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RESEARCH ARTICLE

Dual functional cholinesterase and MAO inhibitors for the treatment of Alzheimer's disease: synthesis, pharmacological analysis and molecular modeling of homoisoflavonoid derivatives

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Abstract

Because of the complexity of Alzheimer's disease (AD), the multi-target-directed ligand (MTDL) strategy is expected to provide superior effects for the treatment of AD, instead of the classic one-drug-one-target strategy. In this context, we focused on the design, synthesis and evaluation of homoisoflavonoid derivatives as dual acetyl cholinesterase (AChE) and monoamine oxidase (MAO-B) inhibitors. Among all the synthesized compounds, compound **10** provided a desired balance of AChE and hMAO-B inhibition activities, with IC₅₀ value of 3.94 and 3.44 μ M, respectively. Further studies revealed that compound **10** was a mixed-type inhibitor of AChE and an irreversible inhibitor of hMAO-B, which was also confirmed by molecular modeling studies. Taken together, the data indicated that **10** was a promising dual functional agent for the treatment of AD.

Keywords

Alzheimer's disease, cholinesterase inhibitors, homoisoflavonoid derivatives, MAO inhibitors

History

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Introduction

Alzheimer's disease (AD), characterized by memory loss and other cognitive impairments, is currently one of the most difficult and baffling diseases to treat^{1–3}. In the past decades, various pathogenesis hypothesis of AD have been proposed, such as cholinergic hypothesis^{4,5}, amyloid cascade hypothesis^{6–8}, oxidative stress hypothesis^{9–12}, tau protein hypothesis^{13,14} and so on. Among them, cholinergic hypothesis was a widely accepted theory, which suggests that the low level of ACh in specific regions of the brain is the major cause results in learning and memory dysfunctions. Based on the cholinergic hypothesis, one possible approach to treat AD is to restore the level of acetylcholine, such as using reversible inhibitors to inhibit acetyl cholinesterase (AChE). Up to now, AChE inhibitors, including tacrine (Cognex[®], Figure 1), donepezil (Aricept[®], Figure 1), rivastigmine (Exelon[®], Figure 1) and galanthamine (Reminyl[®], Figure 1) are approved by Food and Drug Administration to use in clinic, which could effectively improve memory in some AD patients^{15,16}. The crystal structures of AChE with inhibitors revealed that there are two binding sites, a peripheral anionic site (PAS) and a catalytic active site (CAS)^{17–19}. The PAS region of AChE, located at the rim of the gorge, was involved in the interaction with A β peptide, which may induce the deposition of neurotoxic A β fibrils^{20,21}. Recent studies showed that AChE inhibitors, blocking both the catalytic and peripheral sites simultaneously, might not only alleviate the cognitive defect

of AD patients by elevating ACh levels, but also act as disease-modifying agents delaying amyloid plaque formation^{22,23}. For example, donepezil, a mixed-type inhibitor, has a small effect on decreasing A β aggregation induced by AChE (about 22%), which can be attributed to its higher affinity for the active site than for the PAS²⁴.

Monoamine oxidases A and B (MAO-A and MAO-B), which could catalyze the oxidative deamination of neuroactive and vasoactive amines, are thought to play an important role in the metabolism of biogenic amines in the central nervous system and in peripheral tissues^{25,26}. Recent studies have revealed that MAOs are associated with psychiatric and neurological disorders, including depression, Parkinson's disease (PD) and AD^{27,28}. MAO-A inhibitors are used as antidepressants and anti-anxiety agents in the clinic, while MAO-B inhibitors are used as therapeutics for PD²⁹. As oxidative stress has been implicated in central nervous system degenerative disorders, MAO-B inhibitors are also potential candidates as anti-AD drugs due to their regulation of neurotransmitters and capacity to inhibit oxidative damage³⁰.

Because AD is a complex neurodegenerative disorder resulting from multiple factors, molecules that modulate the activity of a single protein target might not be able to significantly alter the progression of the disease. In recent years, a multi-target-directed ligand (MTDL) strategy has been developed to approach this multifaceted disease. Based on the MTDL drug design strategy, Sterling et al. incorporated the pharmacophores of rivastigmine (an AChE inhibitor, Figure 1) and rasagiline (an MAO inhibitor, Figure 2) to a single molecular entity to give a novel dual AChE and MAO-B inhibitors, ladostigil (Figure 2), the Phase II clinical trials of which is in progress³¹.

Homoisoflavonoids, which have been found to possess various biological properties, such as antifungal³², antiviral^{33,34}, anti-

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proliferative³², antioxidant³⁵ and anti-inflammatory³⁶ activities, were also proven to be potent and selective MAO-B inhibitors³⁷. Inspired by the outcomes of ladostigil, we have developed a series of tacrine-contained hybrids targeted to both AChE and MAO-B in earlier studies^{38,39}. Herein, we described the synthesis and evaluation of a series of homoisoflavonoid derivatives as dual AChE and MAO-B inhibitors (Figure 3), which were expected to bind to MAO-B, PAS and CAS of AChE.

Materials and methods

Materials

¹H NMR and ¹³C NMR spectra were recorded on a Bruker BioSpin GmbH spectrometer (Östliche Rheinbrückenstr,

Germany) at 400.132 and 100.614 MHz, respectively, using tetramethylsilane as the internal standard. Coupling constants are given in Hz. MS spectra were recorded on a Agilent LC–MS 6120 instrument (Santa Clara, CA) with an ESI mass selective detector. Flash column chromatography was performed with silica gel (200–300 mesh) purchased from Qingdao Haiyang Chemical Co. Ltd. (Qingdao, China). The chemicals used in the chemistry section were procured from Sinopharm Chemical Reagent Co., Ltd. (Shanghai, China) and all of the chemicals are analytical reagents. The purities of the synthesized compounds were confirmed to be higher than 95% by analytical HPLC (Agilent technologies 1200 series system) with a TC-C8 column (4.6 × 150 mm, 5 μm) and the compounds were eluted with CH₃OH/water (0.1% TFA, w/v) in ratios of 70:30–40:60 at a flow rate of 1.0 mL/min.

General procedures for the preparation of 5–8

To a stirred suspension of compounds **4** (10 mmol) and K₂CO₃ (15 mmol) in butanone (50 mL), KI (1 mmol) and α,ω-dibromo alkane (12 mmol) were added. The mixture was stirred at room temperature for 1–4 h, filtered, and then evaporated under vacuum. The crude product was chromatographed on a silica gel column, eluted with EtOAc/petroleum ether as eluent to afford the proposed compound.

