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Pregnenolone derivatives for the treatment of Alzheimer's disease: synthesis, and *in vitro* inhibition of amyloid β_{1-42} peptide aggregation, acetylcholinesterase and carbonic anhydrase-II[†]

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The amyloid state, which is a specific conformation of proteins, offers valuable information about both functional protein structures and the pathological assemblies associated with various diseases. One of the major hallmarks of Alzheimer's disease includes primarily the extracellular build-up of a peptide known as amyloid- β , which has a sequence consisting of 39 to 42 amino acid residues, and the formation of intracellular neurofibrillary tangles mostly consisting of hyperphosphorylated tau protein. Drugs that are expected to reduce A β production, prevent A β aggregation, and promote A β clearance are promising approaches for treating AD. Current work is focused on identifying the compounds that have balanced even mild biological activities against multiple targets instead of finding one-target compound with high potency. We synthesized pregnenolone derivatives and evaluated their potential against inhibition of eeAChE/eqBChE, hCA-II and self-mediated A β_{1-42} peptide aggregation. Our synthesized derivatives 23, and 25–27 exhibited concomitant inhibition of all the tested macromolecular targets. All the active compounds were found to be BBB penetrants in the PAMPA assay. Furthermore, these selected compounds were found to be non-neurotoxic in the MTT assay on neuroblastoma SH-SY5Y cells. Docking studies support dual binding site (PAS and CAS) inhibition of AChE which showed A β_{1-42} aggregation and AChE inhibition. Moreover, docking studies carried out on the 3D crystallographic structure of A β_{1-42} peptide (PDB ID = 1YT) showed significant interactions with amino acid residues Asp 23 and Lys 28, and hydrophobic interactions with the Phe19, Phe20, and Ala 30 effectively impeding the formation of β -sheet structures.

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Introduction

The amyloid state, which is a specific conformation of proteins, offers valuable information about both functional protein structures and the pathological assemblies associated with various diseases.¹ Amyloids are characterized by their insoluble, fibrillary deposits consisting of β -sheet secondary structures.² These structures are observed in numerous diseases and tend to accumulate as plaques and inclusions with advancing age. Notable examples of key proteins that form amyloid fibrils in specific diseases include A β in Alzheimer's disease (AD), huntingtin in Huntington's disease, and α -synuclein in Parkinson's disease.³ AD, a neurodegenerative condition, is

characterized by the accumulation of a peptide called Amyloid- β_{1-42} peptide.^{4,5}

Compared to other fragments formed when amyloid precursor protein (APP) is sliced, A β is chemically "stickier". Gradually, it builds up into tiny amyloid plaques, which are recognized as a hallmark of an Alzheimer's-affected brain.⁶ First, the fragments form tiny clusters known as oligomers, followed by chains of clusters known as fibrils, and finally, "mats" of fibrils known as beta-sheets. Plaques, the last stage, are composed of various substances and beta-sheet clumps. These phases of beta-amyloid accumulation consequently disrupt neuronal function, interfere with cell-to-cell communication and trigger immunological responses, as per the amyloid hypothesis. The brain cells are ultimately destroyed when inflammation is brought on by these immune cells. Thus, the two main pathogenic AD mechanisms are plaque deposition and massive amyloid beta (A β) aggregates.⁷⁻⁹

The complex mechanisms responsible for A β aggregation involve genetic, environmental, and metabolic factors. As amyloid plaques are a hallmark of AD because the aggregate

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formation of A β sets off a series of actions that eventually cause brain cell degradation and cognitive loss as well as memory decline.¹⁰

Scientists are devoting an enormous amount of time and energy to finding out the potential therapeutic approaches that stop beta-amyloid aggregation as a possible therapy for the condition.^{11,12} Amyloid aggregation inhibitors, including poly-oxometalates (POM), metallohelices, nanomaterials, and peptides, have also been rationally designed. However, there are still hazards associated with BBB permeability alterations, even with the effectiveness of therapeutic medicines in brain uptake.^{13,14} Additionally, they are searching for strategies to stop the first contact between A β and nerve cells that cause toxicity since some data indicates that the harmful effects of beta-amyloid happen before plaques and oligomers develop.¹⁵⁻¹⁷ Alvarez in 1997 and Inestrosa in 1996 suggested that the "peripheral anionic site" (PAS) of AChE is involved in the aggregation and formation of toxic A β oligomers. The 42-mer peptide, A β ₁₋₄₂, is the more toxic among these two (*i.e.*, A β ₁₋₄₂ and A β ₁₋₄₀).¹⁸⁻²⁰ Therefore, blocking the PAS could serve to reduce A β aggregation in order to increase the chance to prevent harmful effects. Moreover, a number of authors studied the effect of AChE PAS inhibitors (propidium and fasciculin) and active site inhibitor (edrophonium) on A β aggregation process experimentally.²¹⁻²⁵ Therefore, small molecules that can inhibit the PAS of AChE may also prevent A β from aggregating.²⁶ Moreover, the A β -dependent toxicity is elevated by the AChE-A β and it depends on the bulk of complexes that are formed. The amyloidogenic pathway is one of the pathways for the breakdown of amyloid precursor protein or APP.^{27,28} Beta secretase which is also known as BACE-1 and gamma secretases are responsible for this commonly named amyloidogenic pathway. So, by inhibiting the BACE-1 and gamma secretases, A β levels can be degraded.^{29,30}

