


 Cite this: *RSC Adv.*, 2021, 11, 38703

Chemical synthesis, inhibitory activity and molecular mechanism of 1-deoxynojirimycin–chrysin as a potent α -glucosidase inhibitor†

 Ran Zhang,[‡] Yueyue Zhang,[‡] Gaiqun Huang,^{ac} Xiangdong Xin,^a Liumei Tang,^a Hao Li,^{ab} Kwang Sik Lee,^d Byung Rae Jin^d and Zhongzheng Gui^{‡*ab}

Hyperglycemia can be efficaciously regulated by inhibiting α -glucosidase activity and this is regarded as an effective strategy to treat type 2 diabetes. 1-Deoxynojirimycin, an α -glucosidase inhibitor, can penetrate cells rapidly to potently inhibit α -glucosidase in a competitive manner. However, the application of 1-deoxynojirimycin is limited by its poor lipophilicity and low bioavailability. Herein, three 1-deoxynojirimycin derivatives 4–6 were designed and synthesized by linking 1-deoxynojirimycin and chrysin to ameliorate the limitations of 1-deoxynojirimycin. Among them, compound **6**, a conjugate of 1-deoxynojirimycin and chrysin linked by an undecane chain, could better bind to the α -glucosidase catalytic site, thereby exhibiting excellent α -glucosidase inhibitory activity ($IC_{50} = 0.51 \pm 0.02 \mu\text{M}$). Kinetics analyses revealed that compound **6** inhibited the activity of α -glucosidase in a reversible and mixed competitive manner. Fluorescence quenching and UV-Vis spectra showed that compound **6** changed the conformation of the α -glucosidase *via* complex formation, which triggered a static fluorescence quenching of the enzyme protein.

 Received 20th October 2021
 Accepted 17th November 2021

DOI: 10.1039/d1ra07753h

rsc.li/rsc-advances

1. Introduction

Diabetes mellitus, one of the most common life-threatening illnesses worldwide, is a metabolic disorder that is characterized by persistent hyperglycemia.¹ Owing to unhealthy lifestyles, the number of people suffering from type 2 diabetes mellitus (T2DM) has increased dramatically globally. In recent years, various new and effective hypoglycemic drugs, such as α -glucosidase inhibitors, were developed for the treatment of this chronic disease. As known, α -glucosidase inhibitors can delay the release of D-glucose from dietary complex carbohydrates and retard glucose absorption in the small intestine by inhibiting α -glucosidase activity, resulting in reduced postprandial plasma glucose levels and suppression of postprandial hyperglycemia.^{2,3} Currently, the commercial α -glucosidase inhibitors, such as voglibose⁴ and acarbose,⁵ play a significant role in treating diabetes mellitus in clinics. However, these drugs are

limited by some adverse effects, including flatulence, diarrhea, vomiting, and abdominal pain.⁶ Therefore, there is an urgent need to develop novel potent α -glucosidase inhibitors with high efficiency and safety for T2DM patients. Natural products provide more effective choices for developing safe and efficient α -glucosidase inhibitors for T2DM.⁷

1-Deoxynojirimycin, a characteristic natural compound of mulberry leave, has various biological properties, including antidiabetic, anti-lipidemic, antimicrobial, and anticancer.^{8,9} 1-Deoxynojirimycin exerts its antidiabetic effects by inhibiting the activity of α -glucosidase, decreasing serum insulin and glucose and improving carbohydrate metabolism.^{10,11} However, there are some drawbacks such as low bioavailability and poor lipophilicity, making 1-deoxynojirimycin impossible to maintain an efficient and lasting hypoglycemic effect.¹² Therefore, various derivatives of 1-deoxynojirimycin were synthesized to ameliorate lipophilicity and to improve bioavailability. For example, 1-deoxynojirimycin–kaempferol exhibited excellent lipophilicity and better α -glucosidase inhibitory activity than 1-deoxynojirimycin.¹³ Hybrid of 1-deoxynojirimycin and quinazoline was regarded as a high active dual inhibitor of epidermal growth factor receptor (EGFR) and α -glucosidase.¹⁴

Chrysin, a natural occurring flavonoid, is present in various plants and is also abundant in honey and propolis.¹⁵ It has been reported that chrysin exerts a wide range of physiological effects, such as antidiabetic, anticancer, antioxidant, anti-inflammatory, and antihypertensive properties.^{16,17} Due to its excellent antioxidant effect and anti-dyslipidemia activities,

^aSchool of Biotechnology, Jiangsu University of Science and Technology, Zhenjiang 212100, Jiangsu, People's Republic of China. E-mail: srizzgui@hotmail.com

^bSericultural Research Institute, Chinese Academy of Agricultural Sciences, Zhenjiang 212100, Jiangsu, People's Republic of China

^cSericultural Research Institute, Sichuan Academy of Agricultural Sciences, Nanchong 637000, Sichuan, People's Republic of China

^dCollege of Natural Resources and Life Science, Dong-A University, Busan 49315, Republic of Korea

† Electronic supplementary information (ESI) available. See DOI: 10.1039/d1ra07753h

‡ These authors equally contributed to this work.



chrysin has been regarded as an antidiabetic agent with cardiac and hepatic protective effects.¹⁸ This antidiabetic action might be associated with the suppression of 11 β -hydroxysteroid dehydrogenase type I, lowering cortisol production and enhancing insulin sensitivity.¹⁹ Additionally, chrysin and its derivatives exhibit α -glucosidase inhibition activity and have a high potential for the treatment of T2DM.^{20–23} Nevertheless, chrysin is seriously restricted in application owing to its poor absorption in intestinal and the rapid metabolism of glycosylation.²⁰ A common way for designing novel chrysin derivatives is that small molecules are attached to the C-7-OH of chrysin *via* alkane chain, offering an effective and practical strategy for overcoming the above problem.^{24–26}

Therefore, three 1-deoxynojirimycin derivatives 4–6 were designed and synthesized, in which the C-7-OH of chrysin was linked to the amino group of 1-deoxynojirimycin by an alkyl chain linker, aiming at improving the lipophilicity of 1-deoxynojirimycin. Furthermore, because chrysin has an effect on inhibiting the activity of α -glucosidase, we investigated whether the introduction of chrysin would improve the inhibitory activity of 1-deoxynojirimycin against α -glucosidase. Finally, the molecular mechanism of 1-deoxynojirimycin–chrysin inhibition of α -glucosidase activity was studied.

2. Experimental section

2.1 Materials and methods

1-Deoxynojirimycin, 1,5-dibromopentane, 1,8-dibromooctane and 1,11-dibromoundecane were acquired from Energy-Chemical Co., Ltd (Shanghai, China). Chrysin was obtained from Tianjin Heowns Biochem LLC. 4-Nitrophenyl- α -D-glucopyranoside (pNPG), and α -glucosidase (*Saccharomyces cerevisiae*, EC 3.2.1.20) were purchased from Solarbio. All other reagents were analytically pure unless otherwise specified. The ¹H-NMR and ¹³C-NMR of all intermediates and products were recorded in Bruker spectrometer. The spectrum data were anatomized by the MestReNova software. The Bruker SolanX 70 FT-MS instrument was used to determine the HR-MS of compounds 4–6. The UV and fluorescence data were recorded by Infinite F50 (TECAN UV-Vis spectrophotometer) and F-7000 FL fluorescence spectrophotometer (Hitachi Scientific Co., Japan), respectively.

