

Table S4. Chemical composition of the VCs obtained by microwave extraction from the aerial parts of *Veronica* taxa collected on wet habitats.

Table S1. Chemical composition of the VOs obtained by microwave extraction from the aerial parts of <i>V. officinalis</i> L. collected in wet habitats.							
	<i>V. anagallis-aquatica</i>			<i>V. anagalloides</i>	<i>V. beccabunga</i>	<i>V. catenata</i>	<i>V. longifolia</i>
Component	RI ^a	RI ^b	VC±SD	VC±SD	VC±SD	VC±SD	VC±SD
Oxygenated monoterpenes			12.52	4.82	30.1	-	-
1,8-Cineole	1026	1210	0.46±0.01	-	-	-	-
γ-Terpinene	1057	1225	7.58±0.01	3..81±0.01	-	-	-
Linalool	1095	1506	0.71±0.01	-	-	-	-
Camphor	1151	1499	0.91±0.01	-	-	-	-
Terpinen-4-ol	1174	1686	-	1.01±0.04	-	-	-
<i>trans-p</i> -Mentha-1(7),8-dien-2-ol	1187	1803	1.78±0.01	-	0.82±0.01	-	-
Piperitone	1250	1719	-	-	29.28±0.01	-	-
Menthyl acetate	1294	1550	1.08±0.01	-	-	-	-
Sesquiterpene hydrocarbons			5.1	10.16	3.69	2.67	2.98
<i>E</i> -Caryophyllene*	1424	1585	3.29±0.01	4.01±0.01	2.95 ±0.04	2.48±0.01	1.43±0.01
<i>allo</i> -Aromadendrene	1465	1662	-	2.11±0.01	0.32±0.02	0.19±0.01	1.55±0.01
Germacrene D	1481	1692	0.88±0.01	3.07±0.01	0.42±0.01	-	-
δ-Selinene	1492	1756	0.22±0.04	0.97±0.01	-	-	-
δ-Cadinene	1517	1745	0.71±0.01	-	-	-	-
Oxygenated sesquiterpenes			29.45	31.19	11.18	24.39	12.37
Spathulenol	1577	2101	0.65±0.01	-	-	0.65±0.1	0.81±0.01
Caryophyllene oxide*	1581	1955	2.55±0.01	8.58±0.01	1.62±0.01	6.52±0.01	1.53±0.01
Viridiflorol	1592	2099	-	0.81±0.01	-	-	0.75±0.01
α-Bisabolol	1685	2210	0.28±0.01	1.91±0.01	-	-	-

α -Bisabolol oxide	1748	2511	-	0.77±0.02	-	-	-
Hexahydrofarnesyl acetone*	1839	2113	25.97±0.01	19.12±0.01	9.56±0.01	17.22±0.01	9.28±0.01
Oxygenated diterpene			14.56	14.88	34.54	42.26	37.18
Phytol*	1942	2610	14.56±0.01	14.88±0.01	34.54±0.01	42.26±0.01	37.18±0.01
Phenolic compounds			3.43	0.33	-	-	-
Methyl eugenol	1403	2005	3.43±0.01	0.33±0.07	-	-	-
Acids, alcohols and esters			25.93	18.17	9.54	12.98	30.01
Isopentyl acetate	863	1127	-	-	-	-	0.22±0.03
Benzaldehyde	952	1508	1.63±0.01	-	-	-	-
Benzene acetaldehyde	1036	1633	9.67±0.01	-	2.48±0.01	-	4.17±0.01
<i>n</i> -Nonanal	1100	1389	1.82±0.01	2.12±0.01	0.89±0.02	-	2.68±0.01
Hexyl 2-methyl butanoate	1233	1425	-	0.44±0.01	-	-	-
<i>n</i> -Decanol	1266	1711	1.28±0.01	-	-	-	0.58±0.01
2,4-Decadienal	1304	1764	0.28±0.05	-	-	-	-
(<i>E</i>)- β -Damascenone	1384	1819	0.12±0.16	0.52±0.07	-	0.32±0.09	-
β -Ionone	1487	1935	6.36±0.01	4.22±0.01	1.43±0.01	0.27±0.01	16.22±0.01
Hexadecanoic acid*	1959	2912	4.77±0.01	9.17±0.01	4.74±0.01	5.81±0.01	6.14±0.01
Oleic acid	2133	2998	-	1.02±0.01	-	2.25±0.01	-
Octadecanol acetate	2209	-	-	0.68±0.03	-	4.33±0.01	-
Hydrocarbons			5.3	18.12	8.26	15.1	10.71
Eicosane*	2000	2000	0.72±0.01	1.12±0.01	-	-	-
Heneicosane*	2100	2100	0.18±0.01	0.33±0.04	-	-	-
Docosane*	2200	2200	2.22±0.01	5.62±0.01	4.15±0.01	-	2.82±0.01

Tricosane*	2300	2300	0.75±0.01	-	-	3.03±0.01	-
Tetracosane*	2400	2400	1.15±0.01	-	-	12.07±0.01	-
Pentacosane*	2500	2500	-	5.43±0.01	1.13±0.01	-	4.71±0.01
Hexacosane*	2600	2600	-	3.90±0.01	0.27±0.09	-	1.48±0.01
Heptacosane*	2700	2700	0.28±0.1	0.95±0.01	1.45±0.01	-	1.70±0.01
Octacosane*	2800	2800	-	0.77±0.15	1.26±0.01	-	-
Total identification (%)			97.29	96.79	97.31	97.4	93.25

Retention indices (RIs) were determined relative to a series of n-alkanes (C8–C40) on capillary columns VF5-ms (RI^a) and CPWax 52 (RI^b); Identification method: RI, comparison of RIs with those in a self-generated library reported in the literature [41] and/or with authentic samples; comparison of mass spectra with those in the NIST02 and Wiley 9 mass spectral libraries; *co-injection with reference compounds; -, not identified; SD, standard deviation of triplicate analysis.