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Identification of “sarsasapogenin-aglyconed” timosaponins as novel Aβ-lowering modulators of amyloid precursor protein processing†

Lai-King Sy, Chun-Nam Lok, Juan-Yu Wang, Yungen Liu, Lu Cheng, Pui-Ki Wan, Chi-Ting Leung, Bei Cao, Wai-Lun Kwong, Raymond Chuen-Chung Chang and Chi-Ming Che*a

The inhibition of amyloid β (Aβ) peptide production is a key approach in the development of therapeutics for the treatment of Alzheimer’s disease (AD). We have identified that timosaponins consisting of sarsasapogenin (SSG) as the aglycone can effectively lower the production of Aβ peptides and stimulate neurite outgrowth in neuronal cell cultures. Structure–activity relationship studies revealed that the cis-fused Aβ ring, cis-configuration, spiroketal F-ring and 25S-configuration of SSG are the essential structural features responsible for the Aβ-lowering effects and neurite-stimulatory activity. New synthetic derivatives that retain the SSG scaffold also exhibited an Aβ lowering effect. Treatment of cells with timosaponins led to modulation of amyloid precursor protein (APP) processing through the suppression of β-cleavage and preferential lowering of the production of the 42-amino acid Aβ species (Aβ42) without affecting another γ-secretase substrate. The SSG and “SSG-aglyconed” timosaponins also penetrated brain tissue and lowered brain Aβ42 levels in mice. Our studies demonstrate that timosaponins represent a unique class of steroidal saponins that may be useful for the development of AD therapeutics.

Introduction

Alzheimer’s disease (AD) is a neurodegenerative disorder that starts with a decline in short-term memory and progresses to the loss of cognition and executive functions. The pathology of AD is characterized by synaptic loss, neuronal death, frequent deposition of phosphorylated tau proteins and Aβ aggregation within the brain. Although the underlying cause of AD is complex, the accumulation of Aβ within the brain appears to play a pivotal role in the onset and progression of the disease. Aβ is generated from the proteolysis of amyloid precursor protein (APP) during aging or in subjects with an inherited cause of AD. APP is a transmembrane protein whose proteolysis is mediated by α-, β- and γ-secretases that cleave APP at specific sites. The amyloidogenic process first involves the cleavage of APP to create a c-terminal fragment (CTF), known as β-CTF, which is subsequently cleaved by a multiprotein γ-secretase complex to produce different lengths of Aβ peptides such as Aβ10, Aβ14 and Aβ42. Genetic and mechanistic data strongly suggest that the accumulation of amyloidogenic Aβ42 peptide results in the formation of toxic oligomers and/or fibrils. Accordingly, Aβ42-lowering compounds that target the β- and/or γ-cleavage processes represent a promising strategy for therapeutic intervention. In AD, γ-secretase inhibitors (GSI) initially emerged as effective Aβ-lowering agents, but the side effects resulting from non-specific inhibition of other vital γ-secretase substrates, such as Notch, have complicated the development of these inhibitors. With evidence for the more specific role of Aβ42 in amyloidogenesis, current approaches to AD drug development are focused on γ-secretase modulators (GSM) that preferentially lower Aβ42 without affecting the action of γ-secretase on general APP processing and the cleavage of other substrates. Modulation of β-secretase is also considered to be a viable strategy for lowering Aβ production. A mutation in APP that hinders β-cleavage and lowers the production of Aβ by 40% was demonstrated to be protective against AD and age-related cognitive decline, supporting the idea that a moderate reduction of Aβ in humans is favourable for AD treatment or prevention. Natural products provide opportunities for the development of anti-AD pharmaceuticals (ESI Table S1†). For example, green tea polyphenols, ginsenosides and resveratrol have all been
shown to exhibit promising anti-amyloidogenic effects.10–12 In the course of studying the pharmacological properties of medicinal saponins from natural products, we found that timosaponins consisting of sarsasapogenin (SSG, 1) (Fig. 1) as the aglycone, including timosaponin A III (TAIII, 2) isolated from the rhizome of Anemarrhena asphodeloides Bge. (Liliaceae), could effectively lower Aβ production. We have previously identified TAIII’s anti-cancer and autophagy-inducing properties.13,14 Oral administration of SSG, TAIII and other timosaponins were shown to improve memory dysfunction in animal models of dementia despite the chemical structural differences between these compounds. Total saponins from Anemarrhena asphodeloides Bge. were reported to ameliorate diabetes-associated cognitive decline in rats and mediate Aβ decreases in brain.15 In the present study, we show that SSG (1) and other timosaponins (2–5, Fig. 1) specifically exhibit Aβ-lowering activities and their actions are akin to GSM,7 which preferentially lowers Aβ42 peptide production. We propose a model by which the timosaponins may bind to the steroid binding site of APP,20 possibly modulating the APP secretase properties. Importantly, SSG and several timosaponins showed brain penetration and Aβ42-diminishing activities in vivo.