Zinc-containing metallo-enzyme carbonic anhydrase (CA) was initially discovered in cow RBC in 1933.^{31,32} A number of investigations provided fresh insights into the potential of carbonic anhydrases (CAs) as therapeutic targets for Alzheimer's disease (AD). In prokaryotes and eukaryotes, carbonic anhydrases (CAs) are zinc-bearing metalloenzymes that initiate the reversible reaction between carbon dioxide and bicarbonate ions.³³ The zinc ion at the CA active site is surrounded by hydroxide ions (OH) and possesses the same catalytic activity.^{34,35} Three histidine residues (His 94, His 96, and His 119) as well as side-chain residues that make coordination bonds with the zinc ion are found in the CA enzyme.³⁶ When developing CA inhibitors, the design of the inhibitors that target hCA-II is crucial. The main sulfonamide group (RSO₂NH₂)-containing compounds make up the foundation of all CA inhibitors (hCA-II). Sulfonamide hCA-II inhibitors, including acetazolamide, methazolamide, ethoxzolamide, dichlorphenamide, and others, have demonstrated significant inhibition against the majority of human hCA isoforms at generally low nanomolar doses. So far, a number of compounds have been created to inhibit different isoforms of carbonic anhydrase.³⁷ Clinically available carbonic anhydrase inhibitors, such as acetazolamide (AZA), dorzolamide, methazolamide,

brinzolamide, ethoxzolamide, diclofenamide, indisulam, zonisamide, *etc.*, are used to treat a variety of conditions, including ocular hypertension, glaucoma, and epilepsy.³⁸⁻⁴¹

Steroids include various structurally related compounds primarily originating from plant and animal kingdoms.⁴² Perhydro-cyclopentanophenanthrene is the common structural nucleus present in this class of compounds.⁴³ Steroids either natural or semi-synthetic have been involved in the handling of inflammatory, immunology, infectious and metabolic diseases.⁴⁴ In 2004, Abbate *et al.*, reported hCA-I, II and IX inhibitions by Estrone-3-O-sulfamate (EMATE, 1).⁴⁵ In 2014, the inhibition mechanism of human carbonic anhydrase-II by bile acid (2) was first explored (see Fig. 1).

In 2018 Nocentini *et al.* reported the various carbonic anhydrase isoforms inhibition of 22 steroids having pendants and functional groups of sulfonates, phenols and carboxylates. Among these 22 bile acids/stEROIDS, hyocholic acid (3, $K_i = 38.9 \mu\text{M}$), α -estradiol (4, $K_i = 40.4 \mu\text{M}$), deoxycholic acid (5, $K_i = 51.0 \mu\text{M}$), and oestrone ($K_i = 50.8 \mu\text{M}$) showed good CA-II inhibition.⁴⁶ A. Khalid *et al.* reported that four recognized bases sarcosine, sarcodine, sarcosine, and alkaloid-C as well as a novel steroid alkaloid, isosarcodine 6, were extracted from extract (MeOH) of *Sarcococca saligna*.⁴⁷ After AChE and butyrylcholinesterase (BuChE) inhibition experiments were directed on these compounds, isosarcodine was discovered as a non-competitive inhibitor of AChE (having $K_i = 21.8 \mu\text{M}$). As, carbonic anhydrases' catalytic activity is inhibited by sulfonamides. Compound 7 was reported as a potent CA IX inhibitor by Cecchi, Alessandro *et al.*⁴⁸ In 2022, compound 8 was reported by Peschiulli *et al.* It is an oral bioavailable and highly effective (hA β ₁₋₄₂ cell $IC_{50} = 1.3 \text{ nM}$) molecule that produced a persistent decrease in A β in mouse and dog animal models (see Fig. 1).⁴⁹

Pregnenolone (PREG) is a neurosteroid synthesized from cholesterol in both mammals and invertebrates.⁵⁰ Neurosteroids are known for their rapid activity, as they are synthesized within the brain and for the brain. Within the brain, pregnenolone and its various metabolic derivatives (pregnenolone sulfate, allopregnanolone, and dehydroepiandrosterone), have demonstrated the ability to improve learning and memory, alleviate depression, and regulate cognitive functions.⁵¹

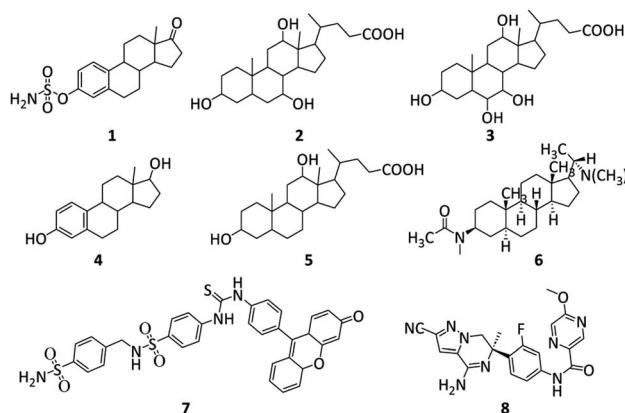


Fig. 1 Structures of reported AChE, A β and CA inhibitors.⁴⁵⁻⁴⁹



Moreover, these compounds play a neuroprotective role in addressing neuroinflammatory diseases like AD and multiple sclerosis (MS), as well as in managing neuropsychiatric disorders such as schizophrenia, depression, and autism.^{29,52}

Our research group is involved in the structural modification of steroid nuclei for various disease treatments. Recently, we reported structural modification of pregnenolone at C-3 and C-17 positions.⁵³ Excellent *in vitro* inhibition results were exhibited by Pyrimidine derivatives of pregnenolone for human dihydrofolate reductase (hDHFR). In another report, we have also synthesized C-3 pregnenolone dihydropyrimidine derivatives for the breast cancer treatment.⁵⁴ In the current research, we planned to modify the pregnenolone core at C-3 and C-17 positions. 4-(Sulfamoylphenyl)thiourea moiety was incorporated at position C-3. While α -bromination of the acetyl group present at position C-17 resulted in the synthesis of various amine derivatives.

