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Introduction

Neurodegenerative diseases are an internationally faced public health problem. Acetylcholinesterase (AChE) catalyzes the hydrolysis of the neurotransmitter acetylcholine, which plays a key role in memory and cognition.¹ It is also the key enzyme target for treatments of myasthenia gravis, glaucoma, Alzheimer's disease, Parkinson's disease, and other neurodegenerative diseases.² Although a few cholinesterase inhibitors from synthetic and natural origins are available in the drug market, they are used only for symptomatic treatment. Moreover, side effects and relatively low bioavailability limit their clinical applications.3 There is still a great demand to discover new cholinesterase inhibitors, which can improve cholinergic

Structurally diverse biflavonoids from Dysosma versipellis and their bioactivity†

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Five pairs of new biflavonoid enantiomers, (\pm) -dysosmabiflavonoids A-E (1-5), two new biflavonoids, dysosmabiflavonoids F-G (6–7), and four biosynthetically related precursors (8–11) were isolated from the roots and rhizomes of Dysosma versigellis. Their structures were elucidated by extensive spectroscopic analysis, including HR-ESI-MS and 2D NMR. Their absolute configurations were determined by comparison of the calculated and experimental ECD spectra. All isolated compounds were evaluated for AChE inhibitory activity. Compounds 6 and 7 exhibited more potent inhibitory activities with IC₅₀ values of 1.42 and 0.73 μ M, respectively, than their biosynthetically related precursors kaempferol (8, 17.90 μ M) and quercetin (9, 3.96 μ M). The preliminary structure-activity relationship study indicated that the connection mode of biflavonoid subunits, oxidation degree of the C ring, and 3,4dihydroxy group of the B ring were important structural factors for AChE inhibitory activity. Racemates 1-5 and their corresponding levorotatory and dextrorotatory enantiomers were tested for their potential to impede the generation of NO in lipopolysaccharide-stimulated RAW264.7 cells, and their mushroom tyrosinase inhibitory effect. Racemate 1 displayed more potent mushroom tyrosinase inhibitory activity (IC₅₀, 28.27 µM) than the positive control kojic acid (IC₅₀, 32.59 µM). D. versipellis may have therapeutic potential for melanogenesis disorders and neurodegenerative diseases.

> transmission in neurodegenerative diseases. Traditional Chinese medicine has become a promising source of structurally novel cholinesterase inhibitors.

> A perennial herb of the Berberidaceae family, Dysosma versipellis (Hance) M. Cheng ex Ying, is widely distributed in the central/south regions of China.4 Its roots and rhizomes, called "Bajiaolian" in Chinese, have the functions of eliminating phlegm, detoxification, removing blood stasis and dispersing knots.⁵ As a common folk medicine, it has a satisfactory effect on the treatment of parotitis,⁴ cough, sore throat, scrofula, nameless swollen venom/carbuncles, snake bite, fall injury, rheumatoid arthritis,5 epidemic encephalitis B, parasites, tumor,6 lumbago, skelalgia,7 ulcer,8 weakness, condyloma accuminata, lymphadenopathy,9 pyogenic infection,10 hemiplegia, joint pain,¹¹ and syphilis¹² since the Han dynasty in China. Its previous phytochemical and pharmacological investigations revealed the presense of bioactive biflavonoids,13-16 and their neuraminidase inhibitory¹³ and cytotoxic properties.¹⁴ In our search for bioactive natural products, five pairs of new biflavonoid enantiomers, (\pm) -dysosmabiflavonoids A-E (1-5), two new biflavonoids, dysosmabiflavonoids F-G (6-7), along with four biosynthetically related precursors (8-11) (Fig. 1) were isolated from the roots and rhizomes of D. versipellis. Reported herein are their isolation, structure elucidation, and bioactive evaluation.

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[†] Electronic supplementary information (ESI) available: UV, IR, HR-ESI-MS, ECD, 1D and 2D NMR spectra for compounds 1-7. See DOI: https://doi.org/10.1039/d2ra06961j



Results and discussion

Compound 1 was obtained as a yellow amorphous powder and possessed a molecular formula C₃₀H₁₈O₁₃ with twenty-two degrees of unsaturation, as revealed from its HR-ESI-MS analysis $(m/z 587.0806 [M + H]^+$, calcd for C₃₀H₁₉O₁₃, 587.0826). The IR spectrum displayed two carbonyl groups (1726, 1635 cm⁻¹), hydroxy groups (3374 cm^{-1}), aromatic rings ($1607, 1512 \text{ cm}^{-1}$) and tertiary alcohol (1179 cm⁻¹). The UV spectrum exhibited absorption maxima characteristic for flavonol at 275 and 360 nm. The ¹³C NMR (Table 1) and HSQC spectra revealed one biflavone skeleton including two carbonyl groups $\delta_{\rm C}$ 176.6, 190.1 (characteristic of a flavonol and a flavanonol, respectively), four benzene rings, two oxygen-bearing olefinic carbons $\delta_{\rm C}$ 147.8, 136.1, two aliphatic quaternary carbons $\delta_{\rm C}$ 117.6, 79.1. The ¹H NMR spectrum (Table 2) showed four sets of aromatic systems including one *para*-disubstituted benzene ring $\delta_{\rm H}$ 8.08 (2H, d, J = 8.5 Hz), 6.94 (2H, d, J = 8.5 Hz), one pentasubstituted benzene ring $\delta_{\rm H}$ 7.01 (1H, s), one 1,3,4-trisubstituted benzene ring $\delta_{\rm H}$ 6.89 (1H, s), 6.72 (1H, d, J = 8.6Hz), 6.75 (1H, d, J = 8.6 Hz), and one 1,2,3,5-tetra-substituted benzene ring $\delta_{\rm H}$ 5.86 (1H, s) and 5.78 (1H, s). The HMBC correlations (Fig. 2) of the aromatic protons $\delta_{\rm H}$ 8.08 (2H, d, J =8.5 Hz, H-2^{'''}, 6^{'''}) with C-2^{''} ($\delta_{\rm C}$ 147.8), in combination with one penta-substituted benzene ring at $\delta_{\rm H}$ 7.01 (1H, s), indicated that compound 1 contained kaempferol as a subunit.17 Another subunit was identified as dihydroquercetin by the HMBC cross