(*E*)-7-(2-bromoethoxy)-3-(4-methoxybenzylidene)chroman-4-one (**5**). Compound **4** was treated with 1,2-dibromoethane according to the general procedure to give the desired product **5** as yellow solid, yield 74%; ¹H NMR (400 MHz, CDCl₃) δ: 7.77 (d, *J* = 8.4 Hz, 1H), 7.65 (s, 1H), 7.30 (d, *J* = 8.6 Hz, 2H), 7.04 (d, *J* = 8.6 Hz, 2H), 6.34 (d, *J* = 8.4 Hz, 1H), 6.31 (s, 1H), 5.30 (s, 2H), 4.51 (t, *J* = 6.0 Hz, 2H), 3.89 (t, *J* = 6.0 Hz, 2H), 3.81 (s, 3H).

(*E*)-7-(3-bromopropoxy)-3-(4-methoxybenzylidene)chroman-4-one (**6**). Compound **4** was treated with 1,3-dibromopropane according to the general procedure to give the desired product **6** as yellow solid, yield 65%; ¹H NMR (400 MHz, CDCl₃) δ: 7.80 (d, *J* = 8.8 Hz, 1H), 7.68 (s, 1H), 7.34 (d, *J* = 8.4 Hz, 2H), 7.09 (d, *J* = 8.4 Hz, 2H), 6.38 (d, *J* = 8.8 Hz, 1H), 6.34 (s, 1H), 5.35 (s, 2H), 4.37 (t, *J* = 6.4 Hz, 2H), 3.83 (s, 3H), 3.70 (t, *J* = 6.4 Hz, 2H), 2.46–2.51 (m, 2H).

(*E*)-7-(4-bromobutoxy)-3-(4-methoxybenzylidene)chroman-4-one (**7**). Compound **4** was treated with 1,4-dibromobutane according to the general procedure to give the desired product **7** as yellow solid, yield 59%; ¹H NMR (400 MHz, CDCl₃) δ: 7.76–7.81

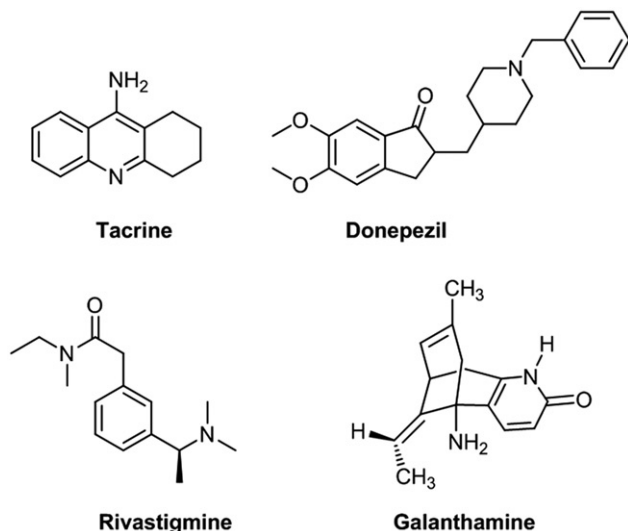


Figure 1. Structure of tacrine, donepezil, rivastigmine and galanthamine.

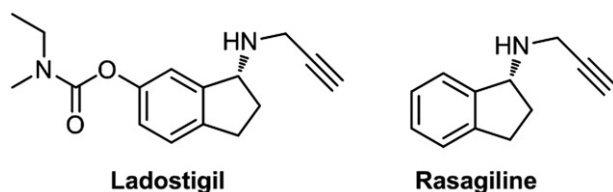
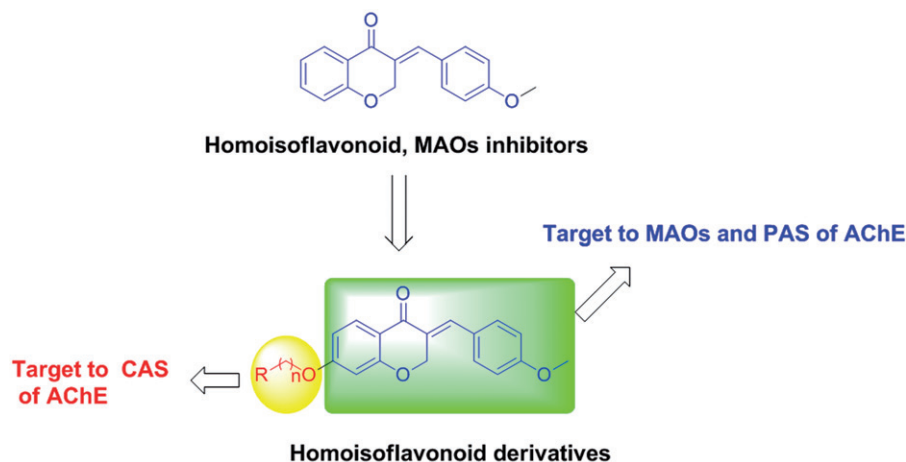


Figure 2. Chemical structures of ladostigil and rasagiline.

Figure 3. Design strategy for homoisoflavonoid derivatives.



(m, 1H), 7.62 (s, 1H), 7.29 (d, $J = 8.6$ Hz, 2H), 7.04 (d, $J = 8.6$ Hz, 2H), 6.31 (d, $J = 8.0$ Hz, 1H), 6.29 (s, 1H), 5.32 (s, 2H), 4.26 (t, $J = 6.4$ Hz, 2H), 3.80 (s, 3H), 3.52 (t, $J = 6.4$ Hz, 2H), 2.11–2.28 (m, 4H).

(*E*)-7-(5-bromopentyloxy)-3-(4-methoxybenzylidene)chroman-4-one (**8**). Compound **4** was treated with 1,5-dibromopentane according to the general procedure to give the desired product **8** as a yellow solid, yield 51%; ^1H NMR (400 MHz, CDCl_3) δ : 7.81 (d, $J = 8.6$ Hz, 1H), 7.64 (s, 1H), 7.33 (d, $J = 8.6$ Hz, 2H), 7.07 (d, $J = 8.6$ Hz, 2H), 6.29–6.31 (m, 1H), 6.30 (s, 1H), 5.34 (s, 2H), 4.22 (t, $J = 6.4$ Hz, 2H), 3.80 (s, 3H), 3.47 (t, $J = 6.8$ Hz, 2H), 1.96–2.05 (m, 4H), 1.57–1.82 (m, 2H).

General procedure for the synthesis of **9–18**

The substituent amine (2.4 mmol) was added to a magnetically stirred solution of **5**, **6**, **7** or **8** (2 mmol) in acetonitrile (30 mL). The mixture was refluxed for 12–24 h, and monitored by TLC. The mixture evaporated under vacuum. The residue was purified by column chromatography (petroleum ether/ethyl acetate, 2:1–10:1, plus 10 mL triethylamine per 1000 mL).

