

## Supporting Information

### **Ultrasound-assisted extraction and dispersive liquid-liquid microextraction coupled with gas chromatography-mass spectrometry for the sensitive determination of essential oil components in lavender**

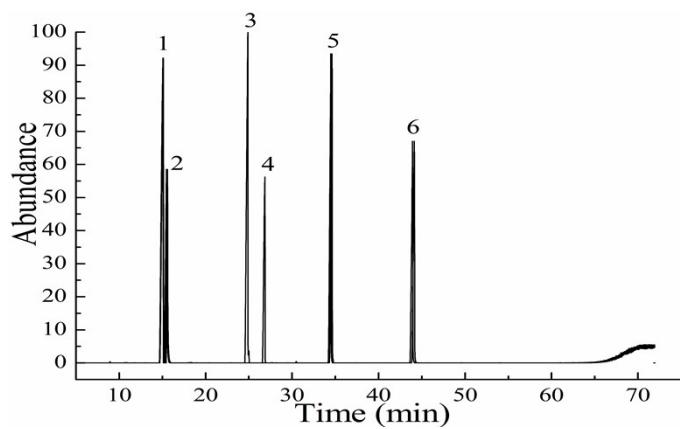
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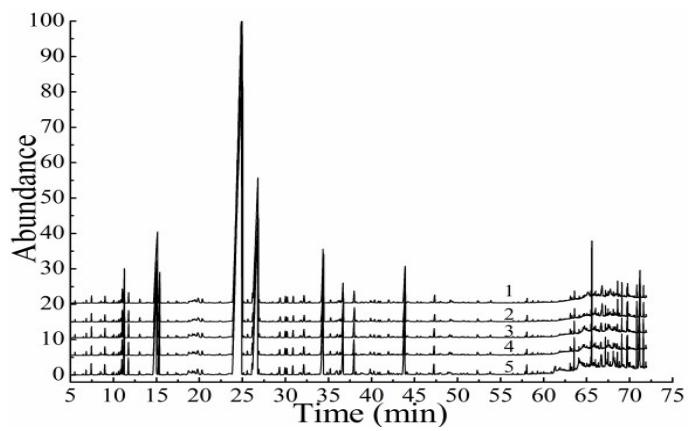
**Table S1** Quantitation results of six main compounds in lavender under optimized UAE-DLLME conditions.

Compounds	$t_R^a$	Regression equation	$R^2$	Linear range μg	LOD <sup>b</sup> ng	Recovery %	Content mg/g	m/z <sup>c</sup>
linalool	15.31, 15.35, 15.37, 15.30, 15.39	$y = 0.53 \times 10^6 x + 3.93 \times 10^6$	0.9991	8.80-400	1.75	97.62	2.61±0.12	71, 93, 55, 43, 41
1-octen-3-yl-acetate	15.50, 15.55, 15.59, 15.41, 15.42	$y = 0.34 \times 10^6 x + 1.09 \times 10^6$	0.9982	0.02-173.40	4.41	90.32	0.46±0.01	43, 99, 54, 41, 67
linalyl acetate	24.93, 24.92, 24.98, 24.95, 24.95	$y = 0.30 \times 10^6 x + 4.32 \times 10^6$	0.9995	8.80-400	2.94	94.38	23.27±1.02	93, 43, 80, 41, 121
lavandulyl acetate	26.72, 26.72, 26.71, 26.72, 26.70	$y = 0.32 \times 10^6 x + 1.05 \times 10^6$	0.9986	4.40-200	3.41	103.84	4.72±0.09	69, 93, 43, 41, 68
caryophyllene	34.67, 34.45, 34.68, 34.47, 34.76	$y = 0.39 \times 10^6 x + 1.26 \times 10^6$	0.9994	2.20-200	7.50	108.38	1.43±0.32	93, 133, 91, 41, 79
caryophyllene oxide	44.14, 44.17, 44.16, 44.22, 43.92	$y = 0.36 \times 10^6 x + 0.63 \times 10^6$	0.9991	1.20-200	3.00	87.91	1.51±0.06	43, 41, 79, 93, 91

<sup>a</sup>Retention time of five replicate experiments.<sup>b</sup>LOD, limit of detection, for a S/N = 3.<sup>c</sup>Five largest peaks in mass spectra.



**Fig.S1** The total ion chromatograms of the standard mixture. Peak identity: 1, linalool; 2, 1-octene-3-yl-acetate; 3, linalyl acetate; 4, lavandulyl acetate; 5, caryophyllene; and 6, caryophyllene oxide.



**Fig.S2** GC-MS total ion chromatograms of five replicate experiments.