Materials and methods

General

The purified reagents/solvents were acquired from viable vendors (standard) and were employed. From Sigma Aldrich and Alfa-Aesar, we have purchased the starting materials. Tetramethyl silane (TMS, internal reference) was employed for recording ^1H and ^{13}C -NMR spectra by utilizing deuterated solvents at 400 MHz and 100 MHz on a Bruker spectrometer. Chemical shifts are specified in δ scale (ppm). TLC was utilized to investigate the progress of all the reactions that is 2.0×5.0 cm aluminium sheets (having silica gel 60F²⁵⁴ on it and 0.25 mm thickness). The purity of the synthesized compounds was analysed by using Shimadzu HPLC system with C18 RP column. The mobile phase was composed of acetonitrile/water (30 : 70 v/v). The system was run in an isocratic mode with at a 1.0 mL min^{-1} flow rate. The compounds with a purity greater than 95% were used for bioevaluation studies. The CHN elemental analysis was carried out by utilizing the LECO-932 CHN analyzer (LECO Corporation, USA).

General method for synthesis of 1-((8S,9S,10R,13S,14S)-3-amino-10,13-dimethyl-2,3,4,7,8,9,10,11,12,13,14,15,16,17-tetradecahydro-1H-cyclopenta[a]phenanthren-17-yl)ethan-1-one (12)

In a round bottom flask, 25 mmol of pregnenolone and 25.5 mmol of *p*-toluene sulfonyl chloride in 30 mL dichloromethane (CH_2Cl_2), trimethylamine (Et_3N , 30 mmol) were added and heat under reflux condition. TLC was utilized to monitor the completion of the reaction. After the completion of the reaction, ice water was poured into it and stirred. The resulting aqueous solution was extracted from ethyl acetate (3×20 mL). Finally, saturated NaHCO_3 was used for washing the combined organic layer and anhydrous NaSO_4 was utilized for drying. The organic layer was removed and the resulting tosylate **11** was dried and used for the next step without any further purification.

The tosylate **11** (20 mmol) was suspended in acetone. Then ammonium acetate (40 mmol) was added and at room temperature for 3–4 hours mixture was stirred. The resulting precipitates were filtered off, diethyl ether was used for washing and drying to afford compound **12**.

Yield = 70%; ^1H NMR (400 MHz, CDCl_3) δ 5.25 (brs, 1H), 4.54 (d, $J = 6.0$ Hz, 2H), 3.74 (t, $J = 5.6$ Hz, 1H), 2.55 (t, 1H), 2.45–2.37 (m, 3H), 2.32–2.27 (m, 1H), 2.21–2.18 (m, 1H), 2.14 (s, 3H), 2.10–2.02 (m, 2H), 1.90–1.86 (m, 1H), 1.77–1.67 (m, 4H), 1.66–1.63 (m, 1H), 1.59–1.52 (m, 1H), 1.50–1.42 (m, 2H), 1.35–1.28 (m, 1H), 1.25 (s, 3H), 1.18–1.14 (m, 2H), 0.68 (s, 3H). ^{13}C NMR (100 MHz, CDCl_3) δ 208.5, 139.2, 120.9, 64.5, 53.8, 51.8, 48.2, 44.7, 41.2, 37.9 (2C), 34.4, 32.5 (2C), 30.5, 29.3, 26.6, 23.8, 21.9, 19.3, 15.9. Analysis calculated for $\text{C}_{21}\text{H}_{33}\text{NO}$: C, 79.95; H, 10.54; N, 4.44; O, 5.07. Observed: C, 80.04; H, 10.52; N, 4.46.

4-((8S,9S,10R,13S,14S)-17-Acetyl-10,13-dimethyl-2,3,4,7,8,9,10,11,12,13,14,15,16,17-tetradecahydro-1H-cyclopenta[a]phenanthren-3-yl)thioureido) benzenesulfonamide (15)

To a stirred solution of **12** (20 mmol) in 30 mL acetone added 20 mmol *o*-(*p*-tolyl)chlorothionoformate **13**. For 2 h, the mixture was kept at continuous stirring at RT. The TLC technique was utilized to check the completion of the reaction. The mixture was filtered to obtain intermediate **14**. In the second step, crude intermediate **14** in 5 mL 1,4-dioxane was added 8 mmol of sulfanilamide followed by 16 mmol triethylamine. It was then stirred at 60 °C for 3 h. The resulting precipitates, after that diethyl ether was employed for washing, and dried to afford compound **15**.

