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2-Benzyl-benzofurans from the tubers of *Ophiopogon japonicus*

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Abstract

Background: The overproduction of nitric oxide (NO) is known to involve in various inflammatory processes. A methanol extract of the tubers of *Ophiopogon japonicus* was found to strongly inhibit NO production. The present paper deals with the isolation, structural identification and NO inhibitory effect of five compounds isolated from the MeOH extract of *O. japonicus* tubers.

Results: Three new compounds were elucidated to be (2*R*)-(4-methoxybenzyl)-5,7-dimethyl-6-hydroxyl-2,3-dihydrobenzofuran (**1**), 2-(2-hydroxyl-4-methoxy-benzyl)-5-methyl-6-methoxyl-2,3-dihydrobenzofuran (**2**), and 2-(4-hydroxy-benzyl)-5,6-dihydroxybenzofuran (**3**). In addition, two known compounds were isolated from a natural source for the first time including 2-(4-methoxy-benzyl)-6,7-dimethoxyl-2,3-dihydrobenzofuran (**4**), and 2-(4-methoxy-benzyl)-6,7-methylenedioxy-2,3-dihydrobenzofuran (**5**). The absolute configuration of compound **1** was determined by experimental and calculated circular dichroism spectra. The effects of the isolated compounds on LPS-induced NO production in RAW264.7 cells were evaluated. Compound **1** and **2** showed the inhibitory activity with IC₅₀ values of 11.4 and 29.1 μM, respectively.

Conclusions: The class of 2-benzyl-2,3-dihydrobenzofuran is uncommon in nature. In this work, three such compounds were isolated from *O. japonicus*. Two of them showed promising anti-inflammatory activity by inhibition of NO production.

Keywords: *Ophiopogon japonicus*, Dihydrobenzofuran, 2-Benzyl-2,3-dihydroxybenzofuran, 2-Benzyl-benzofuran, Inhibition of NO production

Background

Ophiopogon japonicus (L.f) Ker-Gawl (Convallariaceae) occurs widely in Vietnam and it has been used in traditional medicine to treat cough, fever, epistaxis, inflammation, respiratory disease, constipation, and gastrointestinal disorders [1]. Steroidal saponins are among the main characteristic components of *O. japonicus* and have anticancer, anti-inflammatory, antioxidative, and neurotogenic effects [2–4]. Homoisoflavonoids are also

characteristic of *O. japonicus* and possess anti-inflammatory, antioxidative, and cytotoxic activities [4–8]. *O. japonicus* is also rich in polysaccharides that have anti-diabetic, antioxidative, anti-inflammatory, and immunomodulatory properties [4, 7]. In addition, phenolic acids, sesquiterpenes, fatty acids, and lignans have been identified from *O. japonicus* [4, 9, 10].

Nitric oxide (NO) is produced by inducible nitric oxide synthase (iNOS) in macrophages, hepatocytes, and renal cells. When produced in excess, NO directly damages normal tissues and triggers inflammation. Therefore, inhibitors of NO production have potential therapeutic value as anti-inflammatory agents [11]. In our search for anti-inflammatory compounds from natural sources, a methanol (MeOH) extract of the tubers of *O. japonicus* was found to strongly inhibit NO

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production. Phytochemical fractionation of the CHCl_3 -soluble fraction of the MeOH extract led to the isolation of five 2-benzyl benzofurans, including three new (1–3) and two known (4, 5) compounds (Fig. 1). Compound 1 strongly inhibited NO production in lipopolysaccharide (LPS)-induced RAW264.7 cells.