Results and discussion

Synthesis of timosaponins, SSG analogues and SSG derivatives

A. asphodeloides is a medicinal herb that is rich in structurally closely related timosaponins whose isolation and purification are exceptionally difficult.17 Chemical synthesis, on the other hand, is a feasible strategy to obtain timosaponins that are of low natural abundance or commercially unavailable. Preparation of steroidal timosaponins (2–5) (Fig. 1) with different sugar lengths was achieved by reacting aglycone 1 with different glycosyl donors (ESI Fig. S1–S3†).

The chemical structure of the aglycone SSG (1) is rather unique in naturally-occurring sapogenins due to its structural characteristics, including a cis-fused AB ring (or 5b-conformation), 3β-configuration, spiroketal F-ring and 25S-configuration.21 In this study, a series of SSG analogues (6–24) (Fig. 2)
differing in these structural features were employed to study the structure–activity relationship (SAR) in the Aβ-lowering activities. The linkage of AB rings in steroidal sapogenins can be cis-, trans- or Δ^2-double bonded. Δ^2-Double bonded diosgenin (6) and yamogenin (7) were studied in addition to “diosgenin-aglyconed” sapogenins, including synthesized capisoside A7 (8) (ESI Fig. S4†), dioscin (9) and polyphyllin D (10). Hydrogenation of saponins 9 and 10 generated AB ring trans-fused dihydrodioscin (11) and dihydrophyllin D (12), respectively (ESI Fig. S5†). Similarly, hydrogenation of 6 and 7 produced AB ring trans-fused tigogenin (13) and neotigogenin (14), respectively (ESI Fig. S6†). Functionalization at Δ^2 of 6 or Δ^2 of 7 led to production of a pair of isomers for each compound: 5α-H, 6α-OH diosgenin (15), 5β-H, 6β-OH diosgenin (16) and 5α-H, 6α-OH yamogenin (17), 5β-H, 6β-OH yamogenin (18) (ESI Fig. S7†). The hydration products 15 and 17 are AB ring trans-fused, while the new compounds 16 and 18 are AB ring cis-fused.

Compound 1 has a secondary β-OH at C3. Oxidation of 1 to sarsasapogenone (19) followed by reduction gave episarsasapogenin (20) (ESI Fig. S8†). SSG 1 has heterocyclic rings E and F fixed at C22 in which the F-spiroketal ring appears to be a crucial moiety in bioactive saponins (Fig. 2). Reductive cleavage of the F-ring in 1 gave dihydroxysarsasapogenin (dSSG, 21) with a terminal OH group (ESI Fig. S9†). Timosaponin B I (TBI, 22) and timosaponin B II (TBII, 23) are two examples of “dSSG-aglyconed” saponins with two sugar chains substituted at C3 and C26. The contribution of the C25 S-configuration of 1 in decreasing Aβ42 was also investigated by comparing the activity of compound 16 and its epimer, smilagenin (24), both of which have a C25 R-configuration.

We have synthesized new SSG derivatives (28–29, 32–34) in an attempt to modify the biological activity and/or bioavailability (Fig. 3). Reaction of 1 with propargyl bromide yielded 25, containing a propargyl group, which can be linked to other moieties or probes via click chemistry. By reacting 25 with acetonide-protected α-galactose azide (26), via CuI-catalysed click chemistry, triazole SSG (27) was obtained. Removing the acetonide protecting groups of 27 yielded α-OMe triazole SSG (28) (ESI Fig. S10†). The relatively harsh reaction conditions (strong base NaH and long reaction time) used in the preparation of 25 was also attempted for the preparation of carboxylate ethereal SSG (29a) by reacting compound 1 with methyl broomoacetate, but without success. Compound 1 was then reacted with diazoacetate, in the presence of Rh2(OAc)4 as a catalyst, to obtain 29a in good yield via carbene insertion under mild conditions. Hydroxylation of the carboxylate 29a in alkaline conditions, followed by neutralization, produced the ethereal SSG (29) (ESI Fig. S11†). The carboxylic acid of 29 serves as a useful linkage for coupling with other moieties under mild conditions. Reaction of 29 with acetonide-protected α-galactose amine (30) gave amide SSG (31). The acetonide protecting groups of 31 were then removed by acid hydrolysis to yield α-OMe SSG (32), β-OMe SSG (33) and a mixture of α- and β-OH SSG (34) (ESI Fig. S12†).