Light yellow powder. Yield = 61%; R_f = 0.38 (*n*-hexane/EtOAc 8 : 1); HPLC purity = 98.9%, t_r = 5.78 min, ^1H NMR (400 MHz, CDCl_3) δ 8.25 (s, 1H, $-\text{NH}$), 7.90 (d, 2H, $J = 8.4$ Hz, ArH), 7.74 (d, 2H, $J = 8.4$ Hz, ArH), 6.76 (s, 2H, NH_2), 6.51 (s, 1H, NH), 5.27–5.21 (m, 1H), 3.73 (t, 1H, $J = 5.6$ Hz), 2.59–2.54 (m, 1H), 2.41–2.36 (m, 1H), 2.28–2.26 (m, 1H), 2.24–2.21 (m, 1H), 2.15 (s, 3H), 2.10–2.02 (m, 2H), 1.82–1.76 (m, 1H), 1.74–1.66 (m, 4H), 1.53–1.46 (m, 6H), 1.29–1.26 (m, 1H), 1.22 (s, 3H), 1.09–1.04 (m, 2H), 0.69 (s, 3H). ^{13}C NMR (100 MHz, CDCl_3) δ 208.5, 180.6, 141.1, 140.4, 136.1, 125.2 (2C), 122.1 (2C), 63.2, 53.2, 50.2, 48.2, 43.8, 38.5, 35.9, 33.3 (1C), 32.1, 31.5 (2C), 28.1, 26.4, 24.3, 20.5, 18.3 (2C), 15.6 (2C). Analysis calculated for $\text{C}_{28}\text{H}_{39}\text{N}_3\text{O}_3\text{S}_2$: C, 63.48; H, 7.42; N, 7.93; O, 9.06; S, 12.10. Observed: C, 63.36; H, 7.44; N, 8.78.

The general method for the synthesis of α -brominated derivative **16**

A solution of 3-amino pregnenolone derivative (**15**, 10 mmol) in 25 mL of chloroform was stirred. To this solution, bromine (10 mmol) in CHCl_3 was added drop-wise by using a dropping funnel. After that, it was stirred for 15–20 minutes at 50 °C and cooled to RT. The final crude product was recrystallized from acetic acid.

Light brown powder. Yield = 79%; R_f = 0.36 (*n*-hexane/EtOAc 8 : 1); ^1H NMR (400 MHz, CDCl_3) δ 8.24 (s, 1H, $-\text{NH}$), 7.92 (d, 2H, $J = 8.48$ Hz, ArH), 7.74 (d, 2H, $J = 8.48$ Hz, ArH), 6.77 (s, 2H, NH_2), 6.51 (s, 1H, NH), 5.26–5.20 (m, 1H), 3.71 (t, 1H, $J = 5.6$ Hz).



Hz), 2.57–2.53 (m, 1H), 2.41–2.36 (m, 1H), 2.28–2.20 (m, 2H), 2.19 (s, 3H), 2.12–2.06 (m, 2H), 1.82–1.76 (m, 1H), 1.72–1.68 (m, 2H), 1.55–1.48 (m, 4H), 1.46–1.43 (m, 2H), 1.27–1.24 (m, 1H), 1.20 (s, 3H), 1.12–1.06 (m, 2H), 0.71 (s, 3H). ^{13}C NMR (100 MHz, CDCl_3) δ 204.3, 181.8, 141.5, 140.7, 135.2, 129.4 (2C), 122.5 (2C), 61.3, 54.5, 52.6, 49.3, 45.6, 39.0, 37.2, 36.5, 36.1, 35.4, 33.6, 29.7, 27.1, 25.3, 23.8, 20.4, 18.2 (2C), 13.3. Analysis calculated for $\text{C}_{28}\text{H}_{38}\text{BrN}_3\text{O}_3\text{S}_2$: C, 55.25; H, 6.29; Br, 13.13; N, 6.90; O, 7.89; S, 10.53. Observed: C, 55.36; H, 6.24; N, 6.88.

The general method for the synthesis of derivatives 23–28

To a solution of α -brominated **16** (5 mmol) in DMF, 5.25 mmol of trimethylamine (Et_3N) and 7.5 mmol of amines (17–22) were added. The mixture was refluxed overnight. Upon completion of the reaction, the pregnenolone derivatives were purified by column chromatography using petroleum ether/ethyl acetate 20 : 1 for compound **23** and 10 : 1 for **24–28**.

4-((8S,9S,10R,13S,14S)-10,13-Dimethyl-17-((4-sulfa-moylphenethyl)glycyl)-2,3,4,7,8,9,10,11,12,13,14,15,16,17-tetradecahydro-1H-cyclopenta[a]phenanthren-3-yl)-thioureido)benzenesulfonamide (23)

Light yellow powder. Yield = 61%; R_f = 0.40 (*n*-hexane/EtOAc 20 : 1); m.p. = 195–197 °C; HPLC purity = 99%, R_t = 6.72 min, ^{1}H NMR (400 MHz, CDCl_3); δ 8.23 (s, 1H, –NH), 7.94 (d, 2H, J = 8.4 Hz, ArH), 7.77 (d, 2H, J = 8.4 Hz, ArH), 7.41 (d, 2H, J = 8.0 Hz, ArH), 7.24 (d, 2H, J = 8.0 Hz, ArH), 6.98 (s, 4H, 2 \times NH₂), 6.54 (s, 1H, NH), 5.35–5.29 (m, 1H), 4.25–4.20 (m, 1H), 3.42 (d, 2H, J = 5.6 Hz, CH₂), 3.33–3.29 (m, 1H), 2.99–2.94 (m, 2H), 2.78 (t, 2H, J = 5.6 Hz, CH₂), 2.73–2.69 (m, 1H, NH), 2.56–2.48 (m, 2H, CH₂), 1.91–1.88 (m, 1H), 1.75–1.69 (m, 1H), 1.58–1.47 (m, 6H), 1.45 (m, 4H), 1.23 (m, 4H), 1.00 (s, 3H), 0.93 (m, 1H), 0.79 (s, 3H). ^{13}C NMR (100 MHz, CDCl_3) δ 204.3, 180.5, 149.3, 141.5, 140.1, 138.0, 137.1, 136.3, 128.4 (2C), 127.8 (2C), 126.8 (2C), 126.0, 124.5, 121.4, 113.4, 77.0, 76.3, 62.3, 58.2, 55.5, 49.7, 48.1, 40.2, 39.8, 37.9, 37.0, 34.4, 28.8, 24.4, 23.4, 21.5, 17.2, 15.8. Analysis calculated for $\text{C}_{36}\text{H}_{49}\text{N}_5\text{O}_5\text{S}_3$: C, 59.39; H, 6.78; N, 9.62; O, 10.99; S, 13.21. Observed: C, 59.45; H, 6.76; N, 9.59.