Results and discussion

Compound 1 was obtained as a brown solid. Its molecular formula was determined to be $\text{C}_{18}\text{H}_{20}\text{O}_3$ from high-resolution electrospray ionisation mass spectrometry (HRESIMS) (m/z 283.1365 $[\text{M} - \text{H}]^-$). Its ^1H NMR spectrum showed the characteristic resonance of an AA'BB' aromatic ring [δ_{H} 7.19 (2H, d, $J = 8.5$ Hz, H-2', 6'), and 6.86 (2H, d, $J = 8.5$ Hz, H-3', 5')], one aromatic singlet [δ_{H} 6.67 (1H, s, H-4)], one oxygenated methine proton [δ_{H} 4.86 (partially overlapped with HDO signal, H-2)], one methoxyl group [δ_{H} 3.78 (3H, s, 4'-OMe)], two methylene groups [δ_{H} 3.07 (1H, dd, $J = 15.0, 8.5$ Hz, Ha-3), 2.82 (1H, dd, $J = 15.0, 7.5$ Hz, Hb-3), 3.02 (1H, dd, $J = 14.0, 7.0$ Hz, Ha-7'), 2.84 (1H, dd, $J = 14.0, 6.5$ Hz, Hb-7')] and two aromatic methyl groups [δ_{H} 2.12 (3H, s, Me-5) and 2.05 (3H, s, Me-7)] (Table 1). The ^{13}C NMR and DEPT spectra revealed the presence of two methyl carbons at δ_{C} 9.2 (7-Me) and 16.5 (5-Me), two methylene carbons at δ_{C} 35.9 (C-3) and 42.0 (C-7'), one methoxy carbon at δ_{C} 55.7 (4'-OMe), one oxygenated methine carbon at δ_{C} 85.1 (C-2), five methine carbons at δ_{C} 123.9 (C-4), 131.4 (C-2', 6'), and 114.7 (C-3', 5'), and seven quaternary carbons at δ_{C} 153.8 (C-6, observed from HMBC spectrum), 158.1 (C-7a), and 159.8 (C-4'), 118.0 (C-3a), 117.3 (C-5), 108.1 (C-7), and 131.2 (C-1') (Table 2). These data suggested the 2-benzyl-2,3-dihydrobenzofuran skeleton of 1

[12, 13]. The HMBC correlations from aromatic singlet H-4 to C-3, C-3a, C-5, C6, C-7a, and from Me-5 to C-4, C-5, C-6, as well as from Me-7 to C-6, C-7, C-7a indicated the presence of a dihydrobenzofuran skeleton with a hydroxyl group located at C-6 and two methyl groups located at C-5 and C-7. The methoxyl group was placed on C-4' based on the HMBC correlation of the proton of this group with C-4' (Fig. 2). From these data, 1 was identified as 2-(4-methoxybenzyl)-5,7-dimethyl-6-hydroxyl-2,3-dihydrobenzofuran. The quantum chemical electronic circular dichroism (ECD) calculation method, based on time-dependent density functional theory (TDDFT), was used to determine of the absolute configuration at C-2 [14]. The predicted ECD patterns for 2R were consistent with the experimentally measured ECD of 1 (Fig. 3). Thus, compound 1 was assigned as (2R)-(4-methoxybenzyl)-5,7-dimethyl-6-hydroxyl-2,3-dihydrobenzofuran.

Compound 2 was obtained as a brown solid. Analysis of the HRESIMS spectrum indicated that compound 2 has molecular formula $\text{C}_{18}\text{H}_{20}\text{O}_4$ (m/z 301.1436 $[\text{M} + \text{H}]^+$). The ^1H NMR spectrum of 2 showed the presence of two aromatic singlets (δ_{H} 6.87 and 6.39), an aromatic ABX spin system [δ_{H} 6.48 (1H, d, $J = 2.5$ Hz, H-3'), 6.43 (1H, dd, $J = 8.0, 2.5$ Hz, H-5'), and 6.98 (1H, d, $J = 8.0$ Hz, H-6')], one aromatic methyl (δ_{H} 2.10) and two methoxy groups (δ_{H} 3.75 and 3.76). The ^{13}C NMR and DEPT spectra of 2 showed the characteristics of 2-benzyl-2,3-dihydrobenzofuran, which are similar to those of 1 with one oxygenated methine carbon at δ_{C} 85.8 (C-2) and two methylenes at 34.0 (C-3) and 36.9 (C-7) (Table 2). The HMBC correlations confirmed the positions of the methyl and methoxy groups (Fig. 2). In