(*E*)-3-(4-methoxybenzylidene)-7-(2-(piperidin-1-yl)ethoxy)chroman-4-one (**9**). Intermediate **5** was treated with piperidine according to the general procedure to give the desired product **9** as a yellow solid, yield 69%; ^1H NMR (400 MHz, CDCl_3) δ : 7.81 (d, $J = 8.8$ Hz, 1H), 7.65 (s, 1H), 7.10 (d, $J = 8.2$ Hz, 2H), 6.80 (d, $J = 8.3$ Hz, 2H), 6.49 (d, $J = 8.7$ Hz, 1H), 6.27 (s, 1H), 5.19 (s, 2H), 3.98 (t, $J = 5.7$ Hz, 2H), 3.69 (s, 3H), 2.63 (t, $J = 5.7$ Hz, 2H), 2.36 (s, 4H), 1.51–1.28 (m, 6H). ^{13}C NMR (101 MHz, CDCl_3) δ : 180.65, 165.07, 162.78, 160.48, 136.30, 131.87, 129.47, 128.78, 127.07, 115.73, 114.15, 110.63, 101.33, 67.95, 66.39, 57.56, 55.28, 54.97, 25.90, 24.12. LC/MS (ESI) m/z : $[\text{M} + \text{H}]^+ = 394.2$. Purity: 96.2% (by HPLC).

(*E*)-3-(4-methoxybenzylidene)-7-(3-(piperidin-1-yl)propoxy)chroman-4-one (**10**). Intermediate **6** was treated with piperidine according to the general procedure to give the desired product **10** as a yellow solid, yield 74%; ^1H NMR (400 MHz, CDCl_3) δ : 7.89–7.73 (m, 1H), 7.64 (s, 1H), 7.10 (d, $J = 8.2$ Hz, 2H), 6.80 (d, $J = 8.4$ Hz, 2H), 6.47 (dd, $J = 7.2$, 4.8 Hz, 1H), 6.25 (d, $J = 3.4$ Hz, 1H), 5.18 (d, $J = 2.4$ Hz, 2H), 3.88 (d, $J = 6.3$ Hz, 2H), 3.67 (dd, $J = 9.2$, 5.7 Hz, 3H), 2.42–2.04 (m, 6H), 1.83 (d, $J = 5.8$ Hz, 2H), 1.54–1.24 (m, 6H). ^{13}C NMR (101 MHz, CDCl_3) δ : 180.63, 165.33, 162.81, 160.45, 136.23, 131.86, 129.43, 128.79, 127.05, 115.58, 114.14, 110.59, 101.20, 67.92, 66.85, 55.64, 55.25, 54.56, 26.53, 25.91, 24.39. LC/MS (ESI) m/z : $[\text{M} + \text{H}]^+ = 408.2$. Purity: 98.6% (by HPLC).

(*E*)-3-(4-methoxybenzylidene)-7-(4-(piperidin-1-yl)butoxy)chroman-4-one (**11**). Intermediate **7** was treated with piperidine according to the general procedure to give the desired product **11** as a yellow solid, yield 62%; ^1H NMR (400 MHz, CDCl_3) δ : 7.80 (dd, $J = 8.8$, 1.4 Hz, 1H), 7.65 (s, 1H), 7.10 (d, $J = 7.2$ Hz, 2H), 6.84–6.77 (m, 2H), 6.47 (d, $J = 8.7$ Hz, 1H), 6.24 (s, 1H), 5.19 (s, 2H), 3.85 (dd, $J = 6.1$, 5.1 Hz, 2H), 3.68 (dd, $J = 7.0$, 1.7 Hz, 3H), 2.22 (dd, $J = 14.7$, 6.6 Hz, 6H), 1.70–1.38 (m, 10H). ^{13}C NMR (101 MHz, CDCl_3) δ : 180.64, 165.33, 162.81, 160.45, 136.24, 131.85, 129.44, 128.79, 127.05, 115.56, 114.13, 110.59, 101.14, 68.17, 67.92, 58.86, 55.25, 54.51, 27.11, 25.89, 24.42, 23.28. LC/MS (ESI) m/z : $[\text{M} + \text{H}]^+ = 422.2$. Purity: 99.6% (by HPLC).

(*E*)-3-(4-methoxybenzylidene)-7-(5-(piperidin-1-yl)pentyloxy)chroman-4-one (**12**). Intermediate **8** was treated with piperidine according to the general procedure to give the desired product **12** as a yellow solid, yield 58%; ^1H NMR (400 MHz, CDCl_3) δ : 7.94

(d, $J = 8.8$ Hz, 1H), 7.79 (s, 1H), 7.26 (d, $J = 9.0$ Hz, 2H), 6.95 (d, $J = 8.8$ Hz, 2H), 6.60 (dd, $J = 8.8$, 2.4 Hz, 1H), 6.37 (d, $J = 2.3$ Hz, 1H), 5.34 (d, $J = 1.8$ Hz, 2H), 4.02–3.92 (m, 2H), 3.84 (d, $J = 10.1$ Hz, 3H), 2.36 (s, 4H), 2.32–2.28 (m, 2H), 1.84–1.75 (m, 2H), 1.63–1.54 (m, 6H), 1.50–1.42 (m, 4H). ^{13}C NMR (101 MHz, CDCl_3) δ : 180.92, 165.47, 162.91, 160.53, 136.41, 131.86, 129.59, 128.98, 127.23, 115.68, 114.21, 110.66, 101.21, 77.35, 77.04, 76.72, 68.29, 68.00, 59.36, 55.36, 54.68, 28.94, 26.67, 26.01, 24.49, 24.12. LC/MS (ESI) m/z : $[\text{M} + \text{H}]^+ = 436.3$. Purity: 99.0% (by HPLC).

(*E*)-7-(3-(dimethylamino)propoxy)-3-(4-methoxybenzylidene)chroman-4-one (**13**). Intermediate **6** was treated with dimethylamine according to the general procedure to give the desired product **13** as a yellow solid, yield 75%; ^1H NMR (400 MHz, CDCl_3) δ : 7.94 (d, $J = 8.8$ Hz, 1H), 7.80 (s, 1H), 7.28–7.23 (m, 2H), 6.96 (d, $J = 8.7$ Hz, 2H), 6.61 (dd, $J = 8.8$, 2.3 Hz, 1H), 6.39 (d, $J = 2.3$ Hz, 1H), 5.35 (d, $J = 1.8$ Hz, 2H), 4.05 (s, 3H), 3.85 (s, 2H), 2.64–2.59 (m, 2H), 2.36 (s, 6H), 1.98 (s, 2H). ^{13}C NMR (101 MHz, CDCl_3) δ : 180.96, 165.16, 162.91, 160.56, 136.55, 131.89, 129.63, 128.89, 127.18, 115.84, 114.22, 110.57, 101.30, 68.01, 66.34, 55.65, 55.37, 44.48, 26.36, 22.39. LC/MS (ESI) m/z : $[\text{M} + \text{H}]^+ = 368.2$. Purity: 95.1% (by HPLC).