2.2 General procedure for synthesis of 1–3

To the solution of chrysin (2.36 mmol) in acetone, K₂CO₃ (4.72 mmol) and 1,5-dibromopentane or 1,8-dibromooctane or 1,11-dibromoundecane (11.80 mmol) were then added. The mixture was stirred at 60 °C overnight and monitored by thin layer chromatography (TLC). After the completion of the reaction, the crude compound was obtained by vacuum concentration. Then the desired compounds 1–3 as a pale-yellow solid were obtained by silica gel column chromatography.

7-((5-Bromoethyl)oxy)-5-hydroxy-2-phenyl-4H-chromen-4-one (1). Yield: 89%. ¹H NMR (400 MHz, DMSO-*d*₆): δ 12.81 (s, 1H), 8.11 (d, *J* = 6.80 Hz, 2H), 7.65–7.57 (m, 3H), 7.06 (s, 1H), 6.82 (d, *J* = 2.00 Hz, 1H), 6.38 (d, *J* = 2.00 Hz, 1H), 4.11 (t, *J* = 6.40 Hz,

2H), 3.58 (t, *J* = 6.40 Hz, 2H), 1.92–1.85 (m, 2H), 1.81–1.74 (m, 2H), 1.58–1.51 (m, 2H). ¹³C NMR (101 MHz, DMSO-*d*₆): δ 182.05, 164.68, 163.38, 161.11, 157.34, 132.14, 130.56, 129.12, 126.42, 105.29, 104.83, 98.45, 93.18, 68.29, 35.06, 31.82, 27.46, 24.12.

7-((8-Bromoethyl)oxy)-5-hydroxy-2-phenyl-4H-chromen-4-one (2). Yield: 87%. ¹H NMR (400 MHz, DMSO-*d*₆): δ 12.81 (s, 1H), 8.11 (d, *J* = 6.80 Hz, 2H), 7.66–7.58 (m, 3H), 7.06 (s, 1H), 6.83 (d, *J* = 2.00 Hz, 1H), 6.39 (d, *J* = 2.00 Hz, 1H), 4.11 (t, *J* = 6.80 Hz, 2H), 3.53 (t, *J* = 6.80 Hz, 2H), 1.84–1.71 (m, 4H), 1.41–1.23 (m, 8H). ¹³C NMR (101 MHz, DMSO-*d*₆): δ 182.06, 164.77, 163.41, 161.12, 157.37, 132.15, 130.58, 129.13, 126.44, 105.31, 104.83, 98.48, 93.18, 68.46, 35.22, 32.18, 28.48, 28.28, 28.00, 27.43, 25.24.

7-((11-Bromoundecyl)oxy)-5-hydroxy-2-phenyl-4H-chromen-4-one (3). Yield: 89%. ¹H NMR (600 MHz, CDCl₃): δ 12.72 (s, 1H), 7.90 (d, *J* = 7.20 Hz, 2H), 7.58–7.53 (m, 3H), 6.68 (s, 1H), 6.51 (d, *J* = 1.80 Hz, 1H), 6.38 (d, *J* = 1.80 Hz, 1H), 4.05 (t, *J* = 6.60 Hz, 2H), 3.43 (t, *J* = 7.20 Hz, 2H), 1.90–1.81 (m, 4H), 1.51–1.43 (m, 4H), 1.39–1.32 (m, 10H). ¹³C NMR (150 MHz, CDCl₃): δ 182.46, 165.22, 163.91, 162.15, 157.81, 131.80, 131.40, 129.09, 126.28, 105.87, 105.59, 98.62, 93.11, 68.70, 34.06, 32.83, 29.49, 29.44, 29.41, 29.29, 28.95, 28.76, 28.17, 25.93.

2.3 General procedure for synthesis of 4–6

To a solution of 1 or 2 or 3 (0.74 mmol) in anhydrous dimethyl formamide (DMF) were added K₂CO₃ (1.49 mmol) and 1-deoxynojirimycin (0.68 mmol). The reaction was stirred at 80 °C overnight. Then, the solvent was removed by vacuum filtration and the crude compounds were purified by silica gel column chromatography to obtain the desired compounds 4–6 as a pale-yellow solid.

5-Hydroxy-2-phenyl-7-((5-((2R,3R,4R,5S)-3,4,5-trihydroxy-2-(hydroxymethyl)-piperidin-1-yl)pentyl)oxy)-4H-chromen-4-one (4). Yield: 37%. Purity: 97%. ¹H NMR (400 MHz, DMSO-*d*₆): δ 8.11 (d, *J* = 6.80 Hz, 1H), 7.73–7.67 (m, 2H), 7.64–7.58 (m, 2H), 7.06 (s, 1H), 6.83 (d, *J* = 2.00 Hz, 1H), 6.40 (d, *J* = 2.40 Hz, 1H), 4.15–4.13 (m, 2H), 4.12 (s, 1H), 4.02 (d, *J* = 6.40 Hz, 1H), 3.78 (s, 2H), 3.51–3.44 (m, 1H), 3.27 (t, *J* = 8.80 Hz, 1H), 3.11–3.04 (m, 2H), 1.78 (t, *J* = 7.60 Hz, 2H), 1.62 (t, *J* = 6.00 Hz, 2H), 1.35 (t, *J* = 7.60 Hz, 2H), 1.23 (s, 6H). ¹³C NMR (101 MHz, DMSO-*d*₆): δ 182.04, 166.95, 164.71, 161.13, 157.36, 132.15, 131.66, 131.57, 130.56, 129.13, 128.63, 126.42, 105.32, 104.84, 98.47, 93.16, 77.02, 72.69, 71.11, 68.31, 67.35, 28.32, 23.19, 22.37, 13.87, 10.76. HRMS (ESI): *m/z* calculated for C₂₆H₃₁NO₈ [M + H]⁺: 486.2128, found 486.2129.

