SUPPORTING INFORMATION

Antimicrobial Sesquiterpenes from the Cultured Mycobiont Diorygma pruinosum against Methicillin-Resistant Staphylococcus aureus Isolated from Vietnamese Street Foods

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	2		4		Hydroxypruinosone	
No	$O_{4} = H_{15} = H_$		$\begin{array}{c} 2 \\ 0 \\ 3 \\ 4 \\ 15 \\ 15 \\ 12 \\ 12 \\ 12 \\ 14 \\ 9,10H \\ 9,10H \\ 13 \\ 13 \\ 12 \\ 13 \\ 12 \\ 13 \\ 12 \\ 14 \\ 9,10H \\ 13 \\ 13 \\ 12 \\ 13 \\ 13 \\ 12 \\ 13 \\ 13$		$\begin{array}{c} H \\ O \\ 3 \\ 4 \\ 5 \\ 15 \\ HO \\ 12 \end{array} \begin{array}{c} H \\ O \\ 8 \\ 7 \\ 11 \\ 12 \end{array} \begin{array}{c} H \\ O \\ 8 \\ 7 \\ 11 \\ 13 \end{array}$	
	δ н	δc	бн	δc	$\delta_{ m H}$	δc
1	3.04 (d, 9.0)	50.3	3.93 (m)	37.0	3.38 (d, 6.5)	50.5
2	2.51 (m) 2.29 (m)	45.3	2.46 (m) 2.08 (m)	39.3	2.26 (m) 2.46 (d, 10.0)	43.4
3		206.4		208.3	-	208.7
4		141.3		135.7	-	139.1
5		165.7		169.3	-	168.2
6	4.20 (s)	57.0	4.95 (d, 8.5)	68.9	5.02 (s)	66.6
7		70.7		151.6	-	64.9
8	1.65 (m) 1.60 (m)	23.8	5.79 (d, 7.5)	128.4	3.14 (t, 6.5)	57.3
9	2.26 (m) 2.22 (m)	38.6	4.17 (dd, 12.5, 6.0)	70.8	2.26 (m) 2.32 (dd, 7.5, 12.0)	37.7
10		72.1	2.40 (dd, 13.5, 6.0)	37.3	-	73.8
11		70.5	2.32 (hept, 7.0)	41.3	1.88 (hep, 6.5)	30.7
12	1.30 (s)	26.1	1.09 (d, 7.0)	21.5	0.80 (d, 6.5)	17.3
13	1.23 (s)	26.0	1.08 (d, 6.5)	22.3	1.04 (d, 7.0)	17.7
14	0.78 (s)	19.2	0.56 (d, 7.0)	10.5	0.94 (s)	19.9
15	1.76 (d, 1.5)	8.0	1.67 (d, 1.5)	8.2	1.76 (d, 1.5)	9.1
6-OH			5.38, (d, 8.5)		3.63 (brs)	
9-OH			5.19, (d. 5.5)			
10-OH					4.06 (brs)	

Table S1. ¹H (500 MHz) and ¹³C (125 MHz) NMR data of **2**, **4**, and hydroxypruinosone (acetone- d_6 , δ , ppm, J/Hz).

	3		Pruinosone	Pruinosone		
No	$O_{4} = \frac{2}{5} + \frac{14}{5} + \frac{14}{5} + \frac{10}{65} + \frac{8}{5} + \frac{14}{10} + \frac{10}{10} + \frac{8}{10} + \frac{10}{12} + 1$	ł	$O_{4} = \frac{14}{5} + \frac{14}{10} + \frac{14}{10}$			
	бн	δc	бн	δc		
1	3.05 (m)	43.9	3.28 (m)	43.8		
2	2.42 (dd, 18.5, 7.0) 2.03 (m)	40.8	2.08 (m) 2.32 (m)	38.1		
3	2.00 (11)	206.4	2102 (111)	208.1		
4		138.7		137.6		
5		165.6		170.8		
6	3.76 (s)	54.4	5.04 (d, 4.0)	66.8		
7		73.2		66.4		
8	4.23 (ddd, 11.5, 7.0, 3.5)	66.4	3.13 (t, 6.0)	61.4		
9	2.03 (m) 1.88 (m)	40.4	2.08 (m) 1.90 (m)	32.7		
10	2.08 (m)	32.5	2.32 (m)	33.2		
11	2.80 (m)	29.1	1.87 (hep, 6.5)	30.7		
12	0.86 (d, 7.0)	16.2	0.89 (d, 7.0)	17.5		
13	1.10 (d, 7.0)	20.0	1.04 (d, 7.0)	17.7		
14	0.57 (d, 7.0)	11.3	0.81 (d, 7.5)	14.6		
15	1.78 (d, 1.5)	7.7	1.74 (d, 1.5)	8.8		
6-OH			3.53 (d, 4.0)			