(*E*)-7-(3-(diethylamino)propoxy)-3-(4-methoxybenzylidene)chroman-4-one (**14**). Intermediate **6** was treated with diethylamine according to the general procedure to give the desired product **14** as a yellow solid, yield 70%; ^1H NMR (400 MHz, CDCl_3) δ : 7.96 (d, $J = 8.8$ Hz, 1H), 7.82 (s, 1H), 7.31–7.26 (m, 2H), 6.98 (d, $J = 8.8$ Hz, 2H), 6.64 (dd, $J = 8.8$, 2.4 Hz, 1H), 6.42 (d, $J = 2.3$ Hz, 1H), 5.37 (d, $J = 1.8$ Hz, 2H), 4.07 (s, 2H), 3.87 (s, 3H), 2.56 (dt, $J = 21.4$, 7.2 Hz, 6H), 1.94 (dt, $J = 13.2$, 6.4 Hz, 2H), 1.04 (t, $J = 7.1$ Hz, 6H). ^{13}C NMR (101 MHz, CDCl_3) δ : 180.95, 165.49, 162.92, 160.53, 136.42, 131.87, 129.58, 128.99, 127.24, 115.70, 114.21, 110.69, 101.26, 68.01, 66.79, 55.37, 49.22, 47.04, 26.94, 11.83. LC/MS (ESI) m/z : $[\text{M} + \text{H}]^+ = 396.2$. Purity: 99.9% (by HPLC).

(*E*)-7-(3-(dipropylamino)propoxy)-3-(4-methoxybenzylidene)chroman-4-one (**15**). Intermediate **6** was treated with dipropylamine according to the general procedure to give the desired product **15** as a yellow solid, yield 60%; ^1H NMR (400 MHz, CDCl_3) δ : 7.96 (d, $J = 8.8$ Hz, 1H), 7.82 (s, 1H), 7.32–7.24 (m, 2H), 6.98 (d, $J = 8.8$ Hz, 2H), 6.63 (dd, $J = 8.8$, 2.4 Hz, 1H), 6.41 (d, $J = 2.3$ Hz, 1H), 5.37 (d, $J = 1.8$ Hz, 2H), 4.07 (d, $J = 6.3$ Hz, 2H), 3.87 (s, 3H), 2.58 (t, $J = 6.9$ Hz, 2H), 2.43–2.32 (m, 4H), 1.96–1.88 (m, 2H), 1.46 (dd, $J = 14.9$, 7.4 Hz, 4H), 0.88 (t, $J = 7.4$ Hz, 6H). ^{13}C NMR (101 MHz, CDCl_3) δ : 180.94, 165.55, 162.92, 160.53, 136.40, 131.86, 129.57, 129.01, 127.24, 115.67, 114.21, 110.70, 101.23, 68.01, 66.73, 56.33, 55.37, 50.42, 27.08, 20.39, 11.94. LC/MS (ESI) m/z : $[\text{M} + \text{H}]^+ = 424.3$. Purity: 99.5% (by HPLC).

(*E*)-3-(4-methoxybenzylidene)-7-(3-(pyrrolidin-1-yl)propoxy)chroman-4-one (**16**). Intermediate **6** was treated with pyrrolidine according to the general procedure to give the desired product **16** as a yellow solid, yield 62%; ^1H NMR (400 MHz, CDCl_3) δ : 7.96 (d, $J = 8.8$ Hz, 1H), 7.81 (s, 1H), 7.30–7.26 (m, 2H), 6.98 (d, $J = 8.8$ Hz, 2H), 6.64 (dd, $J = 8.8$, 2.3 Hz, 1H), 6.42 (d, $J = 2.3$ Hz, 1H), 5.37 (d, $J = 1.7$ Hz, 2H), 4.09 (t, $J = 6.4$ Hz, 2H), 3.87 (s, 3H), 2.66–2.59 (m, 2H), 2.54 (dd, $J = 9.2$, 3.9 Hz, 4H), 2.06–2.00 (m, 2H), 1.83–1.77 (m, 4H). ^{13}C NMR (101 MHz, CDCl_3) δ : 180.98, 165.43, 162.92, 160.54, 136.46, 131.88, 129.59, 128.98, 127.23, 115.72, 114.22, 110.69, 101.30, 68.01, 66.85, 55.38, 54.22, 52.90, 28.60, 23.47. LC–MS (ESI+APCI) m/z : $[\text{M} + \text{H}]^+ = 394.2$. Purity: 98.4% (by HPLC).

(*E*)-3-(4-methoxybenzylidene)-7-(3-morpholinopropoxy)chroman-4-one (**17**). Intermediate **6** was treated with morpholine according to the general procedure to give the desired product **17** as a yellow solid, yield 75%; ^1H NMR (400 MHz, CDCl_3) δ 7.97 (d, $J=8.8$ Hz, 1H), 7.82 (s, 1H), 7.31–7.25 (m, 2H), 6.98 (d, $J=8.8$ Hz, 2H), 6.64 (dd, $J=8.8$, 2.4 Hz, 1H), 6.42 (d, $J=2.3$ Hz, 1H), 5.37 (d, $J=1.8$ Hz, 2H), 4.09 (t, $J=6.3$ Hz, 2H), 3.86 (d, $J=10.7$ Hz, 3H), 3.75–3.72 (m, 4H), 2.55–2.51 (m, 2H), 2.50–2.46 (m, 4H), 2.03–1.96 (m, 2H). ^{13}C NMR (101 MHz, CDCl_3) δ 180.96, 165.35, 162.92, 160.56, 136.54, 131.89, 129.63, 128.91, 127.20, 115.78, 114.23, 110.65, 101.28, 68.03, 66.96, 66.51, 55.33 (d, $J=10.6$ Hz), 53.74, 26.19. LC/MS (ESI) m/z : $[\text{M} + \text{H}]^+ = 410.2$ Purity: 99.4% (by HPLC).

(*E*)-7-(3-(cyclohexylamino)propoxy)-3-(4-methoxybenzylidene)-chroman-4-one (**18**). Intermediate **6** was treated with cyclohexylamine according to the general procedure to give the desired product **18** as a yellow solid, yield 59%; ^1H NMR (400 MHz, CDCl_3) δ 7.97 (d, $J=8.8$ Hz, 1H), 7.82 (s, 1H), 7.31–7.25 (m, 2H), 6.98 (d, $J=8$ Hz, 1H), 7.81 (s, 1H), 7.27 (d, $J=8.9$ Hz, 2H), 7.01–6.89 (m, 2H), 6.62 (dd, $J=8.8$, 2.4 Hz, 1H), 6.41 (d, $J=2.3$ Hz, 1H), 5.36 (d, $J=1.8$ Hz, 2H), 4.09 (d, $J=6.2$ Hz, 2H), 3.86 (s, 3H), 2.82 (t, $J=7.0$ Hz, 2H), 2.45 (s, 1H), 1.97 (t, $J=6.6$ Hz, 2H), 1.94–1.86 (m, 2H), 1.78–1.70 (m, 2H), 1.63 (d, $J=11.7$ Hz, 2H), 1.32–1.18 (m, 4H), 1.17–1.11 (m, 1H). ^{13}C NMR (101 MHz, CDCl_3) δ 180.94, 165.35, 162.91, 160.54, 136.48, 131.88, 129.61, 128.94, 127.21, 115.76, 114.22, 110.64, 101.28, 68.02, 66.88, 56.85, 55.37, 43.66, 33.60, 29.94, 26.15, 25.06. LC/MS (ESI) m/z : $[\text{M} + \text{H}]^+ = 422.3$. Purity: 98.2% (by HPLC).