5-Hydroxy-2-phenyl-7-((8-((2R,3R,4R,5S)-3,4,5-trihydroxy-2-(hydroxymethyl)-piperidin-1-yl)octyl)oxy)-4H-chromen-4-one (5). Yield: 36%. Purity: 97%. ¹H NMR (600 MHz, DMSO-*d*₆): δ 12.79 (s, 1H), 8.09 (d, *J* = 7.20 Hz, 2H), 7.63–7.57 (m, 3H), 7.02 (s, 1H), 6.80 (s, 1H), 6.37 (s, 1H), 4.71 (s, 2H), 4.34 (t, *J* = 8.40 Hz, 1H), 4.22 (t, *J* = 6.60 Hz, 1H), 4.11–4.07 (m, 2H), 3.72 (d, *J* = 11.40 Hz, 1H), 3.59–3.62 (m, 1H), 3.48–3.45 (m, 1H), 3.11–3.04 (m, 3H), 2.93 (t, *J* = 9.00 Hz, 1H), 2.83–2.81 (m, 1H), 2.66 (t, *J* = 12.00 Hz, 1H), 1.96 (s, 2H), 1.73 (t, *J* = 7.20 Hz, 1H), 1.64 (t, *J* = 7.20 Hz, 1H), 1.40–1.35 (m, 6H), 1.31 (s, 2H), 1.22 (s, 2H). ¹³C NMR (150 MHz, DMSO-*d*₆): δ 182.51, 165.26, 163.90, 161.63,



157.85, 132.60, 131.08, 129.61, 126.91, 105.79, 105.32, 98.96, 93.62, 77.55, 73.21, 69.21, 68.98, 65.48, 57.49, 52.41, 45.19, 30.47, 28.75, 22.86, 19.11, 14.35, 14.01. HRMS (ESI): m/z calculated for $C_{29}H_{37}NO_8$ $[M + H]^+$: 528.2597, found 528.2593.

5-Hydroxy-2-phenyl-7-((11-((2*R*,3*R*,4*R*,5*S*)-3,4,5-trihydroxy-2-(hydroxymethyl)-piperidin-1-yl)undecyl)oxy)-4*H*-chromen-4-one (6). Yield: 39%. Purity: 99%. 1H NMR (600 MHz, DMSO- d_6): δ 12.79 (s, 1H), 8.09 (d, $J = 7.20$ Hz, 2H), 7.64–7.57 (m, 3H), 7.03 (s, 1H), 6.79 (d, $J = 2.40$ Hz, 1H), 6.36 (d, $J = 1.80$ Hz, 1H), 4.70–4.65 (m, 3H), 4.13 (s, 1H), 4.08 (d, $J = 6.60$ Hz, 2H), 3.71 (d, $J = 10.80$ Hz, 1H), 3.54 (d, $J = 11.40$ Hz, 1H), 3.20 (t, $J = 9.00$ Hz, 2H), 3.05–3.03 (m, 1H), 2.92 (t, $J = 8.40$ Hz, 1H), 2.81–2.78 (m, 1H), 2.75–2.69 (m, 1H), 2.38–2.34 (m, 1H), 1.96–1.91 (m, 2H), 1.75–1.70 (m, 2H), 1.41–1.22 (m, 16H). ^{13}C NMR (150 MHz, DMSO- d_6): δ 182.51, 165.26, 163.89, 161.63, 157.86, 132.61, 131.07, 129.61, 126.90, 105.78, 105.31, 98.95, 93.63, 79.65, 71.21, 69.86, 68.97, 67.13, 59.50, 57.34, 52.55, 29.56, 29.50, 29.18, 28.84, 27.49, 25.84, 24.88. HRMS (ESI): m/z calculated for $C_{32}H_{43}NO_8$ $[M + H]^+$: 570.3067, found 570.3069.

2.4 Calculation and measurement of log *P*

The log *P* values of 1-deoxyojirimycin and its derivatives 4–6 were predicted and tested by Molinspiration software (<http://www.molinspiration.com/>) and traditional octanol–water shake-flask method, respectively. Firstly, 1-deoxyojirimycin and its derivatives were dissolved in *n*-octanol and phosphate-buffered saline (PBS) and the absorbance were measured by UV spectrophotometer to establish a standard curve. Secondly, *n*-octanol/water mutual saturation was prepared with a mild mechanical stirring overnight, then separated. The compounds were dissolved in PBS-saturated *n*-octanol, and its solution was prepared to an equivalent volume of *n*-octanol-saturated PBS. The two-phase mixtures were shaken for 3 h. After centrifugation and separation, the concentration of compounds in each phase was determined by UV spectrometer. The concentrations of the compounds in PBS (C_w) and *n*-octanol (C_o) were obtained from the standard curves. The log *P* values were calculated according to the following formula: $\log P = \log C_o/C_w$. All experiments were repeated independently three times.

2.5 Assay of α -glucosidase inhibitory activity

The inhibitory effects of inhibitors (1-deoxyojirimycin, and its derivatives 4–6) on α -glucosidase were determined according to the previous method²⁷ using a TECAN UV-Vis spectrophotometer. Briefly, inhibitors at various concentrations (50 μ L) were preincubated with α -glucosidase (0.02 mg mL⁻¹, 100 μ L) in pH 6.8, 0.1 mol L⁻¹ PBS at 37 °C for 5 min, then pNPG (1 mM, 40 μ L) was added. The mixture was incubated at 37 °C for 30 min, and terminated by addition Na₂CO₃ (100 mM, 60 μ L) to final volume with 250 μ L. Absorption at 405 nm was measured to evaluate the inhibitory activity of the compounds to α -glucosidase. All experiments were repeated independently three times and the IC₅₀ value of each compound was obtained by SPSS 16.0.

2.6 Inhibitory kinetic analysis

The inhibitory mechanism of compound 6 against α -glucosidase was implemented by changing the concentrations of α -glucosidase using the same method as described above. The inhibition type was then assayed by the Lineweaver–Burk plot, and the inhibition constant was determined from the secondary plot. To describe mixed inhibition mechanism, secondary plot can be plotted according to the following equation:²⁸

$$\text{Slope} = K_m/V_{\max} + K_m[I]/K_i V_{\max} \quad (1)$$

$$Y\text{-Intercept} = 1/V_{\max}^{\text{app}} = 1/V_{\max} + [I]/K_{is} V_{\max} \quad (2)$$

where [I] and K_m indicated the concentrations of inhibitor and Michaelis–Menten constant, respectively. K_i and K_{is} were the dissociation constants of the inhibitor binding to the free enzyme and enzyme–substrate complex, respectively.

2.7 Fluorescence spectra measurements

The fluorescence intensity of the interaction between compound 6 and α -glucosidase were measured by F-7000 FL fluorescence spectrophotometer (Hitachi scientific Co, Japan) with an excitation wavelength at 280 nm, and fluorescence emission spectrum was recorded at the wavelength from 300 to 500 nm. Briefly, compound 6 at different concentrations (0–8.0 $\times 10^{-5}$ mol L⁻¹, 10 μ L) were added to α -glucosidase solution (1 U mL⁻¹, 2 mL), respectively. The mixture was incubated at 298 K for 30 s before the test. And fluorescence quenching was described by the following Stern–Volmer equation:²⁹

$$F_0/F = 1 + K_q\tau_0[Q] = 1 + K_{SV}[Q] \quad (3)$$

In the eqn (3), F_0 and F represent the fluorescence intensities in the absence and presence of the quencher, respectively. K_q and K_{SV} are the quenching rate constants of the bimolecular and the Stern–Volmer dynamic quenching constant, respectively. τ_0 is the average life-time of the biomolecule without the quencher ($\tau_0 = 6.2$ ns)³⁰ and [Q] is the concentration of the quencher. In addition, to further illuminate the internal driving force of tannic acid against α -glucosidase, thermodynamic parameters were calculated under different temperatures (273, 298, and 310 K) using the same method as described above.