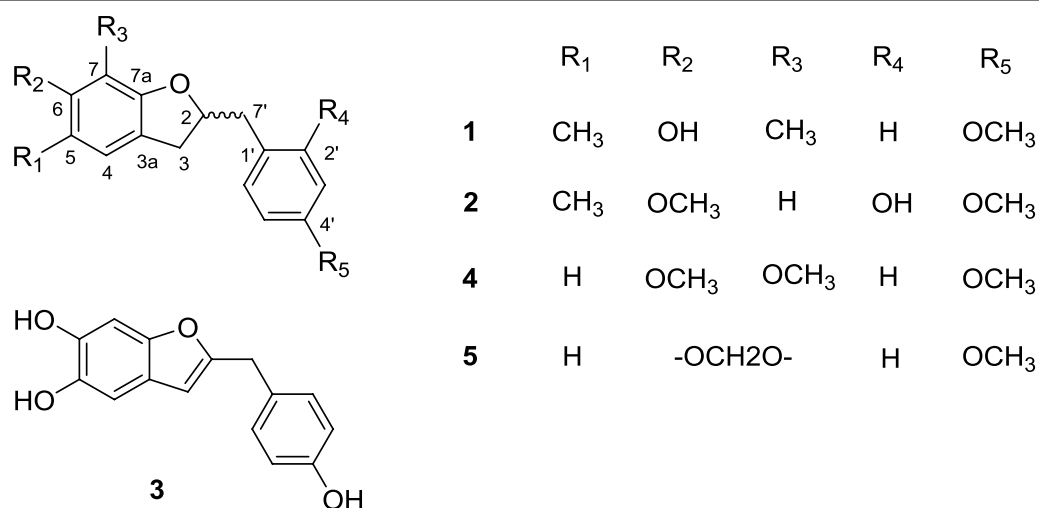


Fig. 1 Structure of compounds 1–5 isolated from *O. japonicus* tubers

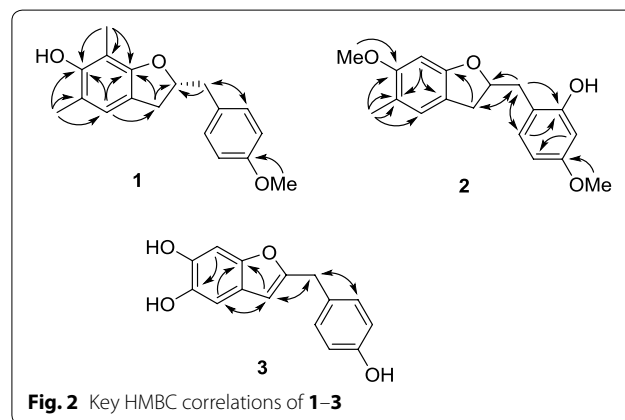
Table 1 ^1H NMR data of compounds 1–5 (δ_{H} in ppm, J in Hz)