Biological activity

In vitro inhibition studies on AChE and BuChE

Acetylcholinesterase (AChE, E.C. 3.1.1.7, from *electric eel*), butylcholinesterase (BuChE, E.C. 3.1.1.8, from equine serum), 5,5'-dithiobis-(2-nitrobenzoic acid) (Ellman's reagent, DTNB), acetylthiocholine chloride (ATC) and butylthiocholine chloride (BTC) were purchased from Sigma-Aldrich (St. Louis, MO). Compounds were dissolved in DMSO (10 mM) and diluted in 0.1 M $\text{KH}_2\text{PO}_4/\text{K}_2\text{HPO}_4$ buffer (pH 8.0) to the desired final concentration. All the compounds are soluble at the tested concentration. DMSO was diluted to a concentration of less than 0.01%, and no inhibitory action on either AChE or BuChE was detected in separate prior experiments at this concentration. The *in vitro* AChE assay was performed as follows: all assays were conducted in 0.1 M $\text{KH}_2\text{PO}_4/\text{K}_2\text{HPO}_4$ buffer (pH 8.0) using a Shimadzu UV-2450 spectrophotometer. The assay medium (1 mL) consisted of phosphate buffer (pH 8.0), 50 μL of 0.01 M DTNB, 10 μL of enzyme and 50 μL of 0.01 M substrate (ACh chloride solution). Test compounds were added to the assay solution and pre-incubated with the enzyme at 37 °C for 15 min, followed by addition of substrate. The activity was determined by measuring the increase in absorbance at 412 nm at 1 min intervals at 37 °C. Calculations were performed according to the method of the equation in Ellman et al. Each concentration was assayed in triplicate.

The *in vitro* BuChE assay was performed similarly to the method described above.

Kinetic characterisation of AChE inhibition

Kinetic characterisation of AChE was performed using a reported method. Test compound was added into the assay solution and pre-incubated with the enzyme at 37 °C for 15 min, followed by addition of substrate (the final concentrations of substrate were

0.05, 0.0625, 0.10, 0.125, 0.25, 0.50 mM, respectively). Kinetic characterisation of the hydrolysis of ATC catalyzed by AChE was performed spectrometrically at 412 nm. A parallel control with no inhibitor in the mixture allowed the change in activities to be measured at various times. The plots were assessed by a weighted least square analysis that assumed the variance of V to be a constant percentage of V for the entire data set. The slopes of these reciprocal plots were subsequently plotted against the concentration of the inhibitors in a weighted analysis, and K_i was determined as the ratio of the replot intercept to the replot slope.

Inhibition of MAO activity

The potential effects of the test compounds on hMAO activity were investigated by measuring their effects on the production of H_2O_2 from *p*-tyramine, using the Amplex Red MAO assay kit (Molecular Probes, Inc., Eugene, OR) and recombinant human MAO-A or MAO-B (Sigma-Aldrich) according to published procedures⁴⁰. Compounds were dissolved in DMSO (10 mM) and diluted in 0.1 M $\text{KH}_2\text{PO}_4/\text{K}_2\text{HPO}_4$ buffer (pH 8.0) to the desired final concentration. All the compounds are soluble at the tested concentration. Adequate amounts of recombinant hMAO-A or hMAO-B (Sigma-Aldrich) were acquired and adjusted to 12.5 $\mu\text{g}/\text{mL}$ for hMAO-A and 75 $\mu\text{g}/\text{mL}$ for hMAO-B. Test drugs (20 μL) and MAO (80 μL) were incubated at 37 °C for 15 min in a flat-black-bottom 96-well microtest plate in dark. The reaction was started by adding 200 μM Amplex Red reagent, 2 U/mL horseradish peroxidase and 2 mM *p*-tyramine for hMAO-A or 2 mM benzylamine for hMAO-B, at 37 °C for 20 min. The reaction was quantified in a multi-detection microplate fluorescence reader based on the fluorescence generated (excitation, 545 nm; emission, 590 nm).

The specific fluorescence emission was calculated after subtraction of the background activity, which was determined from vials containing all components except the hMAO isoforms, which were replaced by a sodium phosphate buffer solution.

Reversibility and irreversibility experiments

To evaluate whether compound **10** is a reversible or irreversible hMAO-B inhibitor, an effective centrifugation ultrafiltration method (so-called "repeated washing") was used⁴⁰. The experiment was performed according to a reported method. Adequate amounts of recombinant monoamine oxidase with or without test drugs were incubated at 37 °C for 15 min. An aliquot from this incubation was stored at 4 °C for later use in the measurement of MAO activity. Another aliquot was transferred to an Amicon Ultra-0.5 Centrifugal Filter Unit with Ultracel-30 membrane (Millipore) and centrifuged (4 °C, 9000g, 20 min) three times. The enzyme was obtained and used for the subsequent measurement of activity using a method similar to that described above (the section "Inhibition of MAO activity"). The corresponding values of percent hMAO-B inhibition were separately calculated for samples with and without repeated washing.

Docking study

The simulation system was built based on the structure obtained from the Protein Data Bank (PDB: 2V60 for MAO-B, 2CMF for AChE). The heteroatoms and water molecules were removed, and all hydrogen atoms were subsequently added to the protein. Then forcefield was assigned to the enzyme. The ligand binding site was defined as 13 Å from the original ligand. Prior to the docking calculations, the original ligand was removed. The 3D structures of compound **10** were generated and optimized with the Discovery Studio 2.1 package (Accelrys Inc., San Diego, CA).

The CDOCKER program of the Discovery Studio 2.1 software, which allows full flexibility of ligands, was used to perform docking simulations. The docking and subsequent scoring were performed using the default parameters of the CDOCKER program. CDOCKER_INTERACTION_ENERGY is used like a score where a lower value indicates a more favorable binding.

