Anti-influenza a virus activity of a new dihydrochalcone diglycosides isolated from the Egyptian seagrass Thalassodendron ciliatum (Forsk.) den Hartog

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Abstract

One new dihydrochalcone diglycosides has been isolated from the EtOAc fraction of the Egyptian seagrass *Thalassodendron ciliatum* (Forsk.) den Hartog, and identified as 6'-O-
rhamnosyl-(1'''→6'')-glucopyranosyl asebogenin for which a trivial name Thalassodendrone was established. Furthermore, five known phenolics were isolated and identified as asebotin, quercetin 3,7-diglucoside, protocatechuic acid, ferulic acid and p-hydroxybenzoic acid. The structures of all isolated compounds were established based on 1D and 2D NMR spectroscopy and HR-Mass spectrometer. The anti-influenza A virus activity of the isolated new compound and asebotin were evaluated, and the obtained results revealed an inhibition dose concentration of asebotin more than Thalassodendrone with IC$_{50}$ = 2.00 & 1.96 µg/mL respectively, and with cytotoxic concentration (CC$_{50}$) of 3.36 & 3.14 µg/mL respectively.

Keywords: *Thalassodendron ciliatum* (Forsk.) den Hartog; Cymodoceaceae; Dihydrochalcones; Phenolics; Antiviral; SAR studies
1. Introduction

Marine natural products have attracted the attention of biologists and chemists all over the world for the last five decades. Several of the compounds isolated from marine sources exhibited significant biological activity, which is accounted for the ocean to be a source of potential drugs (Bhakuni and Rawat, 2005).

In the tropics, shallow coastal areas are characterised by the existence of extensive seagrass meadows. The seagrasses are found in different habitats such as lagoons behind coral reefs that fringe the coast, mangrove bays or estuaries. The seagrass beds provide food and shelter for a variety of other organisms, including commercially important fish species, and thus constitute a valuable component of the nearshore ecosystem (Howard et al., 1989).

*Thalassodendron ciliatum* (Forsk.) den Hartog is commonly known as “Majani kumbi”, it is a very common seagrass species in the Red-Sea, the Western Indian Ocean and the tropical part of the Indo-Pacific region (Den Hartog, 1970).

*T. ciliatum* is used as “mutfusho” and as a treatment to relieve smallpox. A mixture of “short seagrasses”, such as *Thalassia* and *Cymodocea*, is effective against fever and skin diseases. Traditional uses, besides pure medicinal use, people in Chwaka (East Coast of Zanzibar) believe in the power of seagrasses to solve different human problems. Traditional doctors reported seagrasses as an important ingredient for different kinds of magic potions. Moreover, many respondents (45%) reported the use of seagrass beach cast as fertilizer in the cultivated family plots “shambas”. People reported that seagrasses are especially good for the growth of coconut trees (De La Torre-Castro and Rønnbøck, 2004). Seagrasses have showed great ability to accumulate several elements, and are increasingly used as a biological indicator of environmental quality (Pergent-Martini and Pergent, 2000; Pergent-Martini et al., 2005).

The aims of the presented study were to review the phytochemical constituents of the Egyptian seagrass *T. ciliatum*, searching for the presence of dihydrochalcones and other phenolics, accompanied by an evaluation of the antiviral activity of the new isolates, and then highlights their potentials as candidates for new drugs that may be of value in the treatment and prevention of human and livestock diseases. Extensive chromatographic separation and purification resulted in the isolation of one new dihydrochalcone diglycosides identified as 6′′-*O*-rhamnosyl-(1′′′→6′′)-glucopyranosyl asebogenin (compound 1), together with asebotin, quer cetin 3,7-dig lucoside, protocatechuic acid, ferulic acid and *p*-hydroxybenzoic acid.
The anti-influenza A virus activity of the isolated asebotin and the new compound (1) was evaluated and resulted with IC$_{50} = 2.00 \text{ & } 1.96 \mu \text{g/mL}$, respectively.

2. Results and Discussion

The phytochemical investigation of the EtOAc fraction of the MeOH extract of T. ciliatum resulted in the isolation of one new compound (1), together with asebotin (Guang-Min et al., 2005), quercetin 3,7-diglucoside (Mabry et al., 1970), protocatechuic acid (Achenbach et al., 1988), ferulic acid (Zhilan et al., 2006) and $p$-hydroxybenzoic acid (Takeo et al., 2004), which have been structurally elucidated using 1D, 2D-NMR, HR-MS and by comparison with the literature.