Position	1 (CD ₃ OD)	2 (CDCl ₃)	3 (CD ₃ OD)	4 (CDCl ₃)	5 (CDCl ₃)
2	4.86, partially overlapped	5.05, m	–	5.00, m	5.02, m
3	3.07, dd (15.0, 8.5) 2.82, dd (15.0, 7.5)	3.15, dd (15.0, 8.5) 2.94, dd (15.0, 8.5)	6.16, s	3.12, dd (15.0, 8.0) 2.88, dd (15.0, 8.5)	3.09, dd (15.0, 8.5) 2.87, dd (15.0, 8.5)
4	6.67, s	6.87, s	6.82, s	6.74, d (8.0)	6.58, d (8.0)
5	–	–	–	6.38, d (8.0)	6.36, d (8.0)
7	–	6.39, s	6.83, s	–	–
2'	7.19, d (8.5)	–	7.10, d (8.5)	7.18, d (8.5)	7.17, d (8.5)
3'	6.86, d (8.5)	6.48, d (2.5)	6.74, d (8.5)	6.85, d (8.5)	6.85, d (8.5)
5'	6.86, d (8.5)	6.43, dd (8.0, 2.5)	6.74, d (8.5)	6.85, d (8.5)	6.85, d (8.5)
6'	7.19, d (8.5)	6.98, d (8.0)	7.10, d (8.5)	7.18, d (8.5)	7.17, d (8.5)
7'	3.02, dd (14.0, 7.0) 2.84, dd (14.0, 6.5)	3.07, dd (15.0, 3.5) 3.01, dd (15.0, 7.0)	3.91, s	3.16, dd (14.0, 6.0) 2.90, dd (14.0, 6.0)	3.13, dd (14.0, 6.0) 2.90, dd (14.0, 6.0)
5-Me	2.12, s	2.10, s	–	–	–
7-Me	2.05, s	–	–	–	–
4'-OMe	3.78, s	3.75, s	–	3.79, s	3.78, s
6-OMe	–	3.76, s	–	3.82, s	–
7-OMe	–	–	–	3.93, s	–
-OCH ₂ O-	–	–	–	–	5.90, s

Table 2 ^{13}C NMR data of compounds 1–5

Position	1 (CD ₃ OD)	2 (CDCl ₃)	3 (CD ₃ OD)	4 (CDCl ₃)	5 (CDCl ₃)
2	85.1	85.8	158.3	85.3	86.2
3	35.9	34.0	103.5	34.3	34.5
3a	118.0	117.2	122.0	121.2	122.9
4	123.9	126.2	106.0	118.3	116.8
5	117.3	119.0	143.1	109.9	104.3
6	153.8	157.8	144.3	152.1	148.7
7	108.1	93.6	98.6	133.7	130.0
7a	158.1	157.2	150.7	151.4	141.9
1'	131.2	116.2	130.1	129.3	129.1
2'	131.4	156.2	130.8	130.4	130.3
3'	114.7	102.9	116.2	113.9	113.9
4'	159.8	160.2	157.0	158.4	158.3
5'	114.8	106.4	116.2	113.9	113.9
6'	131.4	132.1	130.8	130.4	130.3
7'	42.0	36.9	34.8	40.9	40.7
5-Me	16.5	15.8	–	–	–
7-Me	9.2	–	–	–	–
4'-OMe	55.7	55.6	–	55.3	55.2
6-OMe	–	55.3	–	56.4	–
7-OMe	–	–	–	60.5	–
-OCH ₂ O-	–	–	–	–	100.2

a test of the optical rotation, this compound was optically inactive. Therefore, the structure of compound 2 was determined to be 2-(2-hydroxyl-4-methoxy-benzyl)-5-methyl-6-methoxyl-2,3-dihydrobenzofuran.



Compound 3 was obtained as a brown solid. Its HRESIMS revealed a molecular ion peak at m/z 255.0632 $[\text{M} - \text{H}]^-$ corresponding to a molecular formula of $\text{C}_{15}\text{H}_{12}\text{O}_4$. The ^1H NMR spectrum showed signals due to an olefinic proton [δ_{H} 6.16 (1H, s)], two aromatic protons [δ_{H} 6.82 (1H, s), and 6.83 (1H, s)], a 1,4-disubstituted benzene ring with two apparent doublets [δ_{H} 7.10 (2H, d, $J = 8.5$ Hz), 6.74 (2H, d, $J = 8.5$ Hz)], and a methylene group [δ_{H} 3.91 (2H, s)]. The ^{13}C NMR and DEPT experiments showed one methylene, seven methines, five oxygenated aromatic carbons, and two quaternary carbons. In the HMBC spectrum, the correlations between the olefinic proton (δ_{H} 6.16, H-3) and C-3a (δ_{C} 122.0), C-4 (δ_{C} 106.0), and C-7a (δ_{C} 150.7), H-4/C-3 (δ_{C} 103.5),

