A New Flavone Glycoside from Lumnitzera littorea with in vitro -Glucosidase Inhibitory Activity

Nguyen, Thuy L.T.; Pham, Thuy T.; Bui, Tung T.; Huynh, Loc T.; Truong, Dang T.T.; Le, Ngoc L.; Nguyen, Duc X. ; Le, Dung L.; Hansen, Poul Erik; Nguyen, K. P. Phung

Published in:
Natural Product Communications

DOI:
10.1177/1934578X19851361

Publication date:
2019

Document Version
Publisher's PDF, also known as Version of record

Citation for published version (APA):
Nguyen, T. L. T., Pham, T. T., Bui, T. T., Huynh, L. T., Truong, D. T. T., Le, N. L., Nguyen, D. X., Le, D. L., Hansen, P. E., & Nguyen, K. P. P. (2019). A New Flavone Glycoside from Lumnitzera littorea with in vitro - Glucosidase Inhibitory Activity. Natural Product Communications, 14(6), 1-5. https://doi.org/10.1177/1934578X19851361

General rights
Copyright and moral rights for the publications made accessible in the public portal are retained by the authors and/or other copyright owners and it is a condition of accessing publications that users recognise and abide by the legal requirements associated with these rights.

• Users may download and print one copy of any publication from the public portal for the purpose of private study or research.
• You may not further distribute the material or use it for any profit-making activity or commercial gain.
• You may freely distribute the URL identifying the publication in the public portal.

Take down policy
If you believe that this document breaches copyright please contact rucforsk@ruc.dk providing details, and we will remove access to the work immediately and investigate your claim.
A New Flavone Glycoside From *Lumnitzera littorea* with In Vitro α-Glucosidase Inhibitory Activity

Nguyen T. L. Thuy, Pham T. Thuy, Bui T. Tung, Huynh T. Loc, Truong T. T. Dang, Le L. Ngoc, Nguyen X. Duc, Le T. Dung, Poul E. Hansen, and Nguyen K. P. Phung

Abstract

A new flavone glycoside, lumnitzerone (1), was isolated from leaves of *Lumnitzera littorea*, together with 9 known flavonoids. Their structures were elucidated by spectroscopic (one-dimensional, two-dimensional nuclear magnetic resonance) and high-resolution mass spectrometry analysis, and comparison with literature data. Extracts and all isolated compounds were evaluated for α-glucosidase inhibitory activity; all the extracts and most of the isolated compounds exhibited better activities than the positive control acarbose.

Keywords

*Lumnitzera littorea*, flavonoids, α-glucosidase inhibitory activity

Received: October 1st, 2018; Accepted: February 19th, 2019.

α-Glucosidase is an intestinal enzyme that breaks down α-1,4 linked polysaccharides to α-glucose, which leads to high blood sugar levels. The development of an α-glucosidase inhibitor derived from a natural product could be an important contribution to diabetes prevention. *Lumnitzera littorea* (family Combretaceae) grows in Can Gio mangrove forest in Vietnam. There have been 2 reports about the phytochemistry of this species. The antimicrobial activities of *n*-hexane, ethyl acetate, and methanol extracts of the leaves were evaluated against 6 human pathogenic microbes and the first extract was the most active. Recently, we reported the isolation of gallic acid and naringenin. Here, we present the chemical constituents as well as the α-glucosidase inhibitory activity of extracts and compounds isolated from *L. littorea* leaves.

The ethyl acetate extract of *L. littorea* leaves yielded one new and 9 known flavonoids. The new flavonoid glycoside (1) was obtained as a light yellow solid and appeared purple on thin-layer chromatography (TLC) under UV light at 365 nm. Its molecular formula was established as C_{25}H_{24}O_{11} through the pseudomolecular ion peak in the high-resolution electrospray ionization mass spectrometry (HR-ESI-MS) at m/z 501.1398 [M+H]^+ (calculated for C_{25}H_{24}O_{11}+H, 501.1397). The 1^H and 1H−1H correlation spectroscopy nuclear magnetic resonance (COSY NMR) spectra of 1 revealed signals of AA’BB’-type aromatic protons at δ 7.96 (2H, d, 8.5 Hz) and 6.94 (2H, d, 8.5 Hz), and 2 meta coupled protons at δ 6.43 (1H, d, 2.0 Hz) and 6.78 (1H, d, 2.0 Hz), which were assigned to H-2’/H-6’, H-3’/H-5’, H-6 and H-8, respectively. In addition, an olefinic proton at δ 6.87 (1H, s, H-3), and a chelated hydroxy proton and a phenolic proton at δ 12.96 (1H, s) and 10.40 (1H, s), respectively, were observed. All these data suggested the presence of an apigenin aglycone. Moreover, the 1^H NMR spectrum showed the presence of a hexopyranose unit with the anomeric proton at δ 5.13 (d, 7.5 Hz, H-1”) and signals due to sugar protons in the region of δ 3.15 to 5.50. Furthermore, characteristic signals of an (E)-propenyl group were observed at δ 5.83 (1H, dd, 15.5, 1.5 Hz, H-2’”), 6.85 (1H, dq, 15.5, 7.0 Hz, H-3’”), and 1.64 (3H, d, 7.0 Hz, H-4’”). The 13^C NMR spectrum of 1 revealed 25 carbon signals, including signals of 2 benzene rings of the apigenin aglycone at δ 164.3 (C-2), 103.1...
The known compounds were identified from spectroscopic analysis and comparison with literature data as quercetin (2), quercitrin (3), myricetin (4), myricitrin (5), naringenin (6), chrysoeriol (7), pilloin (8), afzelin (9), and myricetin 3-O-(4″-O-galloyl)-α-L-rhamnopyranoside (10) (Figure 1).