4-((8S,9S,10R,13S,14S)-10,13-Dimethyl-17-(phenethylglycyl)-2,3,4,7,8,9,10,11,12,13,14,15,16,17-tetradeca-hydro-1H-cyclopenta[a]phenanthren-3-yl)-thioureido)benzenesulfonamide (24)

Off-white powder. Yield = 66%; R_f = 0.46 (*n*-hexane/EtOAc 8 : 1); m.p. = 152–154 °C; HPLC purity = 98%, R_t = 7.5 min, ^{1}H NMR (400 MHz, CDCl_3) δ 8.23 (s, 1H, –NH), 7.93 (d, 2H, J = 8.4 Hz, ArH), 7.75 (d, 2H, J = 8.4 Hz, ArH), 7.24–7.19 (m, 5H, ArH), 6.75 (s, 2H, NH₂), 6.51 (s, 1H, NH), 5.29–5.24 (m, 1H), 4.13 (d, 1H, J = 7 Hz), 3.97–3.95 (m, 2H), 3.89–3.86 (m, 2H), 3.30 (d, 2H, J = 5.2 Hz, CH₂), 2.50–2.35 (m, 5H), 2.22–2.17 (m, 4H), 1.95–1.85 (m, 6H), 1.76–1.69 (m, 2H), 1.62–1.54 (m, 2H), 1.24–1.19 (m, 2H), 0.96 (s, 3H), 0.77 (s, 3H). ^{13}C NMR (100 MHz, CDCl_3) δ 204.8, 180.6, 149.3, 141.3, 138.0, 136.5, 128.3 (2C), 126.6 (2C), 124.5, 122.0, 121.4, 113.4, 76.3, 77.0, 76.3, 62.3, 58.2, 55.5, 50.5, 48.7, 49.7, 43.4, 40.2, 39.8, 37.6, 37.3, 34.4, 32.2, 28.8, 24.4, 23.2, 21.5, 17.6, 15.5. Analysis calculated for $\text{C}_{36}\text{H}_{48}\text{N}_4\text{O}_3\text{S}_2$: C, 66.63;

H, 7.46; N, 8.63; O, 7.40; S, 9.88. Observed: C, 66.70; H, 7.47; N, 8.61.

4-((8S,9S,10R,13S,14S)-17-(Benzylglycyl)-10,13-di-methyl-2,3,4,7,8,9,10,11,12,13,14,15,16,17-tetradeca-hydro-1H-cyclopenta[a]phenanthren-3-yl)thioureido)benzenesulfonamide (25)

Off white powder. Yield = 68%; R_f = 0.48 (*n*-hexane/EtOAc 8 : 1); m.p. = 163–165 °C; HPLC purity = 97%, R_t = 8.8 min, ^{1}H NMR (400 MHz, CDCl_3) δ 8.27 (s, 1H, –NH), 7.96 (d, 2H, J = 8.4 Hz, ArH), 7.75 (d, 2H, J = 8.4 Hz, ArH), 7.53–7.45 (m, 5H, ArH), 6.73 (s, 2H, NH₂), 6.53 (s, 1H, NH), 5.37–5.31 (m, 1H), 4.88 (d, 2H, J = 5.2 Hz, CH₂), 4.12–4.05 (m, 1H), 3.51–3.47 (m, 2H), 2.98 (t, 1H, J = 5.4 Hz), 2.92–2.86 (m, 1H), 2.53–2.54 (m, 4H), 2.27–2.09 (m, 4H), 1.94–1.77 (m, 2H), 1.76–1.61 (m, 2H), 1.59–1.50 (m, 1H), 1.37–1.30 (m, 2H), 1.25–1.21 (m, 1H), 0.95 (s, 3H), 0.89–0.81 (m, 1H), 0.77 (s, 3H). ^{13}C NMR (100 MHz, CDCl_3) δ 204.9, 180.5, 149.3, 141.6, 139.7, 136.8, 129.4 (2C), 127.6 (2C), 123.4, 122.5, 121.0, 113.6, 77.5, 77.1, 76.6, 62.4 (2C), 58.2, 55.5, 50.5, 49.7, 43.4, 39.2, 37.6, 37.1, 34.4, 32.2, 28.8, 24.4, 23.4, 21.5, 17.8, 15.6. Analysis calculated for $\text{C}_{35}\text{H}_{46}\text{N}_4\text{O}_3\text{S}_2$: C, 66.21; H, 7.30; N, 8.82; O, 7.56; S, 10.10. Observed: C, 66.26; H, 7.31; N, 8.81.