2.8 UV-Vis absorption spectra

The absorption spectra of α -glucosidase (1.0 $\times 10^{-5}$ mol L⁻¹) mixed with compound 6 (0–9.0 $\times 10^{-5}$ mol L⁻¹) were recorded by Infinite F50 TECAN UV-visible spectrophotometer at wavelengths of 200–600 nm.

2.9 Molecular docking

Docking calculations were achieved using MOE on a α -glucosidase model (<http://www.rcsb.org/pdb/home/home.do>). On account of 85% similarity and 73% sequence identity between α -glucosidase and isomaltase, the protein structure of α -glucosidase (from *Saccharomyces cerevisiae*) was employed on the basis of the X-ray crystal structure of the isomaltase from



Saccharomyces cerevisiae (PDB ID: 3aj7).³¹ When preparing the structure of α -glucosidase, it is necessary to remove the water molecules and the original ligand, and add polar hydrogen atoms. The 3D structure of tannic acid was prepared by using ChemBioDraw Ultra 14.0. The 3D interaction plots of α -glucosidase and compounds are generated by Surflex-Dock in SYBYL2.1.1.

3. Results and discussion

3.1 Synthesis and structural characterization of compounds 1–6

Three 1-deoxynojirimycin derivatives 4–6 (Fig. 1) were synthesized in accordance with the reported procedures,^{24,32} and the synthetic routes were summarized in Scheme 1. Briefly, chrysin analogues (compounds 1–3) were obtained by stirring the mixture of chrysin and 1,5-dibromopentane or 1,8-dibromooctane or 1,11-dibromoundecane at 60 °C in the presence of K_2CO_3 . Then, the finally compounds 4–6 were synthesized by treating 1–3 with 1-deoxynojirimycin in dry DMF for 12 h and purified by column chromatography. Their structures were characterized by nuclear magnetic resonance (NMR) and high-

resolution mass spectrometry (HR-MS). The purity over 95% was analysed by high-performance liquid chromatography (HPLC) (all data are presented in Fig. S1–S18†).

3.2 Lipophilicity of compound 6 contributed to inhibiting α -glucosidase activity

Lipid–water partition coefficient ($\log P$), a physicochemical parameter of quantitative structure–activity relationship, has an impact on the pharmacokinetics characteristic of compounds.³³ Therefore, the theoretical $\log P$ values of 1-deoxynojirimycin and 4–6 were obtained by Molinspiration (Milog P), and the actual $\log P$ values ($clog P$) were measured by the traditional shake flask method. $\log P$ values between 0.5 and 3.0 have contributed to the absorption and distribution for a drug candidate.³⁴ As shown in Table 1, compound 6 exhibited the best lipophilicity with an experimental $\log P$ of 2.61 ± 0.06 among compounds 4–6, which might be related to the presence of chrysin and undecane chain.

Besides, it has been reported that 1-deoxynojirimycin and chrysin exhibit an α -glucosidase inhibition activity.^{12,21} Therefore, the inhibition activity of compound 4–6 against α -

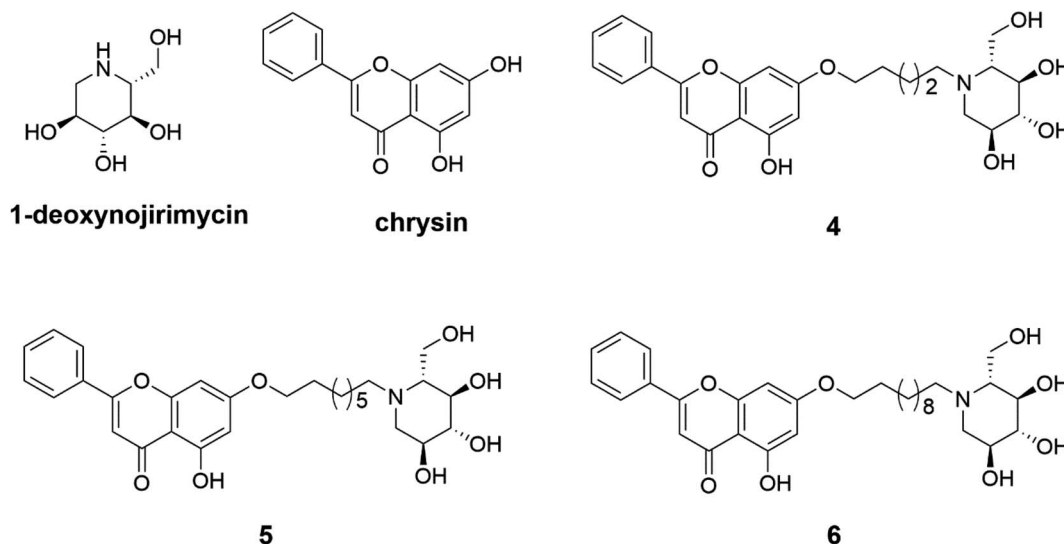
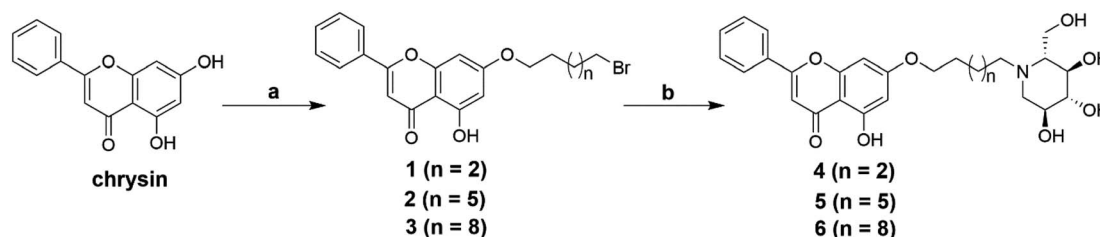


Fig. 1 Chemical structures of 1-deoxynojirimycin, chrysin and target compounds 4–6.



Condition and reagents: (a) 1,5-dibromopentane or 1,8-dibromooctane or 1,11-dibromoundecane, K_2CO_3 , 60 °C, overnight; (b) 1-deoxynojirimycin, K_2CO_3 , DMF, 80 °C, overnight

Scheme 1 Synthetic route for compounds 1–6.