Aβ42-lowering activities

Using Neuro-2A neuroblastoma cells stably transfected with APP with the AD-linked Swedish mutation (N2A-APPswe) as a cell culture model of Aβ production,27 it was found that the SSG 1 treatment modestly decreases Aβ42 production with an IC_{50} of 53 μM (Table 1). Treatment with timosaponins (2–5) markedly lowered Aβ42 production when compared to the aglycone SSG. SSG analogue 18, in addition to the newly synthesized SSG derivatives 28–29 and 32–34, also showed slight to moderate improvement in Aβ42-lowering activity when compared to 1 (Table 1). In rat primary cortical neuronal
Structure–activity relationship

The role of the sugar chain in timosaponins. Treatment of N2A-APPSwe cells with monosaccharide timosaponin A I (TAI, 3) and disaccharide TaIII (2) showed an improved reduction of Aβ42 when compared to aglycone SSG (Table 1), indicating that the presence of the sugar chain is beneficial in lowering Aβ42 levels. However, trisaccharide timosaponin A V (TAV, 4), reported for the first time, showed no further lowering effect (IC50 = ~4.2 μM) when compared to disaccharide 2 (IC50 = ~2.3 μM). In view of the importance of hydrophobicity to the cell permeability of the compounds to be tested, timosaponins with longer sugar chains were not considered in this study.16,17

Tinosaponins 2–4 are galactosyl derived and exhibit prominent Aβ42-level reducing effects. Asporasagrophytene A (AA, 5), a glucosyl-derived timosaponin (Fig. 1), shows a comparable Aβ42-lowering effect (IC50 = ~6.0 μM) to galactosyl 3 (IC50 = ~6.1 μM), revealing that the nature of the monosaccharide coupled to aglycone 1 has a negligible effect on Aβ42 levels.

The aforementioned enhancement in lowering Aβ42 production by the sugar chains in timosaponins has not been observed in the steroidal aglycone diosgenin 6 and its galactosyl product 8 (Table 1). Taken together with the insignificant Aβ42-lowering effects exhibited by diosgenyl saponins (data not shown) 9 and 10 (Δ5 double bond) and their corresponding hydrogenated products 11 and 12 (both AB trans-fused rings), it is suggested that aglycone 1 is critical in lowering Aβ42 production. The structural features associated with 1, including the AB-fused ring and the F-ring, were subjected to investigation and the findings are discussed below.

The roles of the AB-fused ring and the C3 configuration in SSG. Diosgenin 6 and yamogenin 7, both having Δ5 double bonds and different configurations at C25 (R- for 6 and S- for 7), showed no effect on Aβ42 lowering when compared to 1. Their respective hydrogenated products 13 and 14, both of which have AB rings trans-fused, were also ineffective (Table 1). In addition, only compound 18 (AB rings cis-fused) from the 4 hydrated products 15–18 caused reduction of Aβ42 levels (Table 1), indicating that the cis-fused AB ring (or 5β) in 1 is biologically significant in decreasing Aβ42 production.

Compound 19, with a ketone functionality at C3, showed a comparable effect to 1 in Aβ42 lowering, while compound 20, with a C3 α-OH, was inactive (Table 1), implying that the 3α-configuration is unfavourable for Aβ42 lowering.

The roles of the F-spiroketal ring and the C25 configuration in SSG. Steroidal sapogenins, including spirostane-, furostane- and cholestane-types, are widely distributed in the plant kingdom.8,30 SSG 1, which has an intact spirotetral F-ring, is an example of the spirotetral-type (Fig. 2). As mentioned above, this compound exhibits an Aβ42-lowering effect. However, such an effect was not observed for the F-ring cleaved, furostane-type dSSG 21 (IC50 > 100 μM). In addition, timosaponin B I (22) and timosaponin B II (23), possessing dSSG as the aglycone, are also ineffective in lowering Aβ42 (Table 1). Timosaponin B II (23), however, has been shown to inhibit the up-regulation of β-secretase induced by ferric chloride in rat retina.31 Ergosterol (structure not shown) structurally resembles a cholestane-type aglycone. This sterol also elicited no attenuation of Aβ42 production (IC50 > 100 μM). It is therefore concluded that the F-spiroketal ring in 1 should remain intact for Aβ42-lowering activity.