4-((8S,9S,10R,13S,14S)-10,13-Dimethyl-17-(phenylglycyl)-2,3,4,7,8,9,10,11,12,13,14,15,16,17-tetradeca-hydro-1H-cyclopenta[a]phenanthren-3-yl)thioureido)benzenesulfonamide (26)

Off white powder. Yield = 59%; R_f = 0.44 (*n*-hexane/EtOAc 8 : 1); m.p. = 183–185 °C; HPLC purity = 99%, R_t = 8.3 min, ^{1}H NMR (400 MHz, CDCl_3) δ 8.24 (s, 1H, –NH), 7.90 (d, 2H, J = 8.0 Hz, ArH), 7.77 (d, 2H, J = 8.0 Hz, ArH), 7.25–7.22 (m, 2H, ArH), 7.18–7.14 (m, 2H, ArH), 7.11–7.08 (m, 1H, ArH), 6.75 (s, 2H, NH₂), 6.55 (s, 1H, NH), 5.26–5.23 (m, 1H), 4.99 (t, 1H, J = 4.8 Hz, NH), 3.99 (d, 2H, J = 4.8 Hz, CH₂), 3.76–3.72 (m, 1H), 2.56–2.47 (m, 2H), 2.37–2.28 (m, 2H), 2.22–2.16 (m, 2H), 1.99–1.88 (m, 1H), 1.77–1.63 (m, 6H), 1.57–1.43 (m, 4H), 1.29–1.15 (m, 5H), 1.10–1.02 (m, 1H), 0.94 (s, 3H). ^{13}C NMR (100 MHz, CDCl_3) δ 204.7, 180.6, 149.0, 141.6, 138.5, 136.8, 129.3 (2), 127.9 (2), 121.2 (2), 113.7 (2C), 77.4, 77.1, 76.8, 57.3, 55.6, 52.2, 50.5, 49.9, 43.4, 39.3, 37.6, 37.0, 34.6, 32.2, 28.8, 24.4, 23.4, 21.0, 17.6, 15.9. Analysis calculated for $\text{C}_{34}\text{H}_{44}\text{N}_4\text{O}_3\text{S}_2$: C, 65.77; H, 7.14; N, 9.02; O, 7.73; S, 10.33. Observed: C, 65.84; H, 7.16; N, 9.00.

4-((8S,9S,10R,13S,14S)-17-(2-(10H-Phenothiazin-10-yl)-acetyl)-10,13-dimethyl-2,3,4,7,8,9,10,11,12,13,14,15,16,17-tetradeca-hydro-1H-cyclopenta[a]phenanthren-3-yl)thioureido)benzenesulfonamide (27)

Yellow powder. Yield = 69%; R_f = 0.49 (*n*-hexane/EtOAc 8 : 1); m.p. = 170–172 °C; HPLC purity = 99%, R_t = 9.43 min, ^{1}H NMR (400 MHz, CDCl_3) δ 8.24 (s, 1H, –NH), 7.91 (d, 2H, J = 8.4 Hz, ArH), 7.77 (d, 2H, J = 8.4 Hz, ArH), 6.91–6.85 (m, 8H), 6.78 (s, 2H, NH₂), 6.55 (s, 1H, NH), 5.27–5.25 (m, 1H), 3.96 (s, 2H, CH₂), 3.77–3.73 (m, 1H), 3.33 (m, 2H), 2.74–2.63 (m, 2H), 2.42–2.32 (m, 1H), 2.15–1.88 (m, 9H), 1.75–1.71 (m, 2H), 1.60–1.51 (m, 3H), 1.36–1.24 (m, 4H), 0.96 (s, 3H). ^{13}C NMR (100 MHz, CDCl_3) δ 207.2, 181.1, 144.5, 140.1, 139.0, 136.2, 128.7, 127.8,



127.8, 127.3, 127.3, 126.9, 126.9, 122.5, 122.4, 122.2, 120.8, 120.8, 116.7, 116.4, 57.8, 56.4, 55.6, 50.4, 49.6, 43.2, 39.2, 37.7, 37.1, 34.6, 32.2, 32.1, 27.7, 24.4, 24.4, 23.5, 21.1, 19.0, 16.1. Analysis calculated for $C_{40}H_{46}N_4O_3S_3$: C, 66.08; H, 6.38; N, 7.71; O, 6.60; S, 13.23. Observed: C, 66.18; H, 6.40; N, 7.69.

4-((8S,9S,10R,13S,14S)-10,13-Dimethyl-17-((1,2,3,4-tetrahydroanthracen-9-yl)glycyl)-2,3,4,7,8,9,10,11,12,13,14,15,16,17-tetradecahydro-1H-cyclopenta[a]phenanthren-3-yl)thioureido)benzenesulfonamide (28)