Results and discussion

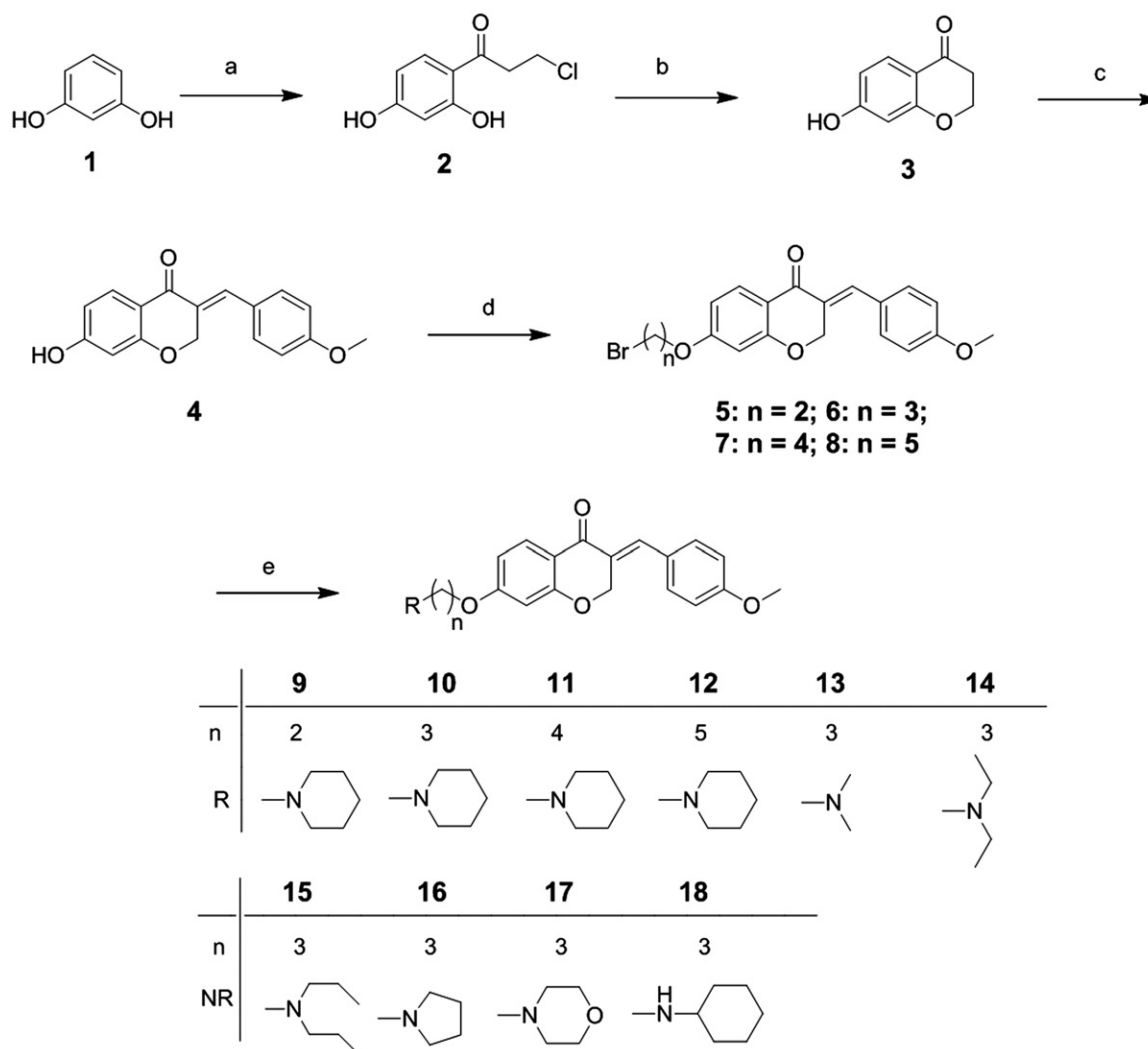
Chemistry

The homoisoflavonoid derivatives **9–18** were synthesized according to the synthetic protocol outlined in Scheme 1. First, commercial available resorcinol (**1**) reacted with 3-chloropropionic acid *via* Fries rearrangement to yield compound **2**, and the followed cyclization of **2** in aqueous NaOH gave 7-hydroxychroman-4-one (**3**). Then, (*E*)-7-hydroxy-3-(4-methoxybenzylidene)-chroman-4-one (**4**) was obtained by the condensation of **3** with *p*-methoxybenzaldehyde³⁹. The reaction of **4** in butanone with α,ω -dibromo alkane in the presence of K_2CO_3 provided responding intermediates **5–8**. Finally, target products **9–18** were got by the reactions of **5–8** with commercially available secondary amines (dimethylamine, pyrrolidine, etc.) in 53–87% yields. All the compounds were characterized using analytical and NMR spectroscopic data (see Supplementary data).

Compounds **5–18** were synthesized by the following procedures. Intermediates **2**, **3** and **4** were synthesized according to reference³⁹.

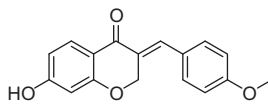
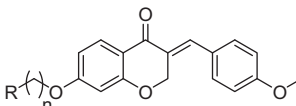






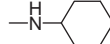
In vitro inhibition of AChE and BuChE

The AChE (E.C.3.1.1.7) inhibitory effects of the homoisoflavonoid derivatives were determined by the spectroscopic method described by Ellman et al. using rivastigmine as the standard⁴¹. As shown in Table 1, most of the hybrids demonstrated potent inhibitory activity against AChE and BuChE with IC_{50} values in the sub-micromolar to micromolar range. Compound **9**, a piperidine moiety linked with homoisoflavonoid by two carbon spacers, gave the most potent inhibitory activity ($IC_{50} = 0.74 \mu M$), which was about 135-fold more potent than lead compound **4** ($IC_{50} > 100 \mu M$). A simple structure–activity relationship analysis showed that the AChE inhibitory potency closely related to the length of the alkylene chain. When the length of the alkylene chain increased to 5, the AChE inhibitory activity decreased dramatically (**12**, $IC_{50} = 14.1 \mu M$). From the IC_{50} values of the tested compounds, it appears that a 3-carbon spacer between homoisoflavonoid moiety and amino group seems to be the proper length for balancing the inhibition activity against AChE and MAO-B. Moreover, amino group is also important for the inhibitory activities. Among the three aliphatic substituted amino derivatives, the *N,N*-dimethyl amino group (**13**,



Scheme 1. Reagents and conditions: (a) 3-chloropropionic acid trifluoromethanesulfonic acid; (b) NaOH, 0 °C; (c) *p*-methoxybenzaldehyde, piperidine, 80 °C; (d) α,ω -dibromo alkane, K_2CO_3 , KI, butanone, rt; (e) amines, acetonitrile, reflux.

Table 1. *In vitro* inhibition IC₅₀ (μM) and selectivity of compounds **4**, **9–18** on ChEs and MAOs.