Smilagenin 24 and 6β-OH substituted 16 were ineffective in lowering Aβ42 (Fig. 2 and Table 1). Thus, one may envisage that

cultures, which produce and secrete low levels of Aβ, chronic exposure to compounds 1–3 also resulted in a moderate diminishment of Aβ42 levels in the medium (Table 2).

Table 1  IC50 values of timosaponins, SSG analogues and SSG derivatives in lowering Aβ42 production in N2A-APPSwe cells. Data represent mean ± standard deviation; n ≥ 3

<table>
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<tr>
<th>Compound</th>
<th>IC50 (μM)</th>
<th>Compound</th>
<th>IC50 (μM)</th>
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<tr>
<td>SSG (1)</td>
<td>53.0 ± 9.0</td>
<td>5β-H, 6β-OH yamogenin (17)</td>
<td>&gt;100</td>
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<tr>
<td>TAI (2)</td>
<td>2.3 ± 0.2</td>
<td>5β-H, 6β-OH yamogenin (18)</td>
<td>45.0 ± 4.0</td>
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<tr>
<td>TAI (3)</td>
<td>6.1 ± 2.8</td>
<td>Sarsasapogenone (19)</td>
<td>50.0 ± 5.0</td>
</tr>
<tr>
<td>TAV (4)</td>
<td>4.2 ± 1.2</td>
<td>Episarsasapogenin (20)</td>
<td>&gt;100</td>
</tr>
<tr>
<td>AA (5)</td>
<td>6.0 ± 1.4</td>
<td>Dihydroxysarsapogenin (21)</td>
<td>&gt;100</td>
</tr>
<tr>
<td>Diosgenin (6)</td>
<td>&gt;100</td>
<td>Timosaponin B I (22)</td>
<td>&gt;100</td>
</tr>
<tr>
<td>Yamogenin (7)</td>
<td>&gt;100</td>
<td>Timosaponin B II (23)</td>
<td>&gt;100</td>
</tr>
<tr>
<td>Capsicose A3 (8)</td>
<td>&gt;100</td>
<td>Smilagenin (24)</td>
<td>&gt;100</td>
</tr>
<tr>
<td>Tigogenin (13)</td>
<td>&gt;100</td>
<td>x-Ome triazole SSG (28)</td>
<td>6.5 ± 2.1</td>
</tr>
<tr>
<td>Neotigogenin (14)</td>
<td>&gt;100</td>
<td>Ethereal SSG (29)</td>
<td>27.0 ± 8.0</td>
</tr>
<tr>
<td>3α-H, 6α-OH diosgenin (15)</td>
<td>&gt;100</td>
<td>x-Ome SSG (32)</td>
<td>7.2 ± 2.2</td>
</tr>
<tr>
<td>3β-H, 6β-OH diosgenin (16)</td>
<td>&gt;100</td>
<td>β-Ome SSG (33)</td>
<td>9.3 ± 3.5</td>
</tr>
<tr>
<td>3α, 6β-OH SSG (34)</td>
<td>7.3 ± 4.0</td>
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Table 2  Effects of SSG, TaIII and TAI on Aβ42 levels in conditioned medium of rat primary cortical neuron culture upon 5-day incubation. A low concentration of SSG was used owing to the compound’s insolubility in the reduced serum medium for the neuronal culture. Data represent mean ± standard deviation; n = 3

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<th>Compound</th>
<th>Aβ42 reduction (%)</th>
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<tr>
<td>SSG (1) (5 μM)</td>
<td>25 ± 4</td>
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<tr>
<td>TAI (2) (5 μM)</td>
<td>42 ± 9</td>
</tr>
<tr>
<td>TAV (4) (10 μM)</td>
<td>28 ± 5</td>
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the 25S-configuration in 1 is one of the vital structural factors contributing to the compound’s Aβ-lowering activities.