Yellow powder. Yield = 64%; R_f = 0.55 (*n*-hexane/EtOAc 8 : 1); m.p. = 214–216 °C; HPLC purity = 99%, R_t = 11.20 min, 1H NMR (400 MHz, $CDCl_3$) δ 8.25 (s, 1H, –NH), 7.91 (d, 2H, J = 8.4 Hz, ArH), 7.78 (d, 2H, J = 8.4 Hz, ArH), 7.63–7.58 (m, 2H), 7.17–7.13 (m, 2H), 6.97 (s, 2H, NH₂), 6.52 (s, 1H, NH), 5.32–5.28 (m, 1H), 3.43 (d, 2H, CH_2), 3.38–3.36 (m, 2H), 3.34–3.26 (m, 2H), 2.58–2.46 (m, 3H), 2.34–2.28 (m, 1H), 2.15–1.88 (m, 9H), 1.94–1.78 (m, 9H), 1.71–1.68 (m, 2H), 1.58–1.55 (m, 1H), 1.53–1.49 (m, 2H), 1.42–1.36 (m, 2H), 1.23–1.18 (m, 1H), 0.96 (s, 3H). ^{13}C NMR (100 MHz, $CDCl_3$) δ 209.5, 181.1, 140.5, 140.1, 139.0, 136.2, 134.2, 128.1, 127.9, 127.8(2), 127.2, 126.1, 125.3(2), 125.1, 123.3, 122.2, 120.8(2), 58.1, 55.8, 54.1, 50.4, 49.6, 43.3, 39.2, 37.7, 37.1, 34.6, 32.6, 32.1, 31.0, 29.5, 27.7, 24.9, 24.4, 24.0, 23.5, 21.1, 19.0, 16.1. Analysis calculated for $C_{42}H_{52}N_4O_3S_2$: C, 69.58; H, 7.23; N, 7.73; O, 6.62; S, 8.84. Observed: C, 69.65; H, 7.24; N, 7.71.

***In vitro* CA-II inhibition assay**

CA inhibition studies were performed using human CA-II (Sigma, Catalogue No. C6165) reported assay procedures with little modification.^{55–57}

***In vitro* acetyl/butyrylcholinesterase inhibition assays**

Ellman method is utilized to evaluate the inhibition potential (IC_{50} values) of synthesized compounds and their derivatives against electric eel acetylcholinesterase (eeAChE).^{57,58} Enzymes AChE is used for the hydrolysis of acetylcholine/butyrylcholine iodide which is used as a basis for the principle of this inhibition assay. An anion 5-thio-2-nitrobenzoate is yielded as a result of this hydrolysis which afterwards form a yellow colour complex with Ellman's reagent; 5,5-dithio-bis-2-nitrobenzoic acid (DTNB) solution and that can be determined by utilizing spectrophotometer.

For the preparation of solution, the compound is dissolved separately in a 0.1 M phosphate buffer and different concentrations are formed that can range from 250 to 1000 micrograms per millilitre. To prepare the solution of phosphate buffer with a certain pH, few micrograms per litre of monopotassium phosphate and dipotassium phosphate are mixed in a ratio and for adjusting the pH, potassium hydroxide is used. Then solid AChE/BChE is diluted in a freshly formed phosphate buffer as far as required final concentration is achieved. Subsequently, the solutions of acetylcholine/butyrylcholine and DTNB are prepared in distilled water and these solutions are placed in the refrigerator after enclosing them in an Eppendorf Cap.

Compound and positive control are dissolved in methanol and then further dilutions of compound are prepared.

Self-mediated $A\beta_{1-42}$ aggregation assay

The impact of synthesized compounds on self-mediated $A\beta$ aggregation was evaluated by using a thioflavin T-based fluorometric test.^{10,59–61} Initially pretreated $A\beta_{1-42}$ peptide was dissolved in DMSO to prepare 100 μ M stock solution. The $A\beta_{1-42}$ peptide stock solution was diluted to a concentration of 10 μ M in 150 mM phosphate buffer (pH 7.4) containing 150 mM NaCl prior to the incubations. Next, 10 μ L of the test compounds and 20 μ L of $A\beta_{1-42}$ was combined, and the mixture was applied to the appropriate wells on 96-well plates with black walls. After that, the mixture was quickly diluted with 70 μ L of thioflavin T solution (ThT) to get a final volume of 100 μ L (2 μ M final $A\beta_{1-42}$ concentration). Every sample was run in triplicate. The sealed 96-well plate was placed in a plate reader (Synergy H4) and heated to 37 °C to start the aggregation process. Every 90 seconds, the ThT fluorescence was monitored through the plate's bottom at an excitation wavelength of 440 nm and an emission wavelength of 490 nm. The media was shaken constantly in between measurements to quantify the development of amyloid fibrils. After 1 hour, the $A\beta_{1-42}$'s ThT emission started to grow, reached a plateau after 8 hours, and then stayed essentially unaltered for an additional 16 hours of incubation. The average fluorescence intensities at the plateau were calculated by summing the fluorescence intensities at the corresponding wells at $t = 0$ h, both in the presence and absence of the various doses of test substances (1, 5, 10, 15, 20, 25 μ M). The amount of inhibition of $A\beta_{1-42}$ self-induced aggregation was measured using the formula:

$$(1 - F_i/F_0) \times 100\%,$$

where F_i is the rise in fluorescence of $A\beta_{1-42}$ treated with the test chemicals and F_0 is the increase in fluorescence of $A\beta_{1-42}$ alone (Control).

Cell viability assay test

To evaluate the $A\beta_{1-42}$ toxicity in the presence or absence of designed compounds, we performed the cell viability assays. Promega cat. CellTiter 96 aqueous non-radioactive cell proliferation assay kit was used to perform this assay. To analyze the cytotoxicity of $A\beta$ under various conditions, SH-SY5Y cell lines were used. RPMI 1640 medium (ATCC formulated) was utilized to culture the SH-SY5Y cells supplemented with 10% Fetal bovine serum (FBS). SH-SY5Y cells were plated in 96-well plates at 10 000 cells per well and then it was cultured in 5% carbon dioxide for 20 hours at 37 °C. After that, pre-disaggregated and purified samples of $A\beta$ were solvated in phosphate-buffered buffered saline (PBS) up to an ultimate concentration of 5 micromolars for preparing the solutions of peptide inhibitors and $A\beta$ mixture. Then at various concentrations, different peptide inhibitors were added. A 0.22 micromolar filter was utilized to filter the mixture solution and then without shaking incubate it for 16 hours at 37 °C. In each