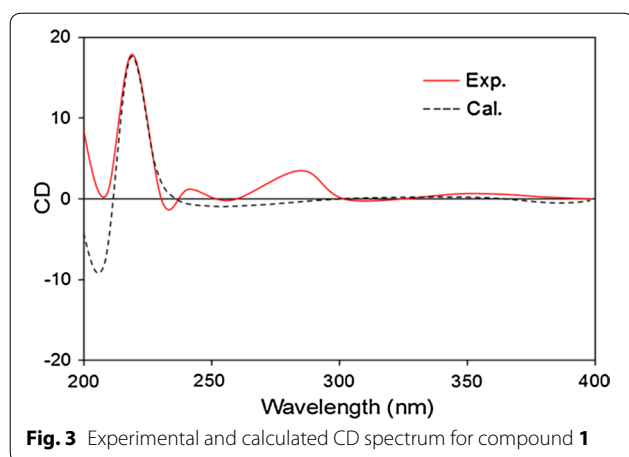


Fig. 3 Experimental and calculated CD spectrum for compound **1**

C-3a, C-5 (δ_C 143.1), C-6 (δ_C 144.3), and C-7a, as well as H-7/C-3a, C-5, C-6, and C-7a indicated the presence of a benzofuran skeleton with two oxygen substituents on ring A (Fig. 2). Similarly, the correlations between the 1,4-disubstituted protons (δ_H 7.10, H-2',6') and the oxygenated aromatic carbon (δ_C 157.0) and the methylene carbon (δ_C 34.8) indicated the presence of a 4-hydroxy-benzyl group. Moreover, the HMBC correlations of the methylene protons (δ_H 3.19, H-7') with C-2, C-3, and C-4 suggested the location of the 4-hydroxy-benzyl group at C-2 (Additional file 1). This compound was also optically inactive. Therefore, the structure of **3** was assigned as 2-(4-hydroxy-benzyl)-5,6-dihydroxybenzofuran.

Compounds **4** and **5** were identified as 2-(4-methoxy-benzyl)-6,7-dimethoxy-2,3-dihydrobenzofuran, and 2-(4-methoxy-benzyl)-6,7-methylenedioxy-2,3-dihydrobenzofuran, respectively [12]. Both compounds were isolated for the first time from the nature. It is noted that the class of 2-benzyl-2,3-dihydrobenzofuran is uncommon and only a few compounds of this type have been isolated so far from natural source [15].

The isolated compounds were tested for their ability to inhibit NO production in LPS-stimulated RAW264.7 cells. NO is an important signalling molecule in various physiological and pathophysiological responses, including the circulation, blood pressure, platelet function, and host defence. The overproduction of NO is also important in inflammation and related processes [16]. High NO levels are used as a marker for the treatment of inflammatory disorders. According to its specific functions and characteristics, inhibition of NO production by immune cells, typically macrophages, is suggested as one strategy for the development of anti-inflammatory agents. Our test revealed that compounds **1** and **2** reduced NO levels in LPS-stimulated RAW264.7 cells. Compound **1** was the most active compound with an IC_{50} of 11.4 μ M, while compound **2** had a moderate

effect (IC_{50} = 29.1 μ M). Compounds **3–5** were inactive up to the highest concentration tested (30 μ M). The cell viability test showed that neither compound had significant toxicity at its effective dose for NO inhibition (data not shown). 2,3-Dihydrobenzofurans have been known as potent anti-inflammatory compounds. Clossé et al. [17] demonstrated that the synthetic analogues of 2,3-dihydrobenzofuran-2-one had powerful anti-inflammatory activity in vivo, and 5-chloro-6-cyclohexyl-2,3-dihydrobenzofuran-2-one was significantly more potent than the reference compound, diclofenac, in all testing models. More recently, a series of dihydrobenzofurans was isolated from the seeds of *Prunus tomentosa*, some of which strongly inhibited NO production in LPS-stimulated BV-2 cells [18]. Consistently, our results suggest that *O. japonicus* is a potential natural source of anti-inflammatory dihydrobenzofurans.