							
		4		9 - 18			
Compound	R	n	IC ₅₀ ^a (μM)		MAO-A ^d inhibition rate at 50 μM ^f	MAO-B ^e IC ₅₀ ^a (μM)	
			AChE ^b	BuChE ^c			
9		—	>100	>100	24.1%	1.06 ± 0.01	
		2	0.74 ± 0.01	16.62 ± 1.30	8.0%	>25	
10		3	3.94 ± 0.05	9.21 ± 0.63	24.7%	3.44 ± 0.04	
11		4	5.73 ± 0.41	9.48 ± 0.26	28.5%	1.63 ± 0.06	
12		5	14.1 ± 0.65	8.45 ± 0.64	30.0%	0.65 ± 0.05	
13		3	3.78 ± 0.07	40.50 ± 4.90	19.1%	8.37 ± 0.80	
14		3	2.45 ± 0.26	23.88 ± 2.39	20.0%	11.11 ± 0.25	
15		3	7.47 ± 0.17	21.47 ± 0.35	22.4%	11.44 ± 1.54	
16		3	2.83 ± 0.40	9.10 ± 0.02	24.8%	6.93 ± 0.49	
17		3	8.51 ± 0.33	5.01 ± 0.40	37.9%	8.96 ± 0.77	
18		3	27.86 ± 0.68	34.15 ± 3.05	1.3%	24.75 ± 0.76	
Rivastigmine	—	—	5.10 ± 0.13	3.11 ± 0.04	n.t. ^g	n.t. ^g	
Ladostigil	—	—	50.0 ± 4.8	n.t. ^g	n.t. ^g	37.1 ± 3.1	
Clorgyline	—	—	n.t. ^g	n.t. ^g	4.1 ± 0.2 nM	n.t. ^g	
Pargyline	—	—	n.t. ^g	n.t. ^g	n.t. ^g	0.19 ± 0.01	

^aMean ± SD of at least three independent measurements.^bAChE from *electric eel* was used.^cBuChE from horse serum was used.^dHuman Recombinant MAO-A was used.^eHuman Recombinant MAO-B was used.^fTest concentration is 50 μM.^gn.t. = not tested.

IC₅₀ = 3.78 μM) and *N,N*-diethyl amino group (**14**, IC₅₀ = 2.45 μM) gave better results than *N,N*-dipropyl amino group (**15**, IC₅₀ = 7.47 μM). On the other hand, among the cyclic substituted amino derivatives, compounds **10** and **16**, piperidine and pyrrolidine linked with berberine by three carbon spacers, gave IC₅₀ values of 3.94 and 2.83 μM, respectively. Compound **17** (IC₅₀ = 7.47 μM), morpholine linked with homoisoflavanoid by three carbon spacers, afforded poor activities compared with compounds **10** and **16**. Surprisingly, the replacement of the piperidine group by a cyclohexylamine group markedly decreased the inhibitory activity (**10**, IC₅₀ = 3.94 μM versus **18**, IC₅₀ = 27.86 μM).

The *in vitro* BuChE (E.C. 3.1.1.8) inhibition activities were also determined using the same method. Most of the tested compounds demonstrated much higher inhibitory potency against BuChE than homoisoflavanoid **4**. Compound **10**, which exhibited a good balance of AChE/MAO-B inhibitory activity, was chosen for further studies.

Kinetic study of AChE

In order to investigate the mechanism of homoisoflavanoid derivatives against AChE, kinetic study was performed using compound **10** and the steady-state inhibition data for AChE are shown in Figure 4. Lineweaver–Burk reciprocal plots revealed that there was an increasing slope and an increasing intercept at higher inhibitor concentrations, indicating a mixed inhibition. Based on the kinetic analysis, we concluded that homoisoflavanoid derivatives exhibited as dual binding sites AChE inhibitors, which target catalytic and peripheral sites of AChE simultaneously.

Molecular modeling studies of AChE

To explore the interaction mode of homoisoflavanoid derivatives with AChE and the structure–activity relationships, molecular docking simulations were performed with the Discovery studio 2.1 software based on the structure of the complex of *Torpedo*

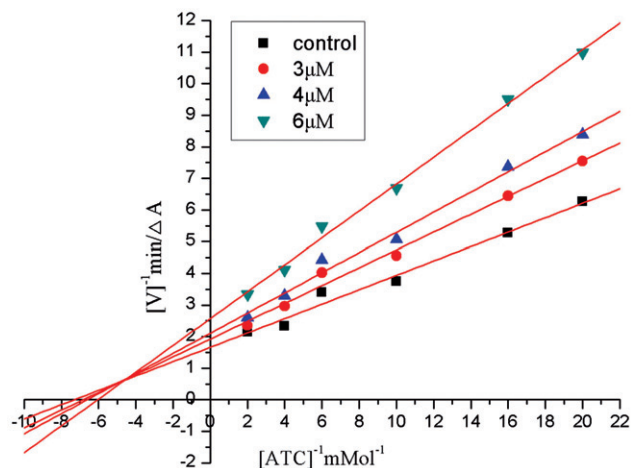


Figure 4. Steady state inhibition by compound **10** of AChE hydrolysis of ACh; the plots show mixed-type inhibition for compound **10** on AChE.

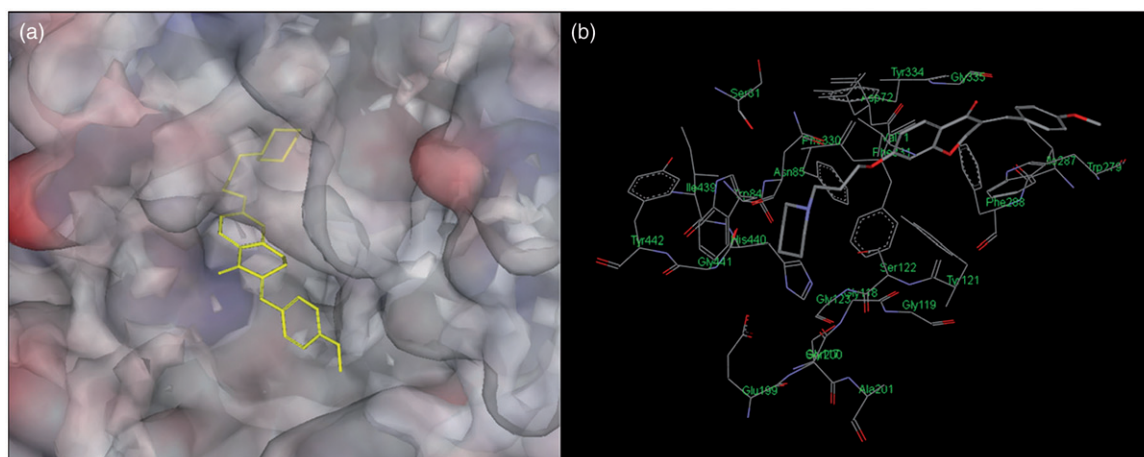


Figure 5. Docking models of the compound–enzyme complex. (a) stereoviews looking down the gorge of TcAChE binding with **10** (yellow). (b) **10** docked into the catalytic gorge of TcAChE, highlighting the protein residues belonging to ACS and PAS that establish the main interactions.

californica enzyme (TcAChE) (PDB entry 2CMF). As shown in Figure 5, compound **10** could simultaneously interact with the CAS and the PAS of AChE, which is consistent with the result of kinetic study. Near the bottom of the gorge, the protonated piperidine moiety of compound **10** interacts with Phe330 and Trp84 *via* hydrophobic interactions to form a cation– π interaction (4.76 and 4.32 Å, respectively). The homoisoflavonoid moiety of **10** reached the top of the cavity and interacted with PAS aromatic residues Trp279, phe331, tyr121 (5.48, 4.40 and 5.36 Å, respectively) through π – π stacking. The homoisoflavonoid moiety snaked along the gorge, interacting with Tyr334, Ser122 and Val71 by hydrophobic interactions.