Collectively, the SSG moiety is essential for Aβ42-lowering activity. Notably, the Aβ42-lowering effect of SSG is significantly enhanced when the compound is glycosylated at C3 to obtain timosaponins 2–5 or carboxylated at C3 to obtain SSG derivatives 28–29 and 32–34 (Fig. 1, 3 and Table 1). It is suggested that chemical modification at the C3 position of 1 is an appealing approach for the generation of versatile SSG derivatives with anti-amyloidogenic effects and thus timosaponins herein represent a class of interesting saponins noteworthy of further investigation related to Aβ42-lowering activities.

**Biochemical mechanisms**

**Timosaponins modulate APP processing with suppression of β-cleavage.** We have investigated whether the timosaponins interfere with APP processing by immunoblot analysis. Treatment of N2A-APPswe cells with TAIII did not elicit a change in the expression of full length APP, but resulted in the alteration of the expression of the CTF and secreted APP fragments (sAPP) (Fig. 4). In general, TAIII treatment decreased the levels of the β-secretase-cleaved CTF (C-99) and secreted APPβ in a concentration-dependent manner. Elevated concentrations (10 μM) of TAIII increased the α-secretase-cleaved CTF (C-83) and secreted APPα (Fig. 4). These changes in the expression of APP cleavage products are indicative of suppression of the amyloidogenic β cleavage and/or enhancement of the α cleavage, which is non-amyloidogenic and is competitive to the former. Other timosaponins (TAI/TAV/AA) that exhibited effective Aβ-lowering activities also elicited changes in APP cleavage products similar to that of TAIII (Fig. 5A). TAIII treatment did not affect the expression of the β-secretase BACE1 (Fig. 4), and β-secretase activities in the cell extracts or the activities of purified BACE1. Furthermore, TAIIII treatment also did not alter the expression of ADAM-10, an α-secretase that is activated by proteolytic cleavage (Fig. 4). Thus, these results suggest that the Aβ-lowering effects mediated by timosaponins are unlikely to be due to changes in the enzyme activities of α- and β-secretases. However, allosteric modulation of the APP processing complexes may be a possible cause for the alterations of APP cleavage elicited by the timosaponins.

We also studied the effect of SSG on the Aβ production and APP processing (Fig. 5C & D). The Aβ-lowering activity of SSG was weaker than those of timosaponins generally. Treatment of N2A-APPswe cells with SSG at 50 μM and 100 μM for 18 h caused a decrease in Aβ42 by 55% and 40%, respectively. Interestingly, treatment of cells with SSG at 100 μM increased both α-CTF and β-CTF expression with marked increase in the sAPPα levels. There was no change in β-secretase activity in the protein extracts from SSG-treated cells compared to that from untreated cells. Though the mechanism accounting for the difference between timosaponins and SSG in APP processing remains to be elucidated, it was concluded that the sugar moiety appears to play a role in modifying the effects of the timosaponins on APP processing.

**Timosaponins preferentially lower Aβ42 production similar to the action of GSM.** Aβ species of variable lengths are generated upon cleavage of the CTFs of APP by γ-secretase; these Aβ profiles can be used for predicting the mechanisms of action of drugs that act on the APP-secretase complexes. As revealed by ELISA specific for individual Aβ species, treatment
of cells with timosaponins (TAIII/TAI/TAV/AA) effectively lowered the levels of secreted Aβ42 while having a much smaller effect on Aβ10 and Aβ18 (Fig. 5B). The timosaponins’ preferential effects on Aβ42, over the shorter forms of Aβ species, resemble the results of γ-secretase modulation.4,7 Currently, identification of novel GSM is of considerable interest in the development of AD therapeutics, because Aβ42 oligomers or fibrils are considered to be the most toxic Aβ species in AD pathology.

GSM are expected to selectively act on the APP complexes, without inhibiting the cleavage activity of γ-secretase on other physiological substrates.6,7 To further investigate the specificity of the impact of timosaponins on γ-secretase-mediated protein processing, the effects of timosaponins on the γ-secretase-mediated cleavage of the transmembrane receptor Notch1 were examined (Fig. 6). γ-Secretase-catalysed cleavage of Notch releases the Notch intracellular domain (NICD), which regulates developmental gene transcription.22 In cells expressing a Notch1 variant containing transmembrane and intracellular domains (NotchΔE), the NICD is constitutively present due to γ-secretase activity (Fig. 6). Treatment of cells with a γ-secretase inhibitor DAPT completely blocked the production of NICD, while treatment of cells with timosaponins at concentrations that effectively lower Aβ levels did not affect NICD levels. Thus, the timosaponins selectively interfere with Aβ production without altering Notch1 processing.