Methods

General experimental procedures

Optical rotation values were recorded on a JASCO P-2000 digital polarimeter (JASCO, Tokyo, Japan). The IR spectra were obtained from a Tensor 37 FT-IR spectrometer (Bruker, Ettlingen, Germany). CD spectra were obtained with a JASCO J-1100 spectropolarimeter. NMR experiments were carried out on a Bruker AM500 FT-NMR spectrometer (Bruker, Rheinstetten, Germany) using residual solvent peak as a reference or tetramethylsilane (TMS) as internal standard. The HR-ESI-MS were recorded on a Waters Q-TOF micromass spectrometer Waters Q-TOF micromass spectrometer and an LTQ Orbitrap XLTM Mass spectrometer. Absorbance of bioassay solutions was read on an xMark microplate spectrophotometer.

Plant materials

The tubers of *O. japonicus* were collected in Feb. 2014 at Me Linh, Hanoi and identified by Prof. Tran Huy Thai, Institute of Ecology and Biological Resources, Vietnam Academy of Science and Technology. The voucher specimens were deposited at the Department of Bioactive Products, Institute of Marine Biochemistry, Vietnam Academy of Science and Technology.

Extraction and isolation

The air-dried and powdered tubers of *O. japonicus* (2.4 kg) were extracted with methanol (4 L \times 3 times) in a sonic bath for 30 min at 40 °C. The combined extracts were concentrated under a vacuum to obtain a crude residue (360 g), which was then resuspended in water (2 L), and extracted by chloroform (1 L \times 3 times) to obtain chloroform (8 g) and water residues. The chloroform residue was chromatographed on a silica gel column eluted with a gradient of 1–100% ethyl acetate in hexane

to afford nine fractions F1–F9. Fraction F1 was fractionated on a silica gel column eluted with hexane–ethyl acetate (20:1 v/v) to give nine subfractions F1.1–F1.9. Compound **5** (69.5 mg) was purified from F1.4 by using a reverse phase C₁₈ column eluted with acetone–water (2:1 v/v). Compound **1** (70.0 mg) and **4** (18.2 mg) were isolated from F1.7 by using a reverse phase C₁₈ column eluted with acetone–water (3:2 v/v). The F1.9 was chromatographed on a silica gel column (hexane–acetone 8:1, v/v) to afford **2** (20.8 mg). Compound **3** (20.0 mg) was obtained from F9 by repeated C₁₈ column (methanol–water 1:1 v/v) and silica gel column (dichloromethane–acetone 6:1, v/v).

(2R)-(4-methoxybenzyl)-5,7-dimethyl-6-hydroxyl-2,3-dihydrobenzofuran (1)

Brown solid. $[\alpha]_D^{24} = +86.5$ (*c* 0.05, MeOH). IR (KBr): 3446, 1615, 1513, 1472, 1247, 1097. CD (MeOH): 220 (+17.8), 288 (+3.2). ¹H-(CD₃OD, 500 MHz) and ¹³C-NMR (CD₃OD, 125 MHz): see the Tables 1 and 2. HR-ESI-MS (neg.): 283.1365 [M – H][–] (calc. for C₁₈H₁₉O₃, 283.1334).

2-(2-Hydroxyl-4-methoxy-benzyl)-5-methyl-6-methoxyl-2,3-dihydrobenzofuran (2)

Brown solid. IR (KBr): 3440, 1610, 1482, 1245, 1120. ¹H-(CDCl₃, 500 MHz) and ¹³C-NMR (CDCl₃, 125 MHz): see the Tables 1 and 2. HR-ESI-MS (pos.): 301.1436 [M + H]⁺ (calc. for C₁₈H₂₁O₄, 301.1440).