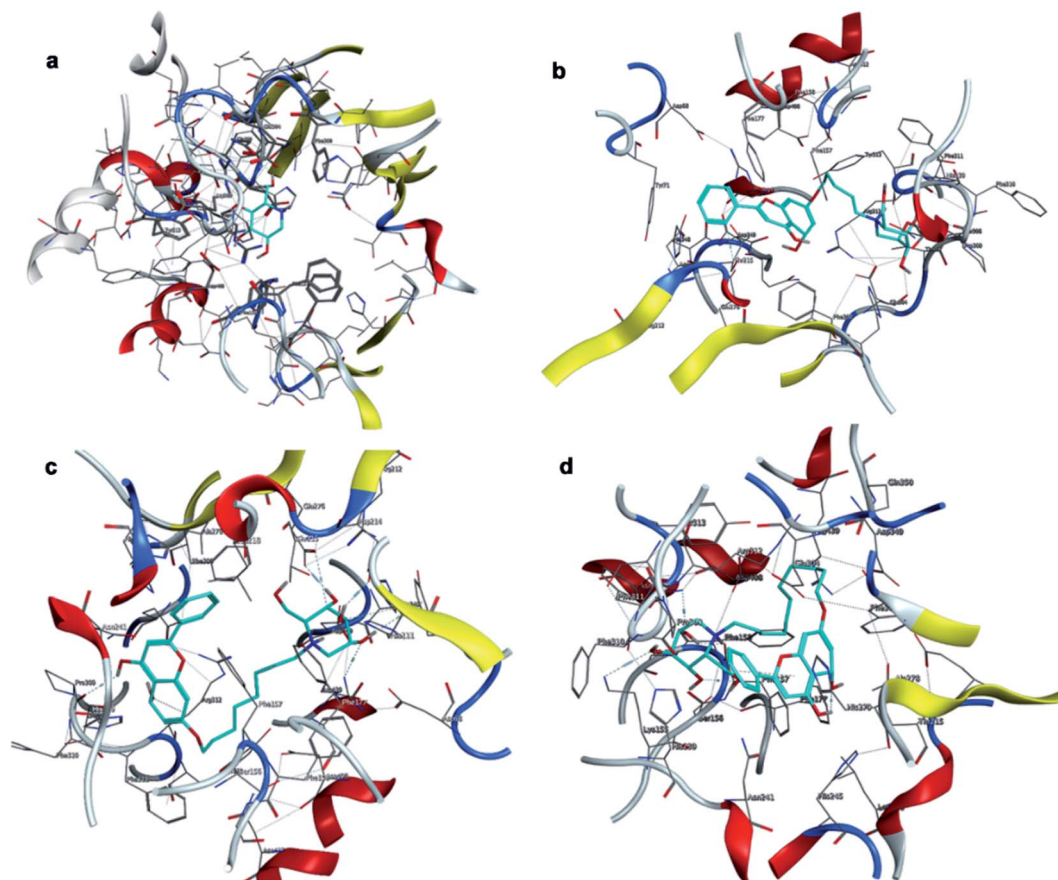
Table 1 Calculated and experimental log *P* values, and the α -glucosidase inhibitory IC₅₀ of 1-deoxynojirimycin compounds

Compound	Milog <i>P</i>	clog <i>P</i>	α -Glucosidase inhibitory IC ₅₀ (μ M)
1-Deoxynojirimycin	-2.40	-0.72 \pm 0.03	8.15 \pm 0.12
4	2.06	0.28 \pm 0.05	>100
5	3.57	1.35 \pm 0.07	5.56 \pm 0.24
6	5.09	2.61 \pm 0.06	0.51 \pm 0.02

glucosidase was assessed by using 1-deoxynojirimycin as a control. As shown in Table 1, all tasted compounds had a potent inhibitory effect on α -glucosidase, among which two 1-deoxynojirimycin derivative 5 and 6 had superior inhibition effects compared to 1-deoxynojirimycin. Especially, the α -glucosidase inhibitory activity of 6 with IC₅₀ values of 0.51 \pm 0.02 μ M was up to 16-fold higher than that of 1-deoxynojirimycin with IC₅₀ of 8.15 \pm 0.12 μ M, revealing that the introduction of chrysin could improve the inhibitory activity of 1-deoxynojirimycin against α -glucosidase. The result suggested that the lipophilicity of compound 6 contributed to enhancing its absorption and subsequent binding to the catalytic site of α -glucosidase, thereby exerting excellent α -glucosidase inhibitory activity.

3.3 Docking simulation between compound 4-6 and α -glucosidase

The affinity between 1-deoxynojirimycin derivatives 4-6 and α -glucosidase might offer more explanation for the above results of the α -glucosidase inhibitory activity. As we all known, 1-deoxynojirimycin needs to bind with the catalytic site of α -glucosidase, thereby exerting α -glucosidase inhibitory activity and lowering the blood glucose level.¹² Therefore, molecular docking was performed *via* Molecular Operating Environment (MOE) and SYBYL2.1.1 software to evaluate the binding ability of compounds 4-6 with α -glucosidase, respectively. Based on the crystal structure of α -glucosidase (PDB ID: 3aj7), we constructed the three-dimensional (3D) model of α -glucosidase by the SWISS-MODEL. 1-Deoxynojirimycin interacted with amino acid residues on α -glucosidase in the main way of hydrogen bonds. As shown in Fig. 2, docking results demonstrated that 1-deoxynojirimycin and its derivatives 4-6 could bind well to α -glucosidase. The docked conformations of 1-deoxynojirimycin its derivatives were shown by 2D interaction graph (Fig. 3). Compound 4 can form three hydrogen bonds, one π -H with some amino acid residues such as Asn241, Glu304 and Arg439, and compound 5 form three hydrogen bonds with Asn241, Glu304 and Gln360, but compound 6 form one hydrophobic bond with Ser308. In addition, compound 5 also interacts with Phe311 and Arg439, and compound 6 interacts with Phe311,

**Fig. 2** 3D interactions of 1-deoxynojirimycin, and 4-6 with α -glucosidase. (a) 3D image of 1-deoxynojirimycin, (b) 3D image of compound 4, (c) 3D image of compound 5, and (d) 3D image of compound 6.