peak of the aromatic protons $\delta_{\rm H}$ 6.89 (1H, s, H-2') and 6.75 (1H, d, J = 8.6 Hz, H-6') with C-2 ($\delta_{\rm C}$ 117.6), one 1,2,3,5-tetrasubstituted benzene ring $\delta_{\rm H}$ 5.86 (1H, s), 5.78 (1H, s), and two aliphatic quaternary carbons $\delta_{\rm C}$ 117.6, 79.1. A careful comparison of the ¹³C NMR spectra of 1 with kaempferol indicated that the subunit of kaempferol was substituted at C-6 and C-7, which was confirmed by the chemical shift change from $\delta_{\rm C}$ 156.8 (C-5' '), 110.3 (C-6''), 164.3 (C-7''), 91.1 (C-8''), and $\delta_{\rm H}$ 7.01 (1H, s, H-8'') in **1** to $\delta_{\rm C}$ 160.7 (C-5), 98.2 (C-6), 163.9 (C-7), 93.5 (C-8), and $\delta_{\rm H}$ 6.48 (1H, d, J = 2.0 Hz, H-8) in kaempferol.¹⁷ An ether bridge C-2-O-C-7" and a C-3-C-6" bond between the two flavonoid subunits were determined by two aliphatic quaternary carbons [one di-oxygenated $\delta_{\rm C}$ 117.6 (C-2), one mono-oxygenated $\delta_{\rm C}$ 79.1 (C-3)].¹⁸ The planar structure of 1 (Fig. 1) was determined by detailed elucidation on 1D and 2D NMR data. Despite repeated experiments, suitable single crystals of 1 for X-ray diffraction were not obtained successfully. NOESY experiments failed to provide any correlations between the two flavonoid subunits. Compound 1 showed no distinct Cotton effects (200-400 nm) at all examined concentrations, indicating it was a racemic mixture. This result is also consistent with the observation that 1 was optically inactive. Racemate 1 was resolved into 1a and 1b in a ratio of approximately 1:1 by a chiral Daicel PAK AD-H column. As expected, the ECD spectra of 1a and 1b showed opposite Cotton effects. The absolute configurations of C-2 and C-3 were extrapolated by comparing the experimental with calculated ECD spectra, the latter performed by density

Table 1 ¹³C NMR data (125 MHz) of 1–7 in DMSO- d_c (δ in ppm)

No.	1	2	3	4	5	6	7
2	117.6 C	117.8 C	117.8 C	117.7 C	117.8 C	146.9 C	146.7 C
3	79.1 C	79.2 C	79.9 C	79.9 C	79.9 C	135.4 C	135.4 C
4	190.1 C	190.7 C	191.0 C	190.5 C	191.0 C	176.2 C	176.1 C
5	163.3 C	163.4 C	163.5 C	163.5 C	163.5 C	158.4 C	158.4 C
6	97.5 CH	96.9 CH	97.0 CH	97.1 CH	97.0 CH	97.9 CH	98.0 CH
7	168.9 C	167.8 C	167.9 C	168.1 C	167.9 C	162.1 C	162.1 C
8	95.4 CH	95.1 CH	95.3 CH	95.3 CH	95.3 CH	104.6 C	104.7 C
9	160.4 C	160.4 C	160.6 C	160.3 C	160.6 C	153.8 C	153.8 C
10	97.8 C	98.4 C	98.6 C	98.3 C	98.6 C	102.8 C	102.6 C
1'	123.9 C	123.7 C	123.7 C	123.4 C	123.7 C	122.1 C	122.5 C
2'	114.5 CH	128.8 CH	114.5 CH	128.3 CH	114.5 CH	129.6 CH	115.2 CH
3'	144.6 C	115.2 CH	144.7 C	114.8 CH	144.7 C	115.4 CH	145.0 C
4'	146.8 C	159.1 C	146.9 C	158.7 C	146.9 C	159.1 C	147.6 C
5'	114.9 CH	115.2 CH	115.5 CH	114.8 CH	115.1 CH	115.4 CH	115.6 CH
6'	118.1 CH	128.8 CH	118.2 CH	128.3 CH	118.2 CH	129.6 CH	120.0 CH
2″	147.8 C	148.0 C	147.5 C	147.5 C	147.5 C	146.9 C	146.6 C
3″	136.1 C	136.2 C	136.1 C	136.0 C	136.1 C	135.4 C	135.3 C
4″	176.6 C	176.6 C	176.3 C	176.2 C	176.2 C	176.2 C	176.1 C
5″	156.8 C	156.9 C	163.9 C	163.8 C	163.8 C	158.4 C	158.3 C
6″	110.3 C	110.0 C	94.8 CH	94.7 CH	94.7 CH	97.9 CH	97.9 CH
7″	164.3 C	164.3 C	164.9 C	164.8 C	164.8 C	162.1 C	162.1 C
8″	91.1 CH	91.1 CH	106.3 C	106.1 C	106.1 C	104.6 C	104.6 C
9″	157.2 C	157.2 C	151.2 C	151.5 C	151.2 C	153.8 C	153.7 C
10″	105.6 C	105.7 C	105.2 C	105.2 C	105.2 C	102.8 C	102.6 C
1‴	121.3 C	121.6 C	121.4 C	121.7 C	121.7 C	122.1 C	122.1 C
2‴	129.7 CH	115.3 CH	130.2 CH	116.6 CH	116.2 CH	129.6 CH	129.5 CH
3‴	115.5 CH	145.2 C	115.3 CH	145.4 C	144.9 C	115.4 CH	115.4 CH
4‴	159.6 C	148.2 C	159.6 C	148.1 C	148.1 C	159.1 C	159.0 C
5‴	115.5 CH	115.7 CH	115.3 CH	115.3 CH	115.3 CH	115.4 CH	115.4 CH
6‴	129.7 CH	120.2 CH	130.2 CH	120.5 CH	120.5 CH	129.6 CH	129.5 CH
8-CH ₂ -8"						16.4 CH_2	16.5 CH_2

functional theory. A comparison of the observed ECD spectra for 1a and 1b with the calculated ECD spectra for the (2S, 3S)-1 and (2R, 3R)-1 is shown in Fig. 3. The overall ECD spectra for (2S, 3S)-1 and (2R, 3R)-1 are in good accordance with the experimental ECD for 1a and 1b, respectively. Therefore, the absolute configurations of 1a and 1b were respectively deduced to be 2S, 3S and 2R, 3R. Thus, compounds 1a and 1b were determined as (2S,3S)-3,5,7,3',4'-pentahydroxyflavanone-(2-O-7":3-6")-3",5",4"'trihydroxyflavone and (2R,3R)-3,5,7,3',4'-pentahydroxyflavanone-(2-O-7":3-6")-3",5",4"'-trihydroxyflavone, and named as (+)-dysosmabiflavonoid A and (-)-dysosmabiflavonoid A, respectively.