2-(4-Hydroxy-benzyl)-5,6-dihydroxybenzofuran (3)

Brown solid. IR (KBr): 3450, 1613, 1514, 1484, 1242, 1142. ¹H-(CD₃OD, 500 MHz) and ¹³C-NMR (CD₃OD, 125 MHz): see the Tables 1 and 2. HR-ESI-MS (neg.): 255.0632 [M – H][–] (calc. for C₁₅H₁₁O₄, 255.0657).

Assay for inhibition of NO production

The effects of compounds on the NO production in LPS-stimulated RAW264.7 cells were evaluated as previously described [19]. The cells were seeded in 96-well plate at 2 × 10⁵ cells/well and incubated for 12 h. The plate were pretreated with compounds in various concentrations (from 1 to 30 μM) for 30 min and then incubated for another 24 h with or without 1 μg/ml LPS. 100 μl of the culture supernatant were transferred to other 96-well plate and 100 μl of Griess reagent were added. The absorbance of the reaction solution was read at 570 nm with a microplate reader (XMark microplate reader, Biorad, USA). The remaining cell solutions in cultured 96-well plate were used to evaluate cell viability by 3-(4,5-dimethylthiazole-2-yl)-2,5-diphenyl tetrazolium bromide (MTT) assay. Cardamonin was used as a positive control (IC₅₀ = 2.80 μM).

Conclusion

In summary, three 2-benzyl-2,3-dihydrobenzofurans and two 2-benzyl-benzofurans were isolated from the CHCl₃-soluble fraction of the MeOH extract of *O. japonicus* tubers. Their structures were found to be (2R)-(4-methoxybenzyl)-5,7-dimethyl-6-hydroxyl-2,3-dihydrobenzofuran (**1**), 2-(2-hydroxyl-4-methoxy-benzyl)-5-methyl-6-methoxyl-2,3-dihydrobenzofuran (**2**), 2-(4-hydroxy-benzyl)-5,6-dihydroxybenzofuran (**3**), 2-(4-methoxy-benzyl)-6,7-dimethoxyl-2,3-dihydrobenzofuran (**4**), and 2-(4-methoxy-benzyl)-6,7-methylenedioxy-2,3-dihydrobenzofuran (**5**). Compounds **1** and **2** inhibited NO production in LPS-stimulated RAW264.7 cells.

Additional file

Additional file 1. NMR spectra of compounds **1–3**.

Authors' contributions

NHD and NTD designed research and wrote the paper; NDC, HMT and NVT performed research; NTH and DL analyzed CD data. All authors read and approved the final manuscript.

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Competing interests

The authors declare that they have no competing interests.

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References

- Do TL (2004) Vietnamese medicinal herbs and remedies. Medicinal Publisher, Hanoi
- Qi J, Hu ZF, Zhou YF, Hu YJ, Yu BY (2015) Steroidal saponins and glycosides from the fibrous roots of *Ophiopogon japonicus* and *Liriope spicata* var. *prolifera* with anti-inflammatory activity. *Chem Pharm Bull* 63:187–194
- Ye Y, Qu Y, Tang R, Cao S, Yang W, Xiang L, Qi J (2013) Three new neurotogenic steroidal saponins from *Ophiopogon japonicus* (Thunb.) Ker-Gawl. *Steroids* 78:1171–1176
- Chen MH, Chen XJ, Wang M, Lin LG, Wang YT (2016) *Ophiopogon japonicus*—a phytochemical, ethnomedicinal and pharmacological review. *J Ethnopharmacol* 181:193–213
- Duan CL, Kang ZY, Lin CR, Jiang Y, Liu JX, Tu PF (2009) Two new homoisoflavonoids from the fibrous roots of *Ophiopogon japonicus* (Thunb.) Ker-Gawl. *J Asian Nat Prod Res* 11:876–879