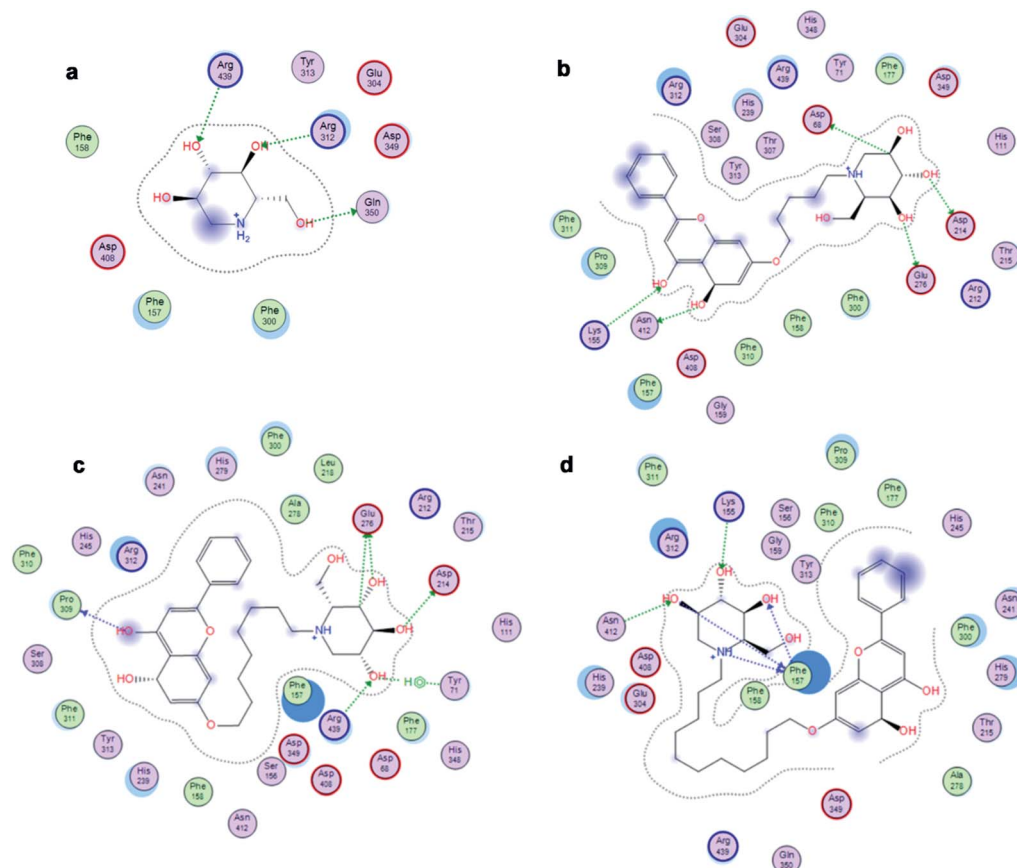


Fig. 3 2D images of the docked conformations of 1-deoxynojirimycin, 4–6 with α -glucosidase. (a) 2D image of 1-deoxynojirimycin, (b) 2D image of compound 4, (c) 2D image of compound 5, and (d) 2D image of compound 6.

His348, Arg439 and Glu304. 1-Deoxynojirimycin binds with α -glucosidase with these residues, suggesting that these residues play significant roles in the interaction between 4–6 and α -glucosidase. Furthermore, the docking score of compounds 4–6 is 6.9072, 8.4506 and 9.3964, further suggesting that compound 6 could be better to bind to the α -glucosidase catalytic site and exert the best α -glucosidase inhibitory activity. Therefore, compound 6 was selected as a representative to further investigate its inhibitory mechanism on α -glucosidase.

3.4 Compound 6 exerted a reversible and mixed-type inhibitory effect on α -glucosidase

It has been reported that 1-deoxynojirimycin could reversibly inhibit the activity of α -glucosidase.³⁵ In order to explore the inhibitory mode of compound 6 on α -glucosidase, enzyme reaction kinetics method was performed. As shown in Fig. 4a, three straight lines (with 0.0, 0.5, and 1.0 μ M of compound 6) passed through the origin. Therefore, the inhibition mechanism of the compound 6 on α -glucosidase was reversible.³⁶ Moreover, with increasing concentration of compound 6, the slope of the curves decreases, indicating that the presence of compound 6 did not reduce the content of α -glucosidase, but only led to a decrease in the activity of enzyme. The inhibition type was then assayed by the Lineweaver–Burk plot, and the

inhibition constant was determined from the secondary plot. As shown in Fig. 4b, a series of lines with different slopes and intersected in the second quadrant, indicating that compound 6 was a mixed-type inhibitor. In other words, compound 6 inhibited α -glucosidase activity not only by binding to free enzyme directly, but also by interfering with the formation of the α -glucosidase-PNPG intermediate through producing an α -glucosidase-PNPG-inhibitor complex in a non-competitive manner. According to eqn (1) and (2), the inhibition constants of compound 6 binding with the free enzyme (K_i) and with the enzyme–substrate complex (K_{is}) were determined to be 0.21 μ M and 0.76 μ M, respectively (Fig. 4c and d). The value of K_i was lower than that of K_{is} , suggesting that the affinity of compound 6 with free enzyme was stronger than that of the enzyme–substrate complex.³⁷

3.5 Fluorescence quenching mechanism of compound 6

To obtain further investigation of the interaction between compound 6 and α -glucosidase, fluorescence quenching experiments were performed. The emission spectra of α -glucosidase in the presence of 6 was recorded in the wavelength range 300–500 nm, by exciting the wavelength at 280 nm. As shown in Fig. 5a, α -glucosidase exhibited a maximum emission wavelength at approximately 337 nm, associated with the Trp



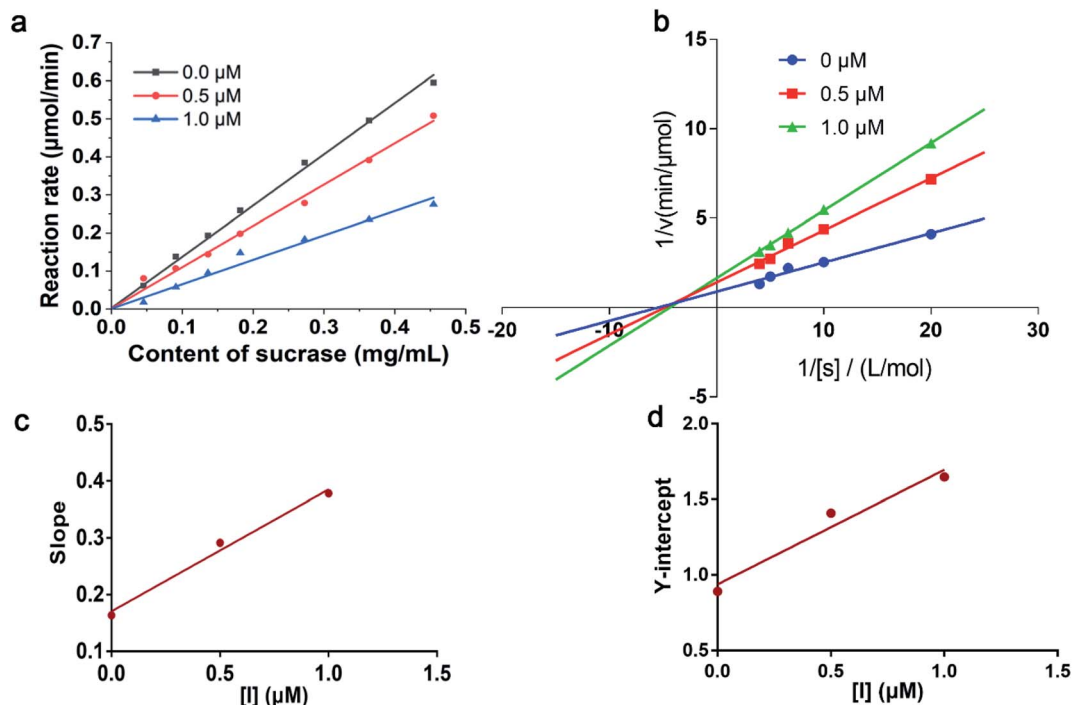


Fig. 4 (a) Inhibitory mechanism of compound 6 on α -glucosidase. (b) Lineweaver–Burk plot for kinetic analysis of α -glucosidase inhibition by compound 6. (c) Inhibition constant (K_i) of compound 6 on α -glucosidase. (d) Inhibition (K_{is}) constant of compound 6 on α -glucosidase–substrate complex.