Compound 2 was obtained as a yellow amorphous powder. It gave the same molecular formula C₃₀H₁₈O₁₃ as that of 1, based

Table 2	¹ H NMR data	(500 MHz) o	f 1–7 in	DMSO-d ₆	(δ in ppm,	J in Hz
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No.	1	2	3	4	5	6	7
6	5.86, s	5.93, d (1.6)	5.90, d (1.6)	5.88, d (1.9)	5.76, s	6.17, s	6.12, s
8	5.78, s	5.86, d (1.6)	5.86, d (1.6)	5.86, d (1.9)	5.73, s		
2'	6.89, s	7.28, d (8.7)	6.90, d (1.6)	7.29, d (8.8)	6.89, d (2.0)	7.97, d (8.7)	7.82, d (2.0)
3'		6.79, d (8.7)		6.79, d (8.8)		6.84, d (8.7)	
5'	6.72, d (8.6)	6.79, d (8.7)	6.74, d (8.4)	6.79, d (8.8)	6.73, d (8.3)	6.84, d (8.7)	6.80, d (8.5)
6′	6.75, d (8.6)	7.28, d (8.7)	6.76, dd (8.4, 1.6)	7.29, d (8.8)	6.76, dd (8.3, 2.0)	7.97, d (8.7)	7.54, dd (8.5, 2.0)
6″			6.71, s	6.69, s	6.67, s		6.16, s
8″	7.01, s	7.00, s					
2‴	8.08, d (8.5)	7.72, d (1.9)	8.16, d (8.8)	7.88, d (2.2)	7.89, d (2.0)	7.97, d (8.7)	8.01, d (8.8)
3‴	6.94, d (8.5)		6.94, d (8.8)			6.84, d (8.7)	6.83, d (9.0)
5‴	6.94, d (8.5)	6.91, d (8.5)	6.94, d (8.8)	6.90, d (8.5)	6.90, d (8.6)	6.84, d (8.7)	6.83, d (9.0)
6‴	8.08, d (8.5)	7.58, dd (8.5, 1.9)	8.16, d (8.8)	7.64, dd (8.5, 2.2)	7.65, dd (8.6, 2.0)	7.97, d (8.7)	8.01, d (8.8)
8-CH ₂ -8"						4.22, s	4.24, s
5-OH						12.53, s	12.52, s
5″-OH							12.53, s



on the positive HR-ESI-MS ion observed at m/z 587.0824 [M + H]⁺ (Calcd 587.0826). Its ¹³C and ¹H NMR spectra (Tables 1 and 2) were quite similar to those of 1, except that quercetin and dihydrokaemferol subunit in 2 were observed instead of kaempferol and dihydroquercetin subunit respectively in 1. Those were further supported by HMBC cross peak of the aromatic protons $\delta_{\rm H}$ 7.72 (1H, d, J = 1.9 Hz, H-2^{'''}) and 7.58 (1H, dd, J = 8.5, 1.9 Hz, H-6^{'''}) with C-2^{''} ($\delta_{\rm C}$ 148.0), of the aromatic proton $\delta_{\rm H}$ 7.28 (2H, d, J = 8.7 Hz, H-2′, 6′) with C-2 ($\delta_{\rm C}$ 117.8). The lack of optical activity and Cotton effect in the ECD spectrum indicated that 2 was a racemic mixture. Chiral HPLC resolution of 2 gave 2a and 2b in a ratio of approximately 1:1. Comparison between the experimental and calculated ECD spectra of two enantiomers allowed the absolute configurations of 2a and 2b to be established as (2S,3S) and (2R, 3R), respectively. Thus, compounds 2a and 2b were determined as (2S,3S)-3,5,7,4'-tetrahydroxyflavanone-(2-O-7":3-6")-3",5",3"",4"''-tetrahydroxyflavone and (2R,3R)-3,5,7,4'-tetrahydroxyflavanone-(2-O-7' ':3-6")-3",5",3",4"'-tetrahydroxyflavone, and named as (+)-dysosmabiflavonoid B and (-)-dysosmabiflavonoid B, respectively.

Compounds 3 and 4 were obtained as yellow, amorphous powders. They possessed the same molecular formula C₃₀H₁₈O₁₃ as that of 1 and 2, as revealed from their HR-ESI-MS analyses $(m/z 587.0808 [M + H]^+ \text{ in } 3; m/z 587.0800 [M + H]^+ \text{ in } 4)$. Their ¹³C and ¹H NMR data (Tables 1 and 2) were quite similar to those of 1 and 2, except that the chemical shift changes of an aromatic carbon and an aromatic proton [from $\delta_{\rm C}$ 94.8 (C-6"), $\delta_{\rm H}$ 6.71 (1H, s, H-6") in 3 to $\delta_{\rm C}$ 91.1 (C-8"), $\delta_{\rm H}$ 7.01 (1H, s, H-8") in 1, from $\delta_{\rm C}$ 94.7 (C-6"), $\delta_{\rm H}$ 6.69 (1H, s, H-8") in 4 to $\delta_{\rm C}$ 91.1 (C-8"), $\delta_{\rm H}$ 7.00 (1H, s, H-8") in 2, indicating the subunits of kaempferol in 3 and guercetin in 4 were substituted at C-7 and C-8. Those were also supported by chemical shift change from $\delta_{\rm C}$ 94.8 (C-6"), 164.9 (C-7"), and 106.3 (C-8") in 3 to $\delta_{\rm C}$ 98.2 (C-6), 163.9 (C-7), and 93.5 (C-8) in kaempferol [17], and from $\delta_{\rm C}$ 94.7 (C-6"), 164.8 (C-7"), and 106.1 (C-8") in 4 to $\delta_{\rm C}$ 98.6 (C-6), 164.3 (C-7), and 93.8 (C-8) in quercetin.¹⁹ Compounds 3 and 4 were also racemic mixtures, and separated into 3a and 3b, and 4a and 4b, respectively. The absolute configurations of 3a, 3b, 4a, 4b were determined as (2S, 3S), (2R, 3R), (2S, 3S), (2R, 3R) by comparison of their experimental and calculated ECD spectra. Thus,