Table S2. ¹H (500 MHz) and ¹³C (125 MHz) NMR data of **3** and pruinosone (acetone- d_6 , δ , ppm, J/Hz).



Figure S1. S. aureus on mannitol salt agar (A) and Baird-Parker agar with tellurite egg yolk (B)



Figure S2. Gram stain of S. aureus



Figure S3. Catalase test result A: negative control B: a presumptive *S. aureus* isolate (catalase positive)



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Figure S12. ¹H NMR (acetone-*d*₆, 500 MHz) spectrum of 2



Figure S13. ¹³C NMR (acetone-*d*₆, 125 MHz) spectrum of 2



Figure S14. HSQC (acetone-d₆, 500 MHz, 125 MHz) spectrum of 2



Figure S15. HMBC (acetone- d_6 , 500 MHz, 125 MHz) spectrum of 2



Figure S16. COSY (acetone-d₆, 500 MHz) spectrum of 2



Figure S17. NOESY (acetone-d₆, 500 MHz, 125 MHz) spectrum of 2



Figure S18. HRESIMS spectrum of 2



Figure S19. HRESIMS spectrum of 3







Figure S21. ¹³C NMR (acetone- d_6 , 125 MHz) spectrum of **3**



Figure S22. ¹H–¹H COSY (acetone-*d*₆, 500 MHz) spectrum of **3**



Figure S23. HSQC (acetone-*d*₆, 500 MHz, 125 MHz) spectrum of 3



Figure S24. HMBC (acetone-*d*₆, 500 MHz, 125 MHz) spectrum of 3



Figure S25. NOESY (acetone-*d*₆, 500 MHz) spectrum of 3



Figure S27. ¹H NMR (acetone-*d*₆, 500 MHz) spectrum of 4



Figure S28. ¹³C NMR (acetone-*d*₆, 125 MHz) spectrum of 4



Figure S29. HSQC (acetone-*d*₆, 500 MHz, 125 MHz) spectrum of 4



Figure S30. HMBC (acetone-d₆, 500 MHz, 125 MHz) spectrum of 4



Figure S31. NOESY (acetone-d₆, 500 MHz) spectrum of 4

High-performance liquid chromatography analysis (HPLC) was conducted using an Agilent 1260 Infinity II system with a Diode Array Detector (DAD). For each prepared sample, 35 μ L (at a concentration of 1 mg/mL) was separately injected. A gradient system of acetonitrile (ACN) and water was used, with the following changes over a 60-minute analysis: 5% to 10% ACN in 5 minutes, 10% to 30% ACN in 15 minutes, 30% to 80% ACN in 10 minutes, 80% to 100% ACN in 5 minutes, followed by 100% ACN for 5 minutes. This analysis employed a Luna C18 column (Phenomenex, 150 mm × 4.6 mm, 5 μ m) and a C18 guard column (Phenomenex, Torrance, CA, USA)



Figure S34. Chromatogram of 4



Figure S35. ¹H NMR (acetone-*d*₆, 500 MHz) spectrum of **5**



Figure S36. ¹H NMR (CDCl₃, 500 MHz) spectrum of 6



Figure S37. ¹H NMR (DMSO-*d*₆, 500 MHz) spectrum of 7



Figure S38. ¹H NMR (DMSO-*d*₆, 500 MHz) spectrum of 8

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