**Tamosaponins stimulate neurite outgrowth.** Intriguingly, treatment of cells with SSG (1) and timosaponins (2–5) also markedly stimulated neurite outgrowth at concentrations that lower Aβ production, as revealed by type III β-tubulin immunostaining (Fig. 7). The neurite outgrowth stimulation was not shared by other steroidal saponins (e.g., 8–12) or SSG analogues (e.g., 6, 7, 13, 14, 20, and 24) investigated in this study. Stimulation of neurite outgrowth is considered to be a favourable pathology.

**APP as a potential molecular target of timosaponins.** Taken together, our data demonstrate that timosaponins modulate APP processing with a suppression of β-cleavage and selective reduction in Aβ42 production. To establish a possible binding model of the molecular targeting of Aβ, molecular simulation of timosaponins to APP was performed. We postulate that timosaponin binds to APP because there is evidence for interactions between a number of GSM and APP, particularly at its transmembrane region that contains the sites of γ-cleavage (Fig. 8A).23 We employed a model of an APP fragment (Protein Data Bank ID: 2LP1) that spans the extracellular juxtamembrane and transmembrane domains (TMD) (Fig. 8). Previous NMR analyses indicate that the extracellular amino terminus includes a surface-embedded “N-helix” followed by a short “N-loop” connecting to the TMD.20 Importantly, a binding pocket for cholesterol, centred around the N-helix/N-loop/TMD structural element, has been identified (Fig. 8A).20 Our preliminary molecular docking analysis revealed that SSG and timosaponins can be positioned within the cholesterol binding pocket (ESI Fig. S1†). High level hybrid quantum mechanics/molecular mechanics (QM/MM) calculation (ESI†) was performed to provide in depth understanding on the timosaponin binding to the transmembrane domain of APP (Fig. 8B and C). The results imply that the binding interaction is selective, which is probably due to the specific polarity of the SSG-aglyconed timosaponins (Fig. 8B and C). The parent SSG part of TAIII containing the lipophilic steroid ring is surrounded by a group of hydrophobic residues (689–692, 695, 696, 704, 705, 707–712), while the polar part of TAIII possessing the hydrophilic galactose moiety is surrounded by several polar residues (697–699) and lies at the surface of the transmembrane. In particular, the binding pose of the SSG motif is proximal to the GXXXG motifs (particularly, the G750AIG754) of the TMD. These motifs have been shown to be important in the production of the long and short forms of the Aβ polypeptides mediated by γ-secretase. In addition, these motifs are involved in the non-steroidal anti-inflammatory drug (NSAID)-derived GSM modulation of γ-secretase.26,27 Nonetheless, further experiments (e.g., crystallography) are needed to validate the specific binding mode. It is noteworthy that the

**Fig. 6** Effects of timosaponins on Notch cleavage by γ-secretase. N2A-APPswe cells were transfected with myc-tagged NotchΔE construct, which is constitutively cleaved by γ-secretase to generate NICD. Cells were then treated with DMSO control (C); the indicated concentrations of timosaponins or DAPT (as a positive control for γ-secretase inhibition) for 18 h and the expression of NotchΔE and NICD were examined by immunoblot analysis.

**Fig. 7** Effects of timosaponins and SSG on neurite outgrowth of Neuro-2A cells. Cells were treated with DMSO (control), TAIII (5 μM), TAI (10 μM), AA (10 μM), SSG (25 μM) and TAV (10 μM) for 18 h, stained with a monoclonal antibody raised against type III β-tubulin and examined by fluorescence microscopy.
modulation of APP processing by some endogenous steroid-like compounds from animals and plants has been recently reported.\textsuperscript{38,39}