compounds **3a**, **3b**, **4a** and **4b** were deduced as (2S,3S)-3,5,7,3',4'-pentahydroxyflavanone-(2-0-7'':3-8'')-3'',5'',4'''-trihydroxyflavone, (2R,3R)-3,5,7,3',4'-pentahydroxyflavanone-(2-0-7'':3-8'')-3'',5'',4'''-trihydroxyflavone, (2S,3S)-3,5,7,4'-tetrahydroxy yflavanone-(2-0-7'':3-8'')-3'',5'',3,4'''-tetrahydroxyflavone, and (2R,3R)-3,5,7,4'-tetrahydroxyflavanone-(2-0-7'':3-8'')-3'',5'',3''',4'''tetrahydroxyflavone, and named as (-)-dysosmabiflavonoid C, (+)-dysosmabiflavonoid D, respectively.

Compound 5 was obtained as a yellow amorphous powder. Its ¹H and ¹³C NMR spectra data (Tables 1 and 2) were quite analogous to those of 4, except that 1,3,4-trisubstituted benzene ring $\delta_{\rm H}$ 6.89 (1H, d, J = 2.0 Hz), 6.73 (1H, d, J = 8.3 Hz), 6.76 (1H, dd, J = 8.3, 2.0 Hz) was observed in 5 instead of 1,4-disubstituted benzene ring in 4. This was further supported by their HR-ESI-MS, which gave an $[M + H]^+$ quasi-molecular ion peak m/z603.0752 (Calcd 603.0775) in 5, being 16 mass-units more than that of 4. The dihydroquercetin subunit was identified as by the HMBC cross peak of the aromatic protons $\delta_{\rm H}$ 6.89 (1H, d, J =2.0 Hz, H-2') and 6.76 (1H, dd, J = 8.3, 2.0 Hz, H-6') with C-2 ($\delta_{\rm C}$ 117.8), one 1,2,3,5-tetra-substituted benzene ring $\delta_{\rm H}$ 5.76 (1H, s), 5.73 (1H, s), and two oxygenated aliphatic quaternary carbons $\delta_{\rm C}$ 117.8, 79.9. The apparent lack of a Cotton effect in ECD spectrum indicated that 5 was a racemic mixture. Compounds 5a and 5b were obtained as a pair of enantiomers by chiral-phase HPLC isolation of 5. The absolute configurations of 5a and 5b were determined as (2S, 3S) and (2R, 3R) by comparison of calculated ECD and experimental data. Thus, compound 5 was deduced as (2S,3S)-3,5,7,3',4'-pentahydroxyflavanone-(2-O-7":3-8")-3",5",3"',4"'-tetrahydroxyflavone and (2R,3R)-3,5,7,3',4'-pentahydroxyflavanone-(2-O-7":3-8")-3",5' ',3''',4'''-tetrahydroxyflavone, (–)-dysosmabiflavonoid E, and (+)-dysosmabiflavonoid E, respectively.

Compound **6** was obtained as a yellow amorphous powder and possessed a molecular formula $C_{31}H_{20}O_{12}$ with twenty-two degrees of unsaturation, as revealed from its HR-ESI-MS analysis (*m*/*z* 585.1023 [M + H]⁺, calcd for $C_{31}H_{21}O_{12}$, 585.1033). The IR spectrum displayed the conjugated carbonyl group (1647 cm⁻¹), hydroxy groups (3361 cm⁻¹), and aromatic ring (1510 cm⁻¹). The UV spectrum exhibited absorption maxima

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Fig. 3 Experimental and calculated ECD spectra of compounds 1a-5a and 1b-5b.

characteristic for flavonol at 271 and 369 nm. The ¹H NMR spectrum showed two sets of aromatic systems including one para-disubstituted benzene ring $\delta_{\rm H}$ 7.97 (2H, d, J = 8.7 Hz), 6.84 (2H, d, J = 8.7 Hz), and one penta-substituted benzene ring $\delta_{\rm H}$ 6.17 (1H, s), and one isolated aliphatic proton $\delta_{\rm H}$ 4.22 (1H, s). The ¹³C NMR and HSQC spectrum revealed one aliphatic carbon $\delta_{\rm C}$ 16.4 and one flavonol skeleton including one carbonyl group δ 176.2, two benzene rings, two oxygen-bearing olefinic carbons $\delta_{\rm C}$ 146.9, 135.4. Its ¹H NMR spectrum was

almost similar to that of kaempferol, except that an extra proton signal at $\delta_{\rm H}$ 4.22 (1H, s) was observed in **6** instead of one aromatic proton signal at $\delta_{\rm H}$ 6.48 (1H, br s) in kaempferol.¹⁷ In combination with the molecular formula C₃₁H₂₀O₁₂, two symmetrical kaempferol moieties were linked at C-8 and C-8' by one methylene group. This was also supported by the HMBC correlations of the methylene group $\delta_{\rm H}$ 4.22 (2H, s) with C-7 and C-7'' ($\delta_{\rm C}$ 162.1), C-8 and C-8'' ($\delta_{\rm C}$ 104.6), and C-9 and C-9'' ($\delta_{\rm C}$ 153.8). Thus, compound **6** was deduced as 3,5,7,4',3'',5'',7'',4'''- octahydroxyl-[8-CH $_2$ -8']-biflavone, and named dysosmabiflavonoid F.