The in vitro α-glucosidase inhibitory activity was evaluated of the extracts and purified compounds (Tables 1 and 2). All the extracts and most of the isolated compounds exhibited better activities than the positive control acarbose. The flavonoid-type structure, the position, and the number of hydroxy groups are determining factors for α-glucosidase inhibition. The A-ring 7-OH and the B-ring 4′-OH groups play an important role in the inhibitory effect. This observation was proved by the most potent inhibitors such as naringenin (6), quercetin (2), and afzelin (9), which possess half-maximal inhibitory concentration (IC50) values of 1.87, 3.42, and 6.26 µg/mL, respectively, compared with that of lumnitzerone (1) (IC50 11.31 µg/
mL). On the contrary, enhancement of the number of hydroxy groups on the B ring or if the hydroxy group is methylated reduces the inhibitory activity, as seen with chrysoeriol (7), pilloin (8), and myricitrin (5) (IC50 53.4, 89.1, and 153.5 µg/mL, respectively). The presence of a sugar unit did not show a clear effect on the inhibition. Careful analysis of the chemical shifts, multiplicities, and coupling constant magnitudes in the 1H NMR spectrum, along with the 1H-1H COSY and 13C NMR spectra of 1 (Table 3) indicated that the sugar was a β-D-glucopyranose unit. The nature of the sugar was further confirmed from the anomeric proton data of the sugar obtained after acid hydrolysis.11

The multiplicity of the protons H3-4‴, H-3‴, and H-2‴ and the 1H-1H COSY spectrum confirmed their contiguous arrangement. The heteronuclear multiple bond correlation (HMBC) experiments showed cross-peaks (Figure 2) of the propenyl protons at δ 5.83 (H-2‴) and 6.85 (H-3‴), as well as the methylene protons at δ 4.41 (H-6‴a) and 4.08 (H-6‴b) to the carbonyl carbon C-1‴ (δ 165.3) suggesting that the butenoyloxy group was attached at C-6‴ of the D-glucose. The attachment of this sugar to C-7 in the apigenin nucleus was confirmed by the HMBC cross-peak of the anomeric proton at δ 5.13 (1H, d, 7.5 Hz) to carbon C-7 (δ 162.6). Accordingly, 1 was identified as apigenin 7-O-[6‴-(E)-butenoyl-β-D-glucopyranoside], for which the trivial name lumnitzerone is proposed.

Experimental

General Experimental Procedure

The NMR spectra were recorded on a Bruker Avance III spectrometer at 500 MHz for 1H NMR and 125 MHz for 13C NMR spectra. HR-ESI-MS were obtained on a Shimadzu +IDA time-of-flight MS. TLC was performed on silica gel 60 F254 (Merck, Darmstadt, Germany). Gravity column chromatography was performed on silica gel 60 (0.040-0.063 mm, Merck) and Sephadex LH-20 (GE Healthcare Bio-Science AB, Uppsala, Sweden). α-Glucosidase (EC 3.2.1.20) from Saccharomyces cerevisiae (750 UN) and p-nitrophenyl-α-D-glucopyranoside were purchased from Sigma Chemical Co. (St Louis, MO, USA). Acarbose and dimethyl sulfoxide were obtained from Merck. Other chemicals were of the highest grade available.

Plant Material

Leaves of Lumnitzera littorea (Jack) Voigt (Combretaceae) were collected at Can Gio mangrove forest, Ho Chi Minh City, Viet Nam in August of 2014. The scientific name of the plant was authenticated by Dr Pham Van Ngot, Faculty of Biology, Ho Chi Minh City University of Pedagogy. A voucher specimen (No US-B012) was deposited in the herbarium of the Department of Organic Chemistry, University of Science.