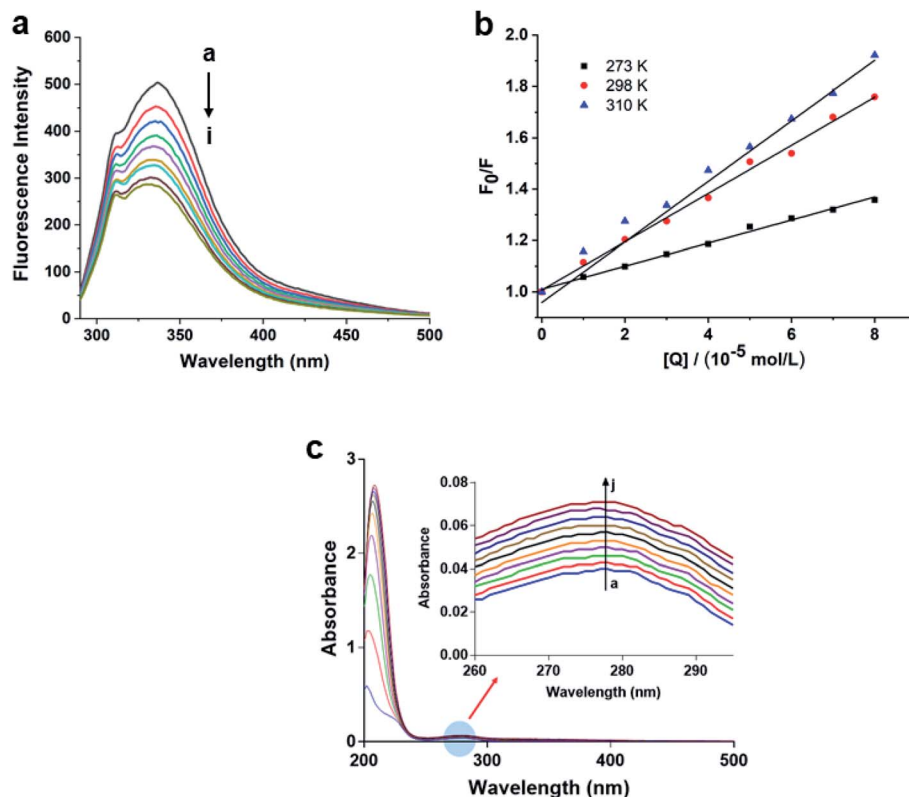


Fig. 5 (a) Fluorescence emission spectra of α -glucosidase in the presence of various concentrations of compound 6. The concentration of compound 6 increases from (a) to (i) ($0, 1, 2, 3, 4, 5, 6, 7, 8 \times 10^{-5} \text{ mol L}^{-1}$). (b) Simple Stern–Volmer plot of the quenching of α -glucosidase fluorescence by compound 6 at different temperatures (273 K, 298 K, and 310 K). (c) UV-Vis absorption spectra of α -glucosidase in the presence of various concentrations of compound 6. The concentration of compound 6 increases from (a) to (j) ($0, 1, 2, 3, 4, 5, 6, 7, 8, 9 \times 10^{-5} \text{ mol L}^{-1}$).



residues of α -glucosidase. However, with the addition of compound **6** to α -glucosidase solution, the fluorescence intensity of the enzyme reduced progressively, indicating that compound **6** can bind to α -glucosidase effectively, which was consistent with the results of molecular docking.

In addition, fluorescence quenching could occur by two mechanisms: dynamic quenching and static quenching. To shed light on the interaction mechanism, the Stern–Volmer equation (eqn (3)) was used to analyze the fluorescence data. From Fig. 5a, the quenching rate constant K_q of **6** was $3.70 \times 10^{11} \text{ L mol}^{-1} \text{ s}^{-1}$, which was much higher than the maximum value possible for diffusion controlled quenching ($2.0 \times 10^{10} \text{ L mol}^{-1} \text{ s}^{-1}$).³⁸ The result suggested that there is a specific interaction between compound **6** and α -glucosidase, and quenching mechanism may be not caused by dynamic quenching but by a static one. Furthermore, with the increasing of temperature, the values of K_q decrease dramatically for α -glucosidase, suggesting that the quenching process was not conducive at elevated temperature and the quenching process of α -glucosidase was an exothermic reaction (Fig. 5b).

3.6 UV-Vis absorption spectrum analysis of compound 6

UV absorption measurement is a very simple but effective method to exploring the structural change and understanding the complex formation. As we all known that for static quenching, the changes in the UV-Vis absorption spectrum are attributed to the formation of the complex.^{39,40} Therefore, the UV-Vis absorption spectra of α -glucosidase in the presence and absence of compound **6** at 298 K were recorded. As shown in Fig. 5c, the absorption intensity of α -glucosidase at 278 nm increased with the increasing concentrations of compound **6**, indicating that the fluorescence quenching of α -glucosidase is mainly caused by complex formation between α -glucosidase and compound **6**. The results further confirmed that the static quenching exists in the interaction of α -glucosidase and compound **6**.

4. Conclusions

In summary, three novel compounds **4–6**, derived from 1-deoxyojirimycin and chrysin, were designed and synthesized. Among them, compound **6** was the most efficient α -glucosidase inhibitor with IC_{50} value of $0.51 \mu\text{M}$. Molecular docking analysis manifested that compound **6** had the best binding conformation with α -glucosidase. Enzyme kinetic analysis indicated that compound **6** inhibited the activity of α -glucosidase in a reversible and mixed competition type. In addition, fluorescence quenching and UV-Vis spectra experiment confirmed that a complex formed from compound **6** and α -glucosidase, further triggering a static fluorescence quenching of the enzyme protein. Overall, this research provides a novel strategy for the development of potential α -glucosidase inhibitors. Further studies of its molecular mechanism in cell are currently underway.

Conflicts of interest

The authors declare no competing financial interests.

Acknowledgements

Financial support for this research was provided by the Jiangsu Province Policy Guidance Project (BX2019072), the Jiangsu Project of Science and Technology (XZ-SZ201925).