Compound (1) was isolated as a yellow amorphous solid, with the molecular formula C$_{38}$H$_{36}$O$_{14}$ as determined by a combination of high resolution MALDI-MS (S6, +ve) ($m/z$ 598.22561 for [M + 2H]$^+$ calc 598.22616 for C$_{38}$H$_{38}$O$_{14}$) together with the $^{13}$C-NMR and HSQC spectra (Experimental Section, S2 & S5) which confirmed the presence of 2 methyl, 3 methylene, 16 methine, and 7 quaternary carbons. The $^1$H-NMR and $^1$H-$^1$H COSY spectra (S1 & S3) of compound (1) suggested the presence of aseboginin (Harborne & Baxter, 1999) and/or asebotin (Nkengfack et al., 2001) derivative, with the characteristic signals as follow; two triplet signals correlate through a cross-peak and appeared at $\delta_H$ 2.88 (2H, $\delta$, $J = 7.5$ Hz, CH$_2$-p) and at $\delta_H$ 3.46 (2H, $\delta$, $J = 7.5$ Hz, CH$_2$-q), which are characteristic for the dihydrochalcones (Hufford and Oguntimewin, 1982), two ortho-coupled doublets correlated together and appeared at $\delta_H$ 7.06 (2H, $d$, $J = 8$ Hz, H-2 & H-6) and at $\delta_H$ 6.68 (2H, $d$, $J = 8$ Hz, H-3 & H-5) which are characteristic for the (AA'BB') system of the B-ring and indicating an oxygen substitution at the 4-position, another correlated meta-coupled doublets appeared at $\delta_H$ 6.29 (1H, $d$, $J = 2$ Hz, H-3') and at $\delta_H$ 6.31 (1H, $d$, $J = 2$ Hz, H-5'). Furthermore, a sharp singlet signal typical for an aromatic methyl group appeared at $\delta_H$ 3.83 (3H, s, H$_3$CO-4') & at $\delta_C$ 56.24 as confirmed from $^{13}$C-NMR and HSQC spectra (S2 & S5). Moreover, a doublet signal appeared at $\delta_H$ 5.02 (1H, $d$, $J = 7$ Hz, H-1') characteristic for the anomeric proton of a hexose moiety identified to be glucose as shown in $^{13}$C-NMR spectrum (S2), and another doublet signal appeared at $\delta_H$ 4.68 (1H, $d$, $J = 1$ Hz, H-1'') characteristic for the presence of rhamnose moiety, which confirmed by the presence of a doublet methyl appeared at $\delta_H$ 1.20 (3H, $d$, $J = 6$ Hz, H$_3$C-6'') and at $\delta_C$ 17.92. A number of multiplets appeared between $\delta_H$ 3.31–4.03 (10H, $m$) and correlated through a number of contiguous cross-peaks in COSY (S3), which are corresponding to the remaining sugars protons. The HMBC experiment (S4, S8) was conducted to get the attachment positions as
well as the inter-linkage of the methoxy group, glucose and rhamnose moieties through the long rang correlations (\(^2J \& \(^3J\)), the obtained results revealed the presence of three distinguishable cross peaks; the methoxy group at \(\delta_{31} 3.83\) with C-4′ of A-ring at \(\delta_{c} 167.66\), the anomic proton of glucose at \(\delta_{31} 5.02\) with C-6′ of A-ring at \(\delta_{c} 161.70\), and the anomic proton of rhamnose at \(\delta_{31} 4.68\) with the methylene hydroxy (\(\text{CH}_2\text{OH}\)) of glucose at \(\delta_{c} 67.84\), confirming \((1′′′ → 6′′′)\) glucorhamnopyranoside. Thus, the structure of compound (1) was elucidated as depicted in (Figure 1) and assigned to 6′-O-rhamnosyl-(1′′′ → 6′′′)-glucopyranosyl aseboigenin trivially named as Thalassodendrone.

![Figure 1. 6′-O-rhamnosyl-(1′′′ → 6′′′)-glucopyranosyl aseboigenin (Thalassodendrone)](image-url)

Infectious viral diseases remain a worldwide problem. Viruses have been resistant to therapy or prophylaxis longer than any other form of life due to their nature because they totally depend on the cells they infect for their multiplication and survival. Currently, there are only few drugs available for the cure of viral diseases including acyclovir, the known antitherpetic drug which is modeled on a natural product parent. In order to combat viruses which have devastating effects on humans, animals, insects, crop plants, fungi and bacteria, many research efforts have been devoted for the discovery of new antiviral natural products. So, as part of our ongoing collaborative effort to discover potential anti-viral agents from natural sources, the isolated new dihydrochalones were tested for their anti-virus activity against influenza A virus using MTT method, and revealed an inhibition dose concentration of asborbin more than compound (1) with IC\(_{50}\) = 2.00 & 1.96 \(\mu\)g/mL respectively, and with cytotoxic concentration (CC\(_{50}\)) of 3.36 & 3.14 \(\mu\)g/mL respectively.