Compound 7 was obtained as a yellow amorphous powder. Its ¹H and ¹³C NMR (Tables 1 and 2) were quite analogous to those of 6, except that one 1,3,4-trisubstituted benzene ring $\delta_{\rm H}$ 7.82 (1H, d, *J* = 2.0 Hz), 6.80 (1H, d, *J* = 8.5 Hz), 7.54 (1H, dd, *J* = 8.5 2.0 Hz) was observed in 7 instead of 1,4-disubstituted benzene ring in 6. This was further supported by their HR-ESI-MS, which gave an $[M + Na]^+$ quasi-molecular ion peak m/z623.0799 (Calcd 623.0802) in 7, being 16 mass-units more than that of 6. The quercetin subunit was identified as by the HMBC cross peak of the aromatic protons $\delta_{\rm H}$ 7.82 (1H, d, J = 2.0 Hz, H-2') and 7.54 (1H, dd, J = 8.5, 2.0 Hz, H-6') with C-2 (δ 146.7), one 1,2,3,4,5-tetra-substituted benzene ring $\delta_{\rm H}$ 6.12 (1H, s), and two oxygenated olefinic carbons $\delta_{\rm C}$ 146.7, 135.4. Thus, compound 7 3,5,7,3',4',3",5",7",4"''-nonahydroxyl-[8-CH₂-8']-biflavone, was and named dysosmabiflavonoid G.

By comparing their physical and spectroscopic data with literature values, the known metabolites were identified as kaempferol (8),¹⁷ quercetin (9),¹⁹ dihydrokaempferol (10),²⁰ dihydroquercetin (11).²⁰

All the isolates (1-11) were evaluated for their acetylcholinesterase inhibitory activity, using a modified spectrophotometric method.²¹ Tacrine was utilized as the positive control. The results are listed in Table 3 as IC_{50} values. Based on the IC_{50} values, all racemates 1-5 exhibited satisfactory AChE inhibitory activity in the range of micromolar concentrations (IC₅₀ values 2.72–6.74 μ M), and more potent inhibitory activity than their corresponding levorotatory or dextrorotatory enantiomers. Among all of biosynthetically related precursors (8-11), only quercetin (9) are more active than racemates 2–5. The combination between quercetin and dihydrokaempferol or between quercetin and dihydroquercetin by a furan ring reduced the inhibitory activity, for example, compared 2, 4 and 5 to 9. However, this phenomenon is inverse the combination between kaempferol and dihydrokaempferol or between kaempferol and dihydroguercetin, for example, compared 1 and 3 to 8. Hydrogenation (10 and 11) at C-2 and C-3 drastically reduced the inhibitory activity of the flavonol derivatives (8 and 9). The hydroxy group at C-3' is important, as 3',4'-dihydroxy derivative 9 showed more potent activity than its corresponding 4'-hydroxy analogue 8. The methylene linkage bridge between flavonol skeletons plays a very important role in maintaining inhibitory activity for this series of biflavonoids, for example, compared 8

 Table 3
 AChE inhibitory activities of all isolated compounds

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No.	$IC_{50} \left(\mu M \right)$	No.	$IC_{50}\left(\mu M\right)$
1	2.72 ± 0.25	5	6.74 ± 0.04
1a	32.95 ± 2.90	6	1.42 ± 0.14
1b	6.25 ± 0.57	7	0.73 ± 0.07
2	4.55 ± 0.36	8	17.90 ± 0.18
2a	4.97 ± 1.80	9	3.96 ± 0.27
3	6.39 ± 0.48	11	43.95 ± 2.43
3a	10.65 ± 1.43	2b, 4a, 4b, 5a, 5b, 10	>50
3b	9.89 ± 0.71	Tacrine	0.21 ± 0.01
4	6.33 ± 0.45		

and **9** to **7**, and **8** to **6**. In terms of the structure, racemates **1–5** are different completely from general biflavonoids. Moreover, due to the similarity in AChE inhibitory activity, they can be used as a template for the development of new drugs against neurodegenerative diseases.

Racemates 1–5 and their corresponding levorotatory and dextrorotatory enantiomers were measured for the potential to impede the generation of NO in lipopolysaccharide-stimulated RAW264.7 cells,²² and mushroom tyrosinase inhibitory effect.²³ Unfortunately, they were inactive (IC₅₀ > 50 μ M) against the inhibition of NO production. Only racemates 1 and 2 were found to display potent mushroom tyrosinase inhibitory activity with IC₅₀ values of 28.27 μ M and 39.75 μ M (the positive control, kojic acid, 32.59 μ M), respectively, whereas racemates 3–5 displayed no inhibitory effect against tyrosinase activity. The variation in inhibitory activity between them indicates the linkage bridges of C-2-O-C-7″:C-3-C-6″ were structurally required for anti-tyrosinase activity. Racemate 1, as the most potent tyrosinase inhibitor, has the therapeutic potential for skin hyperpigmentation and depigmentation in cosmetics.

Conclusions

So far, more than 600 natural biflavonoids have been found, and the vast majority of them are isolated from the plants of families Clusiaceae, Thymelaeaceae, Ochnaceae, and Selaginellaceae, etc.24 Only eleven biflavonoids have been obtained from plants of the genus Dysosma.13-16 Our phytochemical investigations on D. versipellis resulted in the isolation of five pairs of new biflavonoid enantiomers, (\pm) -dysosmabiflavonoids A-E (1-5), two new biflavonoids, dysosmabiflavonoids F-G (6-7), along with four biosynthetically related precursors (8-11). Racemates 1-2 represent the first report of natural C-2-O-C-7' ':C3-C6" biflavonoids. Based upon the IC₅₀ values, compound 7 was the most potent AChE inhibitor among all isolated compounds. Further studies on compound 7 are necessary to action mechanism and in vivo efficacy in neurodegenerative diseases. Above all, the present investigation will broaden the application field of D. versipellis.