References

- J. Wojciechowska, W. Krajewski, M. Bolanowski, T. Kręcki and T. Zatoński, *Exp. Clin. Endocrinol. Diabetes*, 2016, **124**, 263–275.
- S. Kumar, S. Narwal, V. Kumar and O. Prakash, *Pharmacogn. Rev.*, 2011, **5**, 19–29.
- U. Hossain, A. K. Das, S. Ghosh and P. C. Sil, *Food Chem. Toxicol.*, 2020, **145**, 111738.
- P. Nowrouzi-Sohrabi, R. Tabrizi, S. Rezaei, F. Jafari, K. Hessami, M. Abedi, M. Jalali, P. Keshavarzi, S. Shahabi, A. A. Kolahi, K. Carson-Chahhoud, A. Sahebkar and S. Safiri, *Pharmacol. Res.*, 2020, **159**, 104988.
- U. F. Wehmeier and W. Piepersberg, *Appl. Microbiol. Biotechnol.*, 2004, **63**, 613–625.
- W. L. Zhang, W. M. Mu, H. Wu and Z. Q. Liang, *Appl. Microbiol. Biotechnol.*, 2019, **103**, 9335–9344.
- J. Rios, F. Francini and G. Schinella, *Planta Med.*, 2015, **81**, 975–994.
- Y. Liu, J. Zhang, H. Guo, A. Zhao, D. Shao, Z. Dong, Y. Sun, Y. Fan, F. Yang, P. Li, S. Mao, W. Zhong, Z. Ren, H. Wang, Y. Zhang and P. Wang, *J. Funct. Foods*, 2020, **73**, 104117.
- V. K. Ramappa, D. Srivastava, P. Singh, U. Kumar and V. Singh, *J. Horticult. Sci. Biotechnol.*, 2020, **95**, 1–8.
- H. X. Bai, W. Jiang, X. F. Wang, N. Hu, L. N. Liu, X. Li, Y. H. Xie and S. W. Wang, *Food Addit. Contam., Part A*, 2021, 1–13.
- E. Shuang, Y. Kazushi, S. Yu, M. Yui, L. Yui, K. Toshiyuki, N. Kiyotaka, M. Teruo and T. Tsuyoshi, *J. Clin. Biochem. Nutr.*, 2017, **61**, 47–52.
- H. Wang, Y. Shen, L. Zhao and Y. Ye, *Curr. Med. Chem.*, 2021, **28**, 628–643.
- R. Zhang, Y. Y. Zhang, X. D. Xin, G. Q. Huang, N. Zhang, Q. L. Zeng, L. M. Tang, T. Attaribo, K. S. Lee, B. R. Jin and Z. Z. Gui, *J. Nat. Prod.*, 2021, **84**, 1534–1543.
- Y. L. Zhang, H. L. Gao, R. J. Liu, J. Liu, L. Chen, X. B. Li, L. J. Zhao, W. Wang and B. L. Li, *Bioorg. Med. Chem. Lett.*, 2017, **27**, 4309–4313.
- E. Eldutar, F. M. Kandemir, S. Kucukler and C. Caglayan, *J. Biochem. Mol. Toxicol.*, 2017, **31**, e21960.
- E. R. Kasala, L. N. Bodduluru, R. M. Madana, K. V. Athira, R. Gogoi and C. C. Barua, *Toxicol. Lett.*, 2015, **233**, 214–225.
- R. Mani and V. Natesan, *Phytochemistry*, 2017, **145**, 187–196.
- S. Samarghandian, M. Azimi-Nezhad, F. Samini and T. Farkhondeh, *Can. J. Physiol. Pharmacol.*, 2016, **94**, 388–393.



- 19 M. Torres-Piedra, R. Ortiz-Andrade, R. Villalobos-Molina, N. Singh, J. L. Medina-Franco, S. P. Webster, M. Binnie, G. Navarrete-Vázquez and S. Estrada-Soto, *Eur. J. Med. Chem.*, 2010, **45**, 2606–2612.
- 20 D. El-Hussien, G. M. El-Zaafarany, M. Nasr and O. Sammour, *Int. J. Pharm.*, 2021, **592**, 120044.
- 21 H. Gao and J. Kawabata, *Bioorg. Med. Chem.*, 2005, **13**, 1661–1671.
- 22 J. S. Shin, K. S. Kim, M. B. Kim, J. H. Jeong and B. K. Kim, *Bioorg. Med. Chem. Lett.*, 1999, **9**, 869–874.
- 23 J. J. Ramírez-Espinosa, J. Saldaña-Ríos, S. García-Jiménez, R. Villalobos-Molina, G. Ávila-Villarreal, A. N. Rodríguez-Ocampo, G. Bernal-Fernández and S. Estrada-Soto, *Molecules*, 2017, **23**, 67.
- 24 K. Hu, W. Wang, H. Cheng, S. S. Pan and J. Ren, *Med. Chem. Res.*, 2011, **20**, 838–846.
- 25 H. Q. Li, L. Shi, Q. S. Li, P. G. Liu, Y. Luo, J. Zhao and H. L. Zhu, *Bioorg. Med. Chem.*, 2009, **17**, 6264–6269.
- 26 K. S. Babu, T. H. Babu, P. V. Srinivas, K. H. Kishore, U. S. N. Murthy and J. M. Rao, *Bioorg. Med. Chem. Lett.*, 2006, **16**, 221–224.
- 27 T. Ohta, S. Sasaki, T. Oohori, S. Yoshikawa and H. Kurihara, *Biosci., Biotechnol., Biochem.*, 2020, **66**, 1552–1554.
- 28 Q. Huang, W. M. Chai, Z. Y. Ma, C. Ou-Yang, Q. M. Wei, S. Song, Z. R. Zou and Y. Y. Peng, *Int. J. Biol. Macromol.*, 2019, **141**, 358–368.
- 29 J. Toneatto and G. A. Argüello, *J. Inorg. Biochem.*, 2011, **105**, 645–651.
- 30 Y. S. Yuan, L. Yang, S. P. Liu, J. D. Yang, H. Zhang, J. J. Yan and X. L. Hu, *Spectrochim. Acta, Part A*, 2017, **176**, 183–188.
- 31 G. C. Wang, M. Chen, J. Wang, Y. P. Peng, L. Y. Li, Z. Z. Xie, B. Deng, S. Chen and W. B. Li, *Bioorg. Med. Chem. Lett.*, 2017, **27**, 2957–2961.
- 32 Y. X. Zhao, Y. Zhou, K. M. O'Boyle and P. V. Murphy, *Bioorg. Med. Chem.*, 2008, **16**, 6333–6337.
- 33 M. M. Kla and I. Yildiz, *Ankara Univ. Vet. Fak. Derg.*, 2019, **43**, 1–19.
- 34 R. Odi, D. Bibi, T. Wager and M. Bialer, *Epilepsia*, 2020, **61**, 1543–1552.
- 35 A. Hatano, Y. Kanno, Y. Kondo, Y. Sunaga, H. Umezawa, M. Okada, H. Yamada, R. Iwaki, A. Kato and K. Fukui, *Bioorg. Med. Chem.*, 2017, **25**, 773–778.
- 36 M. Z. Lin, W. M. Chai, Y. L. Zheng, Q. Huang and C. Ou-Yang, *Int. J. Biol. Macromol.*, 2019, **122**, 1244–1252.
- 37 J. T. Zhang, L. J. Sun, Y. S. Dong, Z. X. Fang, T. Nisar, T. Zhao, Z. C. Wang and Y. R. Guo, *Food Chem.*, 2019, **299**, 125102.
- 38 P. Kandagal, S. Ashoka, J. Seetharamappa, V. Vani and S. Shaikh, *J. Photochem. Photobiol., A*, 2006, **179**, 161–166.
- 39 B. Zheng, M. Y. Li, G. Gao, Y. Y. He and P. J. Walsh, *Adv. Synth. Catal.*, 2016, **358**, 2156–2162.
- 40 D. Xu, Q. Y. Wang, T. Yang, J. Z. Cao, Q. L. Lin, Z. Q. Yuan and L. Li, *Int. J. Environ. Res. Public Health*, 2016, **13**, 334.