Quantitative structure-activity relationship (QSAR) models are useful in providing a biochemical understanding of the biological activity of natural and synthetic chemicals based solely on molecular structure. Both of the aromatic substituents on both A- & B-rings and the keto-enol functionality of the phenolics can serve as targets for future structure activity
relationship (SAR) studies (Wu et al., 2003). The substituents at the 4-position of the phenyl ring B should have electron-donating properties and most probably this part of the phenolic molecule interacts with the catalytic domain of the enzyme through hydrogen bonds. However, larger substituents in position-4 other than OH are not favourable (Alenka et al., 2002).

It is concluded that the anti-viral inhibitory properties of chalcones and flavonoids are mainly the outcome of electronic interactions between atomic charges within these compounds in both A and B rings and possible receptor-like structures in the cells and prevent the virus attach and penetration to the cell. These agonist-receptor interactions are enhanced by hydrogen bonding contributions and by specific geometrical arrangements associated with each compound.

3. Experimental

3.1. Plant material

Seagrass samples of *Thalassodendron ciliatum* (Forssk.) den Hartog were collected from Magawish city near Hurghada, Egypt in October 2008, and were identified by Prof. Dr. Monir Abd-El Ghaney, Botany Department, Herbarium, Faculty of Science, Cairo University, Cairo, Egypt. A voucher specimen (SAA-41) was deposited in the herbarium section of Pharmacognosy Department, Faculty of Pharmacy, Suez Canal University, Ismailia, Egypt.

3.2. Extraction and isolation

Fresh *T. ciliatum* (800 g) was blended in an electric blender with MeOH, the process was repeated until complete extraction. The resulting extracts were combined, filtered, and the solvent was evaporated under reduced pressure at 45 °C to afford a crude MeOH ext., which is then partitioned between EtOAc and H₂O several times, to afford ethyl acetate fraction (10.61 g). Then the EtOAc fraction was chromatographed on a Sephadex LH-20 column (600 mm) with step gradient elution starting from 30% ethanol in H₂O to 100% ethanol. Fractions of 250 mL each were collected and those exhibiting similar TLC profiles were combined together. Five sub-fractions (I–V) were obtained and subsequently fractionated on Sephadex LH-20 column; fraction I was eluted with sat. BuOH to afford five sub-fractions (I₃) of which sub-fraction (I₃) purified on preparative TLC (MeOH-CHCl₃ 20:80, v/v) gave compound-I (2 mg). Fraction II eluted with 100% EtOH gave two sub-fractions (II₁,₂) on purification on Sephadex LH-20 with 2% EtOH in H₂O resulted in the isolation of asebotin (6 mg). Fraction III eluted with 100% EtOH revealed the isolation of quercetin 3,7-diglucoside (4 mg) and protocatechuic acid (2 mg). Fraction IV eluted with 100% MeOH resulted in the
isolation of fenolic acid (17.5 mg). Fraction V eluted with sat. BuOH to afford p-
hydroxybenzoic acid (4 mg).

3.2.1. 6'-O-rhamnosyl-(1''''→6'')-glucopyranosyl asebogenin (I)

Yellow amorphous solid. $^1$H-NMR (CD$_2$OD, 500 MHz) $^{\delta}$H 6.29 (1H, d, $J$ = 2 Hz, H-3'), $^{\delta}$H 6.31 (1H, d, $J$ = 2 Hz, H-5'), $^{\delta}$H 7.06 (2H, d, $J$ = 8 Hz, H-2 & H-6), $^{\delta}$H 6.68 (2H, d, $J$ = 8 Hz, H-3 & H-5), $^{\delta}$H 2.88 (2H, t, $J$ = 7.5 Hz, CH$_2$-β), $^{\delta}$H 3.46 (2H, t, $J$ = 7.5 Hz, CH$_2$-α), $^{\delta}$H 3.83 (3H, s, H$_3$CO-4'), $^{\delta}$H 5.02 (1H, d, $J$ = 7 Hz, H-1''), $^{\delta}$H 3.57 & $^{\delta}$H 4.03 (2H, brd, $J$ = 10 Hz, CH$_2$-6'''), $^{\delta}$H 1.20 (3H, d, $J$ = 6 Hz, H$_3$C-6'''), $^{\delta}$H 3.31–3.68 (8H, m, CH-2'', 3'', 4'', 5'', 2'''', 3'''', 4'''', 5'''''); $^{13}$C-NMR $^{\delta}$C 107.55 (C-1', C), $^{\delta}$C 167.32 (C-2', C), $^{\delta}$C 95.95 (C-3', CH), $^{\delta}$C 167.66 (C-4', C), $^{\delta}$C 69.13 (C-5', CH), $^{\delta}$C 161.70 (C-6', C), $^{\delta}$C 133.75 (C-1, C), $^{\delta}$C 130.41 (C-2 & -6, CH), $^{\delta}$C 116.10 (C-3 & -5, CH), $^{\delta}$C 156.40 (C-4, C), $^{\delta}$C 29.40 (CH$_2$-β), $^{\delta}$C 47.10 (CH$_2$-α), $^{\delta}$C 206.98 (C-O), $^{\delta}$C 56.24 (H$_3$CO-4'), $^{\delta}$C 102.27 (C-1'', C'), $^{\delta}$C 74.77 (C-2'', CH), $^{\delta}$C 77.26 (C-3'', CH), $^{\delta}$C 71.43 (C-4'', CH), $^{\delta}$C 78.59 (C-5'', CH), $^{\delta}$C 67.84 (CH$_2$-6''), $^{\delta}$C 102.19 (C-1'''', CH), $^{\delta}$C 72.40 (C-2'''', CH), $^{\delta}$C 73.52 (C-3'''', CH), $^{\delta}$C 74.11 (C-4'''', CH), $^{\delta}$C 69.83 (C-5'''', CH), $^{\delta}$C 17.92 (H$_3$C-6'''); HMBC and COSY correlations (S8), MALDI-MS (m/z 598.22561 for [M + 2H]$^+$ calc 598.22616 for C$_{28}$H$_{38}$O$_{14}$).