Experimental method

General experimental procedures

Optical rotations and ECD spectra were measured by a Rudolph AP-IV polarimeter and an Applied Photophysics Chirascanq CD spectropolarimeter, respectively. UV and IR spectra were acquired using a ThermoEVO 300 spectrometer and a Thermo-Nicolet IS 10 spectrometer, respectively. NMR and mass spectra were recorded on a Bruker Avance III 500 spectrometer and a Bruker maXisHD mass spectrometer, respectively. Preparative HPLC separations were performed on a SEP system (Beijing Sepuruisi scientific Co., Ltd, China) equipped with a variable-wavelength UV detector, using a YMC-Pack ODS-A column ($250 \times 20 \text{ mm}, 5 \mu\text{m}$) and a Daicel chiral PAK AD-H column ($250 \times 10 \text{ mm}, 5 \mu\text{m}$). Sephadex LH-20 ($40-70 \mu\text{m}$), and silica gel (160-200 mesh) were acquired from Amersham Pharmacia Biotech AB, Uppsala, Sweden, and Marine Chemical Industry,

Qingdao, China, respectively. Chemical reagents for isolation were of analytical grade and purchased from Tianjin Siyou Co., Ltd, China. Biological reagents were from Sigma Company.

Plant material

The roots and rhizomes of *D. versipellis* were collected in Qingzhen, Guizhou Province, China, in July 2019, and identified by Prof. Cheng-Ming Dong at School of Pharmacy, Henan University of Chinese Medicine, where a voucher specimen (DV 20190706) was deposited.

Extraction and isolation

The powder of air-dried roots and rhizomes of D. versipellis (40 kg) were refluxed with 95% EtOH (v/v 120 L \times 3, 1.5 h each) and 50% EtOH (v/v 120 L \times 1, 1.5 h each) at 95 °C, respectively. The filtrate was concentrated under reduced pressure to yield a dark brown residue (5.4 kg). The residue was adsorbed by silicious earth and fractioned by CH₂Cl₂, EtOAc, and MeOH. The MeOH extract (3.4 kg) was separated into nine fractions M1-M9 by silica gel column chromatography (CC) with a gradient of CH₂Cl₂-MeOH (v/v 100:0, 100:1, 100:3, 100:5, 100:7, 100: 10, 100: 30, 100: 50, 0: 100) based on TLC monitoring results. Fraction M5 (110.0 g) was subjected to sephadex LH-20 CC eluted by methanol to yield subfractions M5-1-M5-8. Subfraction M5-8 (42.5 g) was submitted to silica gel CC eluted by PE-acetone (100:10, 100:20, 100:30, 100:40, 100:50, 100:70, 1:1, 1:2, 1:3) to afford nine subfractions (M5-8-1-M5-8-9). Subfraction M5-8-4 (0.5 g) was isolated by preparative HPLC (MeOH: H_2O , 60: 40) at a flow rate of 3 mL min⁻¹ to give compound 8 ($t_{\rm R}$ 40.5 min, 10.0 mg) and 6 ($t_{\rm R}$ 47.2 min, 6.0 mg). Subfraction M5-8-5 (1.07 g) was separated by preparative HPLC (MeOH: H_2O , 55: 45) at a flow rate of 3 mL min⁻¹ to afford subfractions [M5-8-5-1, (\pm) 2, $t_{\rm R}$ 37.4 min, 8.5 mg], (M5-8-5-2, $t_{\rm R}$ 46.0 min, 78.2 mg), (M5-8-5-3, t_R 72.2 min, 90.1 mg), [M5-8-5-4, (\pm) 3, $t_{\rm R}$ 77.0 min, 12.9 mg]. Subfraction M5-8-5-1 was isolated by preparative chiral HPLC (cyclohexane: isopropanol, 60:40) at a flow rate of 2 mL min⁻¹ to give 2a ($t_{\rm R}$ 7.9 min, 3.6 mg) and 2b ($t_{\rm R}$ 18.0 min, 3.6 mg). Subfraction M5-8-5-2 was chromatographed by preparative HPLC (CH₃CN:H₂O, 45:55) at a flow rate of 3 mL min $^{-1}$ to give subfractions [M5-8-5-2-1, (±) 5, $t_{\rm R}$ 25.7 min, 6 mg], [M5-8-5-2-2, (\pm) 1, $t_{\rm R}$ 29.1 min, 9.4 mg] and compound 11 (t_R 20.6 min, 10.9 mg). Subfraction M5-8-5-2-1 was isolated by preparative chiral HPLC (cyclohexane:isopropanol, 81:19) at a flow rate of 2 mL min⁻¹ to give 5a ($t_{\rm R}$ 31.8 min, 2.4 mg) and 5b (t_R 43.8 min, 2.4 mg). Subfraction M5-8-5-2-2 was subjected to preparative chiral HPLC (cyclohexane : isopropanol, 60:40) at a flow rate of 2 mL min⁻¹ to give 1a ($t_{\rm R}$ 7.8 min, 4.2 mg) and 1b (t_R 21.4 min, 4.2 mg). Subfraction M5-8-5-3 was isolated by preparative HPLC ($CH_3CN: H_2O, 43: 57$) at a flow rate of 3 mL min $^{-1}$ to give subfractions [M5-8-5-3-1, (±) 4, $t_{\rm R}$ 26.9 min, 5.5 mg] and 10 ($t_{\rm R}$ 22.5 min, 5.5 mg). Subfraction M5-8-5-3-1 was isolated by preparative chiral HPLC (cyclohexane: isopropanol, 81:19) at a flow rate of 2 mL min⁻¹ to give 4a ($t_{\rm R}$ 20.8 min, 2.5 mg) and 4b ($t_{\rm R}$ 30.2 min, 2.5 mg). Subfraction M5-8-5-4 was submitted to preparative chiral HPLC (cyclohexane : isopropanol, 70 : 30) at a flow rate of 2 mL min⁻¹

to give **3a** (t_R 9.7 min, 5.0 mg) and **3b** (t_R 12.7 min, 5.0 mg). Fraction M6 (130.0 g) was subjected to sephadex LH-20 CC eluted by methanol to yield subfractions M6-1, M6-2 and **11** (5 mg). The subfraction M6-2 (60.5 g) was applied to silica gel CC with a CH₂Cl₂–MeOH (100:1, 100:3, 100:5, 100:7, 100:10, 100:30) gradient to give six subfractions (M6-2-1–M6-2-6). Further separation of M6-2-4 (1.69 g) using sephadex LH-20 CC eluted by MeOH resulted in six subfractions M6-2-4-1–M6-2-4-6. Subfractions M6-2-4-4 (0.59 g) was purified by preparative HPLC (MeOH:H₂O, 60:40) at a flow rate of 3 mL min⁻¹ to afford compounds 7 (t_R 24.1 min, 3.0 mg) and **9** (t_R 20.4 min, 8.0 mg).