Extraction and Isolation

The fresh leaves were washed under running tap water to remove all sand particles and epiphytes and then were dried and ground into fine powder. The powder (15 000 g) was exhaustively extracted with ethanol at room temperature by maceration. After filtration the ethanol solution was evaporated to dryness under reduced pressure to yield a crude ethanol...
residue (1000 g). This was applied to a silica gel solid phase column and eluted consecutively with n-hexane, ethyl acetate, and finally with ethanol. After evaporation under reduced pressure, 3 extracts were obtained, n-hexane (100 g), ethyl acetate (250 g), and ethanol (550 g). The ethyl acetate extract (250 g) was fractionated by silica gel column chromatography using a mixture of EtOAc−MeOH (99:1 to 0:100) to yield 8 fractions (EA1-EA8). These were then continuously separated using silica gel and Sephadex LH-20 and eluted with appropriate solvent systems of EtOAc−MeOH to give 10 compounds. As a result, fraction EA2 afforded 2 (50 mg), 3 (15 mg), 4 (10 mg), and 5 (10 mg), fraction EA3 gave 6 (3 mg) and 7 (5 mg), EA5 gave 1 (5 mg) and 10 (10 mg), and EA7 gave 8 (5 mg) and 9 (8 mg).

Bioactivity Assay

The α-glucosidase inhibitory activity was determined according to the method of Apostolidis et al. The inhibitory activity was calculated using the following equation:

\[
\% \text{Inhibition} = \left( \frac{A_{\text{control}} - A_{\text{sample}}}{A_{\text{control}}} \right) \times 100
\]

The IC_{50} values were determined from plots of percent inhibition vs log inhibitor concentration and calculated by non-linear regression analysis from the mean inhibitory values.

Lumnitzerone (1)

Light yellow solid. 

\(^1\)H, \(^{13}\)C NMR and HMBC (DMSO-\(d_6\)): Table 3.

HR-ESI-MS: \(m/z\) [M+H] + calculated for \(C_{25}H_{24}O_{11}^+H, 501.1397\); found: 501.1398.

Acid Hydrolysis of 1

Compound 1 (5 mg) was treated with HCl 0.2 M (dioxane/\(H_2O, 1/1, v/v\), 200 µL) at 95°C for 3 hours. After cooling, the reaction mixture was extracted with chloroform (3 × 2 mL) to eliminate the aglycone component. The remaining solution was evaporated to dryness. The obtained residue was dissolved in D_{2}O for subsequent \(^1\)H NMR analysis. The anomeric ratios were obtained by manual integration with \(\delta_H\) 5.23 (d, \(J = 3.5\) Hz, 36.4%) and 4.64 (d, \(J = 8.0\) Hz, 63.6%). These values were highly reminiscent of those of glucose.

Declaration of Conflicting Interests

The author(s) declared no potential conflicts of interest with respect to the research, authorship, and/or publication of this article.

Funding

The author(s) disclosed receipt of the following financial support for the research, authorship, and/or publication of this article: This

| Position | \(\delta_H\) (multiplicity) | \(\delta_C\) | HMBC (\(^1\)H → \(^{13}\)C) |
|----------|-----------------------------|-------------|-----------------------------|
| 2        | 164.3                       |             |                             |
| 3        | 6.87 (1H, s)                | 103.1       | C-2, 4, 10, 1'               |
| 4        | 182.0                       |             |                             |
| 5        | 161.4                       |             |                             |
| 5-OH     | 12.96 (1H, s)               |             | C-5, 6, 10                  |
| 6        | 6.43 (1H, d, 2.0)           | 99.4        | C-5, 7, 8, 10               |
| 7        |                             | 162.6       |                             |
| 8        | 6.78 (1H, d, 2.0)           | 94.7        | C-4, 6, 7, 9, 10            |
| 9        |                             | 156.9       |                             |
| 10       |                             | 105.4       |                             |
| 1'       |                             | 121.0       |                             |
| 2, 6'    | 7.96 (2H, d, 8.5)           | 128.6       | C-2', 3', 4', 5', 6'        |
| 3', 5'   | 6.94 (2H, d, 8.5)           | 116.0       | C-1', 2', 3', 4', 5'        |
| 4'       |                             | 161.1       |                             |
| 4''-OH   | 10.40 (1H, s)               |             | C-3', 4', 5'                |
| 1''      | 5.13 (1H, d, 7.5)           | 99.5        | C-7                         |
| 2''      | 3.28 (,)                    | 72.9        | C-1'', 3''                  |
| 2''-OH   | 5.48 (1H, d, 5.0)           |             | C-2''                       |
| 3''      | 3.30 (, )                   | 76.2        | C-5''                       |
| 3''-OH   | 5.24 (1H, d, 4.5)           |             | C-3''                       |
| 4''      | 3.18 (1H, m)                | 70.0        | C-5''                       |
| 4''-OH   | 5.35 (1H, d, 5.5)           |             | C-3'', 4', 5''              |
| 5''      | 3.79 (1H, t, 8.0)           | 73.8        | C-1'', 3'', 4', 6''         |
| 6''a     | 4.41 (1H, d, 11.0)          | 63.4        | C-1''''                     |
| 6''b     | 4.08 (1H, dd, 11.0, 8.0)    |             | C-1'''', 5''                |
| 1''''    |                             | 165.3       |                             |
| 2''''    | 5.83 (1H, dd, 15.5, 1.5)    | 122.0       | C-1'''', 3'''', 4''''       |
| 3''''    | 6.85 (1H, dq, 15.5, 7.0)    | 145.3       | C-1'''', 2''', 4''''       |
| 4''''    | 1.64 (3H, d, 7.0)           | 17.4        | C-2'''', 3''''              |

