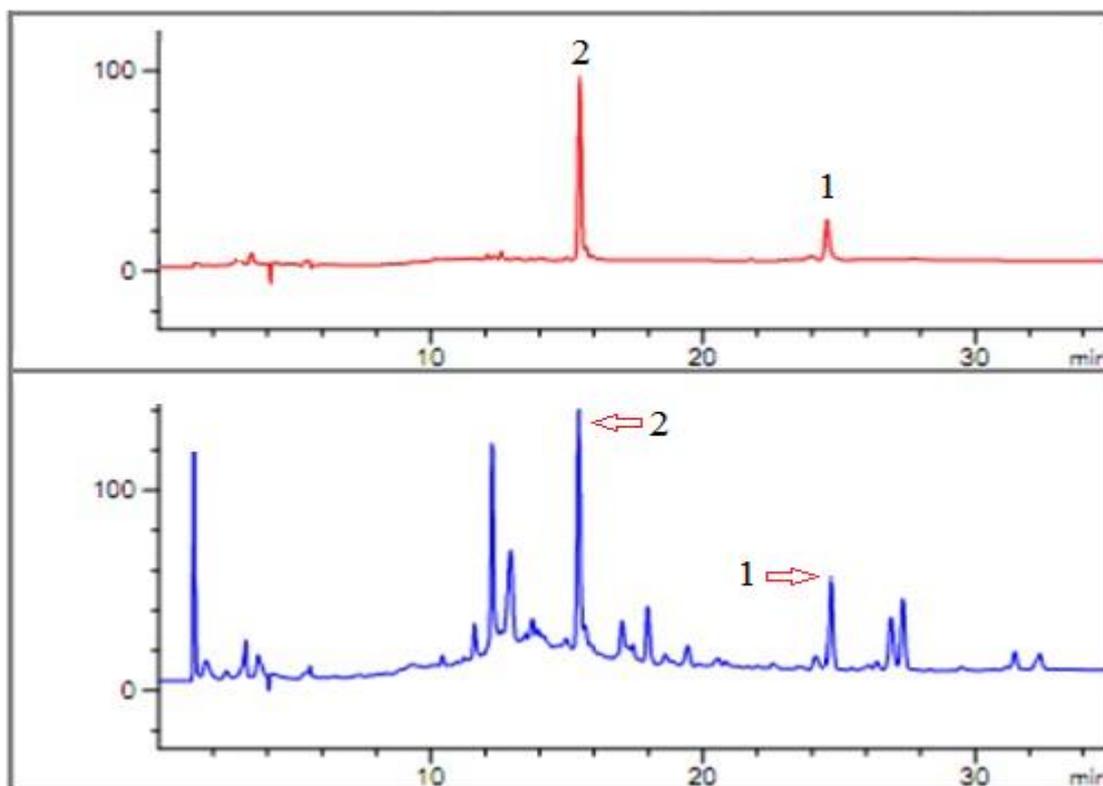


Figure S12: HPLC chromatograms of reference compounds **1** and **2** (upper) and *Amomum aromaticum* ethanol extract (lower).

The dried and powdered fruits (1 g) were extracted with 5 mL ethanol by sonication for 5 minutes. The extract was filtered and injected to an Agilent 1260 series HPLC-DAD system. A ZORBAX Eclipse XDB C18 column (150 mm x 4.6 mm, 5 μ m) was used. The elution was done with a gradient of 10-100% acetonitrile in water for 30 min. The detection wavelength was set at 230 nm for **1**.