(±)-Dysosmabiflavonoid A (1). Yellow, amorphous powder; UV (MeOH) λ_{max} (log ε) 202 (4.33), 275 (3.58), 360 (3.52) nm; IR (iTR) ν_{max} 3374, 2958, 2928, 2853, 1726, 1635, 1607, 1512, 1465, 1374, 1284, 1179, 1123, 1075, 1022 cm⁻¹; HR-ESI-MS (positive): m/z 587.0806 [M + H]⁺ (Calcd for C₃₀H₁₉O₁₃, 587.0826), m/z609.0642 [M + Na]⁺ (Calcd for C₃₀H₁₈O₁₃Na, 609.0645), m/z625.0370 [M + K]⁺ (Calcd for C₃₀H₁₈O₁₃K, 625.0384); NMR data (DMSO- d_6), see Tables 1 and 2

(+)-Dysosmabiflavonoid A (1a). Yellow, amorphous powder; [α] $_{\rm D}^{20}$ + 41.5 (*c* 0.07, MeOH); ECD (MeOH) $\lambda_{\rm max}$ ($\Delta \varepsilon$) 229 (-0.8), 243 (-0.7), 272 (+2.7), 311 (-3.6), 367 (+1.2) nm.

(-)-Dysosmabiflavonoid A (1b). Yellow, amorphous powder; [α] $_{\rm D}^{20}$ – 41.8 (*c* 0.08, MeOH); ECD (MeOH) $\lambda_{\rm max}$ ($\Delta \varepsilon$) 229 (+0.8), 243 (+0.9), 273 (-2.5), 316 (+3.2), 368 (-1.4) nm.

(±)-Dysosmabiflavonoid B (2). Yellow, amorphous powder; UV (MeOH) λ_{max} (log ε) 202 (4.54), 256 (3.92), 298 (3.85), 370 (3.84) nm; IR (iTR) ν_{max} 3374, 2958, 2920, 2850, 1727, 1636, 1519, 1465, 1368, 1282, 1169, 1123, 1075, 1023 cm⁻¹; HR-ESI-MS (positive): m/z 587.0824 [M + H]⁺ (Calcd for C₃₀H₁₉O₁₃, 587.0826), m/z 609.0642 [M + Na]⁺ (Calcd for C₃₀H₁₈O₁₃Na, 609.0645), m/z 625.0383 [M + K]⁺ (Calcd for C₃₀H₁₈O₁₃K, 625.0384); NMR data (DMSO- d_6), see Tables 1 and 2.

(+)-**Dysosmabiflavonoid B (2a).** Yellow, amorphous powder; $[\alpha]_{D}^{20}$ + 39.0 (*c* 0.07, MeOH); ECD (MeOH) λ_{max} ($\Delta \varepsilon$) 219 (-2.9), 246 (-0.5), 272 (+3.3), 312 (-4.0), 370 (+1.8) nm.

(-)-Dysosmabiflavonoid B (2b). Yellow, amorphous powder; [α] $_{\rm D}^{20}$ – 42.0 (*c* 0.05, MeOH); ECD (MeOH) $\lambda_{\rm max}$ ($\Delta \varepsilon$) 219 (+3.4), 246 (+0.6), 273 (-3.3), 314 (+4.0), 369 (-2.0) nm.

(±)-Dysosmabiflavonoid C (3). Yellow, amorphous powder; UV (MeOH) λ_{max} (log ε) 203 (4.64), 271 (3.87), 299 (3.88), 367 (3.79) nm; IR (iTR) ν_{max} 3382, 2959, 2931, 2872, 1725, 1638, 1607, 1514, 1466, 1385, 1353, 1282, 1124, 1075, 1024 cm⁻¹; HR-ESI-MS (positive): m/z 587.0808 [M + H]⁺ (Calcd for C₃₀H₁₉O₁₃, 587.0826), m/z 609.0639 [M + Na]⁺ (Calcd for C₃₀H₁₈O₁₃Na, 609.0645), m/z 625.0375 [M + K]⁺ (Calcd for C₃₀H₁₈O₁₃K, 625.0384); NMR data (DMSO- d_6), see Tables 1 and 2

(-)-Dysosmabiflavonoid C (3a). Yellow, amorphous powder; $[\alpha]_{D}^{20} - 63.9 \ (c \ 0.06, MeOH); ECD \ (MeOH) \ \lambda_{max} \ (\Delta \varepsilon) \ 235 \ (+2.1), 290 \ (+4.0), 327 \ (-6.9), 379 \ (-0.2) \ nm.$

(+)-**Dysosmabiflavonoid** C (**3b**). Yellow, amorphous powder; $[\alpha]_{D}^{20}$ + 66.0 (*c* 0.05, MeOH); ECD (MeOH) λ_{max} ($\Delta \varepsilon$) 236 (-2.4), 291 (-5.0), 327 (+7.9), 379 (+0.3) nm.

(±)-Dysosmabiflavonoid D (4). Yellow, amorphous powder; UV (MeOH) λ_{max} (log ε) 202 (4.42), 257 (3.91), 299 (3.85), 379 (3.71) nm; IR (iTR) ν_{max} 3364, 2925, 2853, 1738, 1640, 1614, 1519, 1466, 1326, 1258, 1172, 1035 cm⁻¹; HR-ESI-MS (positive): m/z 587.0800 [M + H]⁺ (Calcd for C₃₀H₁₉O₁₃, 587.0826), m/z609.0636 [M + Na]⁺ (Calcd for C₃₀H₁₈O₁₃Na, 609.0645), m/z625.0371 [M + K]⁺ (Calcd for C₃₀H₁₈O₁₃K, 625.0384); NMR data (DMSO- d_6), see Tables 1 and 2.