\textit{In vivo} \(\beta\)-lowering activities of timosaponins. The \textit{in vivo} \(\beta\)-lowering activities of timosaponins were examined in mice. A group of 3–5 month old C57BL/6 mice was dosed with SSG (1), TAI (2), TAIII (3) or \(\alpha\), \(\beta\)-OH SSG (34) at 100 mg kg\(^{-1}\) by oral gavage for three times in 2 days. The results showed that these compounds elicited a reduction of \(\beta\)-42 levels in the brain (77% \(\pm\) 4% for SSG, \(P < 0.05\); 83% \(\pm\) 13% for TAI, \(P < 0.05\); 87% \(\pm\) 14% for TAI, \(P = 0.09\); 87% \(\pm\) 15% for 34; \(P = 0.09\)) (Fig. 9). Such a moderate degree of \(\beta\)-42 reduction has been demonstrated by many \(\beta\)-lowering natural compounds (ESI Table S1†). We also determined the levels of SSG 1, timosaponins 2–3, compound 34 and their metabolites in the plasma and brain of the mice by ultra-performance liquid chromatography tandem mass spectrometry. The timosaponins and/or their deglycosylated products (TAI and SSG) at low micromolar concentrations could be detected in the plasma and brain at the end of the experiments (ESI Table S2†). No further metabolites except the deglycosylated products (TAI, SSG from TAIII; SSG from TAI) were detected. The reason for the scarcity of TAIII in the brain is uncertain but may be attributed to a lower ability of the glyco-sylated compounds to cross the blood brain barrier and/or an elevated glycohydrolase activity in the neurons.\textsuperscript{40} Compound 34 was present at a much lower level in the mouse plasma and brain compared to its closest analogue TAI, suggesting that the nature of C3 linkage may have an impact on the bioavailability and/or tissue distribution. Further pharmacokinetics studies of SSG and the timosaponins and their derivatives are required to elucidate the structural features of the timosaponins required for optimal bioavailability and efficacy. Collectively, our results reveal that SSG and certain timosaponins display oral bioavailability, brain penetration capacity and \(\beta\)-lowering activity \textit{in vivo}.

\textbf{Conclusion}

The timosaponins investigated in the current study are preferentially able to lower \(\beta\)-42 production and stimulate neurite outgrowth, largely due to the presence of the effective aglycone SSG 1. They contain structural features including a cis-fused AB ring, 3\(\beta\)-configuration and an intact F-spiroketal ring with a 25\(S\)-configuration. These characteristics are indispensable

Fig. 8  APP transmembrane domain as a potential molecular target of timosaponins. (A) An overview of the portion of APP that is cleaved by \(\alpha\), \(\beta\), and \(\gamma\)-secretases. Also shown are the juxtamembrane and transmembrane regions harbouring binding sites for NSAID-like GSM\textsuperscript{24} and cholesterol.\textsuperscript{28} Numbering is according to the full length of APP770. (B) QM/MM calculations of TAI binding to the transmembrane domain of APP. The surface representation of the transmembrane region of APP, showing the calculated binding pose of TAIIII. (C) The hydrophobic residues and polar residues around TAIIII are portrayed with stick display mode. Carbon atoms of hydrophobic residues are highlighted in green and carbon atoms of polar residues are highlighted in light blue. TAIII is represented by sticks with the carbon atoms in yellow. Colour code: carbon (yellow, green or light blue), nitrogen (dark blue), oxygen (red), and hydrogen (white).

Fig. 9  \textit{In vivo} \(\beta\)-42-lowering activities of SSG and timosaponins. 3–5 month old C57BL/6 mice were dosed with vehicle (C), TAI, TAIII, SSG or \(\alpha\), \(\beta\)-OH SSG (34) at 100 mg kg\(^{-1}\) by oral gavage for three times in 2 days. The levels of \(\beta\)-42 in the mouse brain were determined by ELISA. The number (n) of animals for vehicle control, TAI, TAIII and SSG = 10; for 34, n = 6. Data represent means \(\pm\) standard deviations. Statistical significance in differences between vehicle control and treatment groups was determined by Student’s test, #, \(P < 0.1\); *, \(P < 0.05\).
structural requirements for the compound’s dual properties. The Aβ-lowering activities of “SSG-aglycone” timosaponins are generally associated with decreases in β-cleavage and/or increases in γ-cleavage of APP. These are accompanied by a preferential reduction of Aβ42 levels without affecting the processing of other γ-secretase substrates, resembling the action of GSM. Thus, the “SSG-aglycone” timosaponins are novel agents that modulate APP processing and subsequently lower Aβ production. We have also shown here that some timosaponins and SSG exhibit Aβ-lowering activities in vivo. It is envisaged that, when properly modified and formulated, timosaponins will be intriguing compounds for the development of AD therapeutics.

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Notes and references