3.3. General experimental procedures

NMR: 1D-spectra were obtained using a pulse sequence supplied from JEOL JNM-
AL-400 MHz NMR spectrometer for $^{1}$H, $^{13}$C-NMR, DEPT-45, -90, -135 and $^{1}$H-$^{1}$H COSY) in DMSO-d$_6$. 2D-spectra (HSQC and HMBC) were obtained using a pulse sequence supplied from Varian Gemini VNMR-500 MHz NMR spectrometer. Chemical shifts were given in values (ppm) relative to trimethylsiline (TMS) as an internal reference. High-
resolution MALDI-MS: High resolution mass spectra were obtained on JEOL JMS-700N for
electron ionization or on JEOL JMS-T100 TD for electrospray ionization, using α-Cyano-4-
hydroxycinnamic acid (CHCA) as a matrix (m/z 189.17).

3.4. Assay for Antiviral Activity

3.4.1. Cells and viruses

Mardin-Darby canine kidney cells (MDCK) were grown in minimum essential
medium (MEM) supplemented with 5% fetal bovine serum (FBS) and 1%
penicillin/streptomycin (unless otherwise stated) at 37°C in a 5% CO$_2$ incubator. Original
virus solution: influenza virus A/WSN/33 (3.72×10$^7$ TCID$_{50}$/mL), with 100 TCID$_{50}$/well
infection.

3.4.2. Cytotoxic effect inhibition assay
MDCK cells were seeded (100 µL/well = 3.0×10⁴ cells/well) in 96-well plates and cultured in MEM/10% FBS for 2 days at 37°C to >90% confluence. 40 µL of the test sample (1mg sample dissolved in 1mL of DMSO) (4% solution, 2 fold dilutions) were added to 960 µL of MEM (-), and then from the mixture 120 µL/well were added to each well of 96-wells. Cells were then washed with PBS and infected with approximately 50 plaque forming units (PFU) of influenza virus (1000TCID₅₀/mL). 100 µL aliquot of the cell suspension was added to each well of a 96-well flat-bottomed microtitre tray containing 100 µL of various concentrations of the test sample. After 3-days incubation at 37°C in 5% CO₂, the number of viable cells was determined by the MTT method (Pauwels et al., 1988). The cytotoxicity of the each compound was evaluated in parallel with the antiviral activity, which was based on the viability of mock-infected cells, as monitored by the MTT method (See supplementary material S7). The 50% antiviral effective dose (EC₅₀) and the 50% cytotoxic dose (CC₅₀) of the sample were determined. The absorbances were determined with Tecan Infinite® 200 PRO Modular Microplate Readers at a test wavelength of 560 nm.

4. Conclusions

In the present study, the Egyptian seagrass T. ciliatum was phytochemically investigated for its secondary metabolites, which resulted with the isolation of one new dihydrochalcone diglycosides together with asebotin, flavonoid diglycosides, and three phenolic acids. The anti-influenza A virus activity was evaluated for the newly isolated dihydrochalcone diglycosides along with its mono glycoside derivative, which resulted with virus inhibition with IC₅₀ = 2.00 & 1.96 µg/mL respectively, and with cytotoxic concentration (CC₅₀) of 3.36 & 3.14 µg/mL respectively.

Supplementary material

Original NMR data (S1 – S5), high resolution MALDI/MS (S6) of compound (I), detailed protocol for anti-influenza A virus (S7), and some selected 2D correlations (S8) are available online.

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