(-)-Dysosmabiflavonoid D (4a). Yellow, amorphous powder; [α] $_{\rm D}^{20}$ – 64.4 (*c* 0.05, MeOH); ECD (MeOH) $\lambda_{\rm max}$ ($\Delta \varepsilon$) 233 (+1.5), 288 (+1.2), 322 (-1.9), 380 (-0.3) nm.

(+)-**DysosmabiflavonoidD** (4b). Yellow, amorphous powder; $[\alpha]_{D}^{20}$ + 64.8 (*c* 0.03, MeOH); ECD (MeOH) λ_{max} ($\Delta \varepsilon$) 234 (-1.9), 287 (-1.4), 317 (+2.3), 377 (+0.4) nm.

(±)-**Dysosmabiflavonoid** E (5). Yellow, amorphous powder; UV (MeOH) λ_{max} (log ε) 203 (4.41), 257 (3.82), 297 (3.78), 380 (3.63) nm; IR (iTR) ν_{max} 3377, 2924, 2853, 1739, 1638, 1608, 1463, 1305, 1265, 1173, 1124, 1027 cm⁻¹; HR-ESI-MS (positive): m/z 603.0752 [M + H]⁺ (Calcd for C₃₀H₁₉O₁₄, 603.0775), m/z625.0593 [M + Na]⁺ (Calcd for C₃₀H₁₈O₁₄Na, 609.0594), m/z641.0313 [M + K]⁺ (Calcd for C₃₀H₁₈O₁₄K, 641.0334); NMR data (DMSO- d_6), see Tables 1 and 2.

(–)-Dysosmabiflavonoid E (5a). Yellow, amorphous powder; [α] $_{\rm D}^{20}$ – 50.4 (*c* 0.05, MeOH); ECD (MeOH) $\lambda_{\rm max}$ ($\Delta \varepsilon$) 219 (+0.7), 235 (+0.8), 291 (+1.1), 321 (–1.8), 380 (–0.2) nm.

(+)-**Dysosmabiflavonoid** E (5b). Yellow, amorphous powder; $[\alpha]_{D}^{20}$ + 51.0 (*c* 0.04, MeOH); ECD (MeOH) λ_{max} ($\Delta \varepsilon$) 220 (-0.6), 237 (-0.7), 292 (-0.9), 326 (+1.2), 382 (+0.2) nm.

Dysosmabiflavonoid F (6). Yellow, amorphous powder; UV (MeOH) λ_{max} (log ε) 202 (4.89), 271 (4.69), 317 (4.46), 369 (4.51) nm; IR (iTR) ν_{max} 3361, 1647, 1563, 1510, 1379, 1266, 1188, 1159, 1022 cm⁻¹; HR-ESI-MS (positive): m/z 585.1023 [M + H]⁺ (Calcd for C₃₁H₂₁O₁₂, 585.1033); NMR data (DMSO- d_6), see Tables 1 and 2.

Dysosmabiflavonoid G (7). Yellow, amorphous powder; UV (MeOH) λ_{max} (log ε) 202 (5.05), 270 (4.80), 328 (4.56), 372 (4.70) nm; IR (iTR) ν_{max} 3247, 1650, 1604, 1559, 1512, 1424, 1372, 1316, 1259, 1159, 1114 cm⁻¹; HR-ESI-MS (positive): m/z 623.0799 [M + Na]⁺ (Calcd for C₃₁H₂₀O₁₃Na, 623.0802); NMR data (DMSO- d_6), see Tables 1 and 2.

AChE inhibitory assay in vitro

The AChE inhibitory activity was measured by a classical spectrophotometric method with minor modifications.²¹ A 200 μ L volume reaction system containing phosphate buffer (pH 8.0), tested compounds at different concentrations, and AChE (0.02 U mL⁻¹), was incubated for 20 min at 37 °C. Then, the reaction was initiated by the addition of 40 μ L of solution containing DTNB (0.625 mM) and acetylthiocholine iodide (0.625 mM) for AChE inhibitory activity assay, respectively. The hydrolysis of acetylthiocholine was monitored at 405 nm every 30 seconds for one hour. All the reactions were performed in triplicate. The IC₅₀ value of each compound was calculated by the Reed-Muench's method.

Assay for LPS-induced NO production

Test protocols were carried out as previously described.²² RAW 264.7 macrophages cells (2×10^5 cells per well) were precultured in 96-well microplates for 24 h. The tested compounds (50

 μ M) and L-NMMA with 1 μ g mL⁻¹ LPS were added and incubated at 37 °C for another 18 h. Nitric oxide production was determined by the Griess Reagent System. L-N^G-monomethyl arginine (L-NMMA) was used as a positive control.

Assay for tyrosinase inhibitory activity

Tyrosinase inhibitory activity was measured using an established assay protocol, with slight modification.²³ The tested compound was dissolved phosphate buffer solution containing 0.2% DMSO. The 60 μ L of phosphate buffer (20 mM, pH 6.8) and 50 μ L of the tested compounds at different concentrations, and 50 μ L of mushroom tyrosinase solution (20 μ g mL⁻¹) were placed into each well of a 96-well cell culture plate. After incubation for 10 min at 25 °C, 40 μ L of L-dopa (2.0 mM) was added to the reaction system. The reaction mixture was incubated for another 5 min, and the absorbance was measured at 490 nm. Kojic acid was used as standard agent. The IC₅₀ value of each compound was calculated as the concentration of the sample when the residual enzyme activity is 50%.

ECD computational method

All the quantum chemical ECDs were calculated by the literature method.²⁵ Conformational searches of compounds **1a–5a** and **1b–5b** were investigated by Spartan software using the MMFF94S force field, which gave four stable conformers, respectively. Then, those selected conformers were further optimized by density functional theory method at the B3LYP/6-31G* level. The ECD of these optimized conformers were calculated by the TDDFT method at the B3LYP/6-31+G* level with a CPCM model in MeOH solvent.

Author contributions

R. J. Han, H. Y. Bai and H. J. Wang performed the experiments, data analysis, and experimental planning. M. Li, Y. Y. Si, J. M. Wang, J. H. Gong and H. Chen screened the biological activities. The project was conceived and supervised by Y. J. Sun and W. S. Feng. The manuscript was written by Y. J. Sun. All authors reviewed the manuscript.

Conflicts of interest

There are no conflicts to declare.

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