well that contains a 90 microlitre medium, a pre-incubated mixture of 10 microliters was added to begin the cell viability assay and then for 24 hours it was incubated in 5% carbon dioxide at 37 °C. Then, in each well a dye solution of 15 microliters was added. After incubating it at 37 °C for four hours, 100 microliters stop mix or solubilization solution was put into a solution. It was followed by further incubation for 12 hours at room temperature and then at 570 nm reading of absorbance was taken along with the background reading at 700 nm. Quadruplicate readings for each sample were collected in parallel. Then, the rate for cell survival was normalized by utilizing the cells treated with PBS at 100% and 0.02 per cent cells treated with SDS at 0% viability.

Docking studies

Molecular operating environment (MOE, 2016.0802) was employed for docking simulations on human carbonic anhydrase II complexed with acetazolamide (PDB accession: 3HS4), AChE complexed with bis-tacrine (PDB accession: 2CKM) and $\text{A}\beta_{1-42}$ (PDB accession: 1IYT) and β -sheet pentamer or $\text{A}\beta_{42}$ protofibril oligomer (PDB ID = 2BEG).

The downloaded enzyme was subjected to 3D protonation and energy minimization for stable conformation. Protonation of synthesized compound was done and after that energy minimization was employed by utilizing parameter at default operation (Gradient 0.00001, Amber 10EHT Force field). After the preparation of enzyme and ligand structures, the reliability of the docking procedure to reproduce the correct binding orientation was determined by the re-docking of the extracted ligand from the crystal structure. The protocol with an RMSD value less than 2 Å was used for docking runs. For the current study, triangle matcher (placement), affinity dG (scoring), GBVI/WSA (re-scoring) protocol was used. Ten conformations for each ligand were generated and top-ranked conformations were selected for analysis. Three-dimensional/two-dimensional ligand interactions were visualized by using MOE interaction plots and discovery studio visualizer.

Results and discussion

Design rational

Structural modification of the steroid nucleus has got considerable attention in recent years. This is because steroid-based therapeutics have several advantages and possess interesting structural features for the treatment of various diseases.

Multitarget-directed ligands (MTDLs) have emerged as a successful strategy for the design of therapeutics for multi-factorial diseases like Alzheimer's disease. The combination of two or more pharmacophoric moieties into a large molecular weight single multipotent entity has been reported to have increased therapeutic advantages and chances of success. Initially, several predictive docking analyses were conducted on target enzyme X-ray crystal structures to anticipate optimum moieties on the left and right sides of pregnenolone. Steroidal cores are complex lipophilic molecules and are responsible for the improvement of BBB permeability. At the C-3 position, 4-(sulfamoylphenyl)thiourea moiety was chosen due to its ability to form hydrogen bond interactions with mid-gorge amino acid residues of AChE and carbonic anhydrase-II. Moreover, the sulfonamide group, also known as the zinc binding group (ZBG), is considered highly important for designing CA inhibitors. We then pursued exploring the influence of different amines with diverse sizes (tricyclic/monocyclic systems) and linker lengths (17–22) in the C-17 acetyl group (Fig. 2). This iterative rational approach allows for designing a pregnenolone-based framework that enhances cholinesterase, carbonic anhydrase and $\text{A}\beta$ aggregation inhibition.

Chemistry

Synthesis of the 3-amino derivative of pregnenolone **12** is outlined in Scheme 1. Pregnenolone **9** was reacted with *p*-toluene sulfonyl chloride **10** to give tosylate **11**. For the synthesis of C-3 amine derivative of pregnenolone,

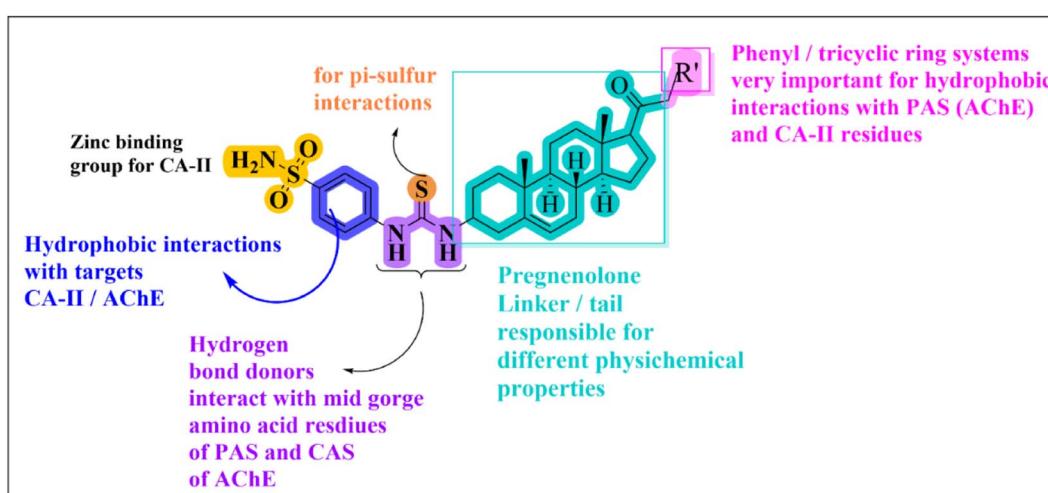