6. Hung TM, Thu CV, Dat NT, Ryoo SW, Lee JH, Kim JC, Na M, Jung HJ, Bae K, Min BS (2010) Homoisoflavonoid derivatives from the roots of *Ophiopogon japonicus* and their in vitro anti-inflammation activity. *Bioorg Med Chem Lett* 20:2412–2416
7. Fan Y, Ma X, Zhang J, Ma L, Gao Y, Zhang W, Song X, Hou W, Guo C, Tong D (2015) *Ophiopogon* polysaccharide liposome can enhance the non-specific and specific immune response in chickens. *Carbohydr Polym* 119:219–227
8. Lin Y, Zhu D, Qi J, Qin M, Yu B (2010) Characterization of homoisoflavonoids in different cultivation regions of *Ophiopogon japonicus* and related antioxidant activity. *J Pharm Biomed Anal* 52:757–762
9. Cheng ZH, Wu T, Bligh SW, Bashall A, Yu BY (2004) *cis*-Eudesmane sesquiterpene glycosides from *Liriope muscari* and *Ophiopogon japonicus*. *J Nat Prod* 67:1761–1763
10. Liu C, Ju A, Zhou D, Li D, Kou J, Yu B, Qi J (2016) Simultaneous qualitative and quantitative analysis of multiple chemical constituents in YiQiFuMai injection by ultra-fast liquid chromatography coupled with ion trap time-of-flight mass spectrometry. *Molecules* 21:E640
11. Luiking YC, Engelen MP, Deutz NE (2010) Regulation of nitric oxide production in health and disease. *Curr Opin Clin Nutr Metab Care* 13:97–104
12. Qi P, Jiang J, Qi H, Jin Y, Shen Q, Wu Y, Song H, Zhang W (2013) Synthesis and antiproliferative activity of new polyoxo 2-benzyl-2,3-dihydrobenzofurans and their related compounds. *Lett Drug Des Discov* 10:886–894
13. Ortega N, Urban S, Beiring B, Glorius F (2012) Ruthenium NHC catalyzed highly asymmetric hydrogenation of benzofurans. *Angew Chem Int Ed* 51:1710–1713
14. Kwon J, Hiep NT, Kim DW, Hong S, Guo Y, Hwang BY, Lee HJ, Mar W, Lee D (2016) Chemical constituents isolated from the root bark of *Cudrania tricuspidata* and their potential neuroprotective effects. *J Nat Prod* 79:1938–1951
15. Tsai YC, Hsu CC, El-Shazly M, Chiang SY, Wu CC, Wu CC, Lai WC, Yen MH, Wu YC, Chang FR (2015) Phytochemicals and estrogen-receptor agonists from the aerial parts of *Liriope platyphylla*. *Molecules* 20:6844–6855
16. Aktan F (2004) iNOS-mediated nitric oxide production and its regulation. *Life Sci* 75:639–653
17. Closse A, Haefliger W, Hausser D, Gubler HU, Dewald B, Baggiolini M (1981) 2,3-Dihydrobenzofuran-2-ones: a new class of highly potent antiinflammatory agents. *J Med Chem* 24:1465–1471
18. Liu QB, Huang XX, Bai M, Chang XB, Yan XJ, Zhu T, Zhao W, Peng Y, Song SJ (2014) Antioxidant and anti-inflammatory active dihydrobenzofuran neolignans from the seeds of *Prunus tomentosa*. *J Agri Food Chem* 62:7796–7803
19. Dat NT, Binh PT, Quynh LTP, Huong HT, Minh CV (2012) Sanggenon C and O inhibit NO production, iNOS expression and NF- κ B activation in LPS-induced RAW264.7 cells. *Immunopharmacol Immunotoxicol* 34:84–88

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