HMBC, heteronuclear multiple bond correlation; DMSO, dimethyl sulfoxide.

(*) overlapped in the solvent signal.

Table 3. Nuclear Magnetic Resonance Data of Compound 1.

Figure 2. Key COSY and HMBC correlations of compound 1.

COSY: correlation spectroscopy; HMBC: heteronuclear multiple bond correlation.

Acid Hydrolysis of 1

Compound 1 (5 mg) was treated with HCl 0.2 M (dioxane/\(H_2O, 1/1, v/v\), 200 µL) at 95°C for 3 hours. After cooling, the reaction mixture was extracted with chloroform (3 × 2 mL) to eliminate the aglycone component. The remaining solution was evaporated to dryness. The obtained residue was dissolved in D_{2}O for subsequent \(^1\)H NMR analysis. The anomeric ratios were obtained by manual integration with \(\delta_H\) 5.23 (d, \(J = 3.5\) Hz, 36.4%) and 4.64 (d, \(J = 8.0\) Hz, 63.6%). These values were highly reminiscent of those of glucose.

Declaration of Conflicting Interests

The author(s) declared no potential conflicts of interest with respect to the research, authorship, and/or publication of this article.

Funding

The author(s) disclosed receipt of the following financial support for the research, authorship, and/or publication of this article: This
research is funded by HoChiMinh City Open University under the grant number E2017.2.11.1.

**Supplemental Material**

Supplemental material for this article is available online.

**References**

1. Saad S, Taher M, Susanti D, Qaralleh H, Rahim NABA. Anti-microbial activity of mangrove plant (*Lumnitzera littorea*). *Asian Pac J Trop Med*. 2011;4(7):523-525.

2. Thuy NTL, Hung QT, Duc NT, et al. Flavonoids from the leaves of *Lumnitzera littorea*. *Viet J Chem*. 2017;55:606-610.

3. Liu H, Mou Y, Zhao J, et al. Flavonoids from *Halostachys caspica* and their antimicrobial and antioxidant activities. *Molecules*. 2010;15(11):7933-7945.

4. Ishikawa T, Navarro LB, Donatini RS, et al. Gastroprotective property of *Plinia edulis* (Vell.) Sobral (Myrtaceae): the role of triterpenoids and flavonoids.. *Pharmacol Online*. 2014;1:36-43.

5. He D, Gu D, Huang Y, et al. Separation and purification of phenolic acids and myricetin from black currant by high speed countercurrent chromatography. *J Liq Chromatogr Relat Technol*. 2009;32(20):3077-3088.

6. Sajitha M, Indhumathi S, Subramani K, Anbarasan PM. Chemical investigation and anti-microbial study of *Cleome gynandra*. *Nat Prod Indian J*. 2015;11:49-53.

7. Younghee P, Byoung HM, Heejung Y, Youngshim L, Eung-jung L, Yoongho L. Spectral assignments and reference data. Complete assignments of NMR data of 13 hydroxymethoxy-flavones. *Magn Reson Chem*. 2007;45(12):1072-1075.

8. Devkota HP, Yoshizaki K, Yahara S. Pilloon 5-O-β-D-glucopyranoside from the stems of *Diplomorpha ganpi*. *Biosci Biotechnol Biochem*. 2012;76(8):1555-1557.

9. Jang DS, Kim JM, Lee YM, et al. Flavonols from *Houttuynia cordata* with protein glycation and aldose reductase inhibitory activity. *Nat Prod Sci*. 2006;12:210-213.

10. Samy MN, Sugimoto S, Matsunami K, Otsuka H, Kamel MS. Bioactive compounds from the leaves of *Eugenia uniflora*. *J Nat Prod*. 2014;7:37-47.

11. Giner J-L, Feng J, Kiemle DJ. NMR tube degradation method for sugar analysis of glycosides. *J Nat Prod*. 2016;79(9):2413-2417.

12. Apostolidis E, Kwon Y-I, Shetty K. Inhibitory potential of herb, fruit, and fungal-enriched cheese against key enzymes linked to type 2 diabetes and hypertension. *Innov Food Sci Emerg Technol*. 2007;8(1):46-54.