In vitro inhibition studies of hMAO-A and hMAO-B

To confirm the multipotent biological profile of homoisoflavonoid derivatives, inhibitory activity against MAOs was determined using ladostigil as reference compound and the results are summarized in Table 1. Most of these target compounds exhibited high selective inhibition to hMAO-B in the micromolar range. Among them, compound **12** presented the most potent inhibition for hMAO-B with IC_{50} value of 0.65 μ M, which is 2-fold and 57-fold more potent than lead compound **4** and ladostigil, respectively. A structure–activity relationship analysis showed that the length of carbon spacer also played an important role in inhibition activity. However, unlike that of AChE, a longer linker resulted in

Table 2. Reversibility and irreversibility of hMAO-B inhibition of hybrid **10** and pargyline^a.

Compound	% hMAO-B inhibition ^b	
	Before washing	After repeated washing
10 (3 μ M)	40.3 \pm 1.33	34.5 \pm 0.89
10 (5 μ M)	75.2 \pm 3.84	65.9 \pm 1.351
Pargyline (200 nM)	64.4 \pm 5.21	51.7 \pm 3.94

^aPargyline is an irreversible hMAO-B inhibitor.

^bData are the mean \pm SD of three independent experiments.

better inhibitory activity, the inhibition activity against MAO-B increased as the increasing of the carbon spacer length. For example, compound **9**, in which piperidine group linked with homoisoflavonoid *via* a 2-carbon spacer, provides IC_{50} value of more than 25 μ M. Its homologous compound **12**, with 5-carbon spacer, gave 0.65 μ M of the IC_{50} value, which was 40 times as that of compound **9**. On the other hand, all of the target compounds showed very weak inhibition activity against MAO-A. Overall, among the synthesized compounds, compound **10** exhibited best balance of inhibition for both AChE and MAO-B.

Reversibility and irreversibility study of hMAO-B

The reversibility and irreversibility of inhibitory activities of **10** was assessed according to a method of reference⁴⁰. Pargyline, known as irreversible MAO-B inhibitor, was used as a reference compound. Data are listed in Table 2. Similarly with pargyline, MAO-B inhibition of **10** was irreversible for the enzyme activity, which did not restore after repeated washing.

Molecular modeling studies of MAO-B

To evaluate the binding modes of the homoisoflavonoid derivatives with MAO-B, docking simulation of compound **10** was carried out using the CDOCKER program in the Discovery studio 2.1 software based on the X-ray crystal structure of human

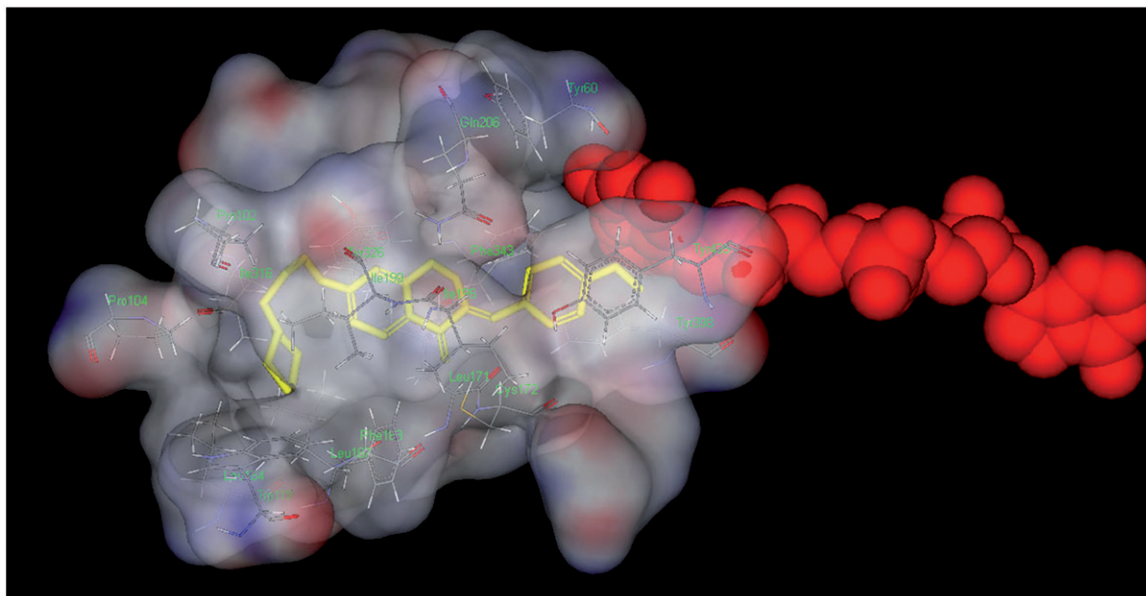


Figure 6. Docking pose of derivative **10** (yellow) into human MAO-B (2V60) highlighting the protein residues that establish the main interactions with **10**. The FAD cofactor was depicted using space fill representation.

MAO-B (PDB entry 2V60). As shown in Figure 6, the benzyl group of homoisoflavonoid moiety is properly oriented to the flavin adenine dinucleotide (FAD) cofactor adopting between Tyr398 (4.25 Å) and Tyr435 (4.33 Å) form face-to-face p-p stacking interactions in a “sandwich” form. Besides, the pyridine moiety and carbon linker are embedded into a hydrophobic pocket delimited by Pro102, Pro104, Trp119, Leu164, Leu167, Phe186, Ile316.

Conclusion

In summary, we have designed, synthesized and evaluated a series of homoisoflavonoid derivatives as dual functional cholinesterase and MAO inhibitors for the treatment of AD. Most of these compounds were potent inhibitors of ChE, which are more active than lead compound **4**. Meanwhile, compound **12** presented the most potent inhibition for hMAO-B with IC₅₀ value of 0.65 μM. Compound **10**, with a pyridine group linked with homoisoflavonoid moiety by a three carbon linker, provided a good balance of AChE and hMAO-B inhibition activities, with IC₅₀ value of 3.94 and 3.44 μM, respectively. Kinetic studies revealed that compound **10** was a mixed-type inhibitor of AChE and an irreversible inhibitor of hMAO-B. Molecular modeling studies indicated that **10** could interact with both CAS and PAS of AChE, which is consistent with kinetic studies. Based on these results, compound **10** is a promising “one-compound-multi-targets” candidate for AD treatment and needs further study.

Declaration of interest

The authors declare no conflicts of interest.

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Supplementary material available online
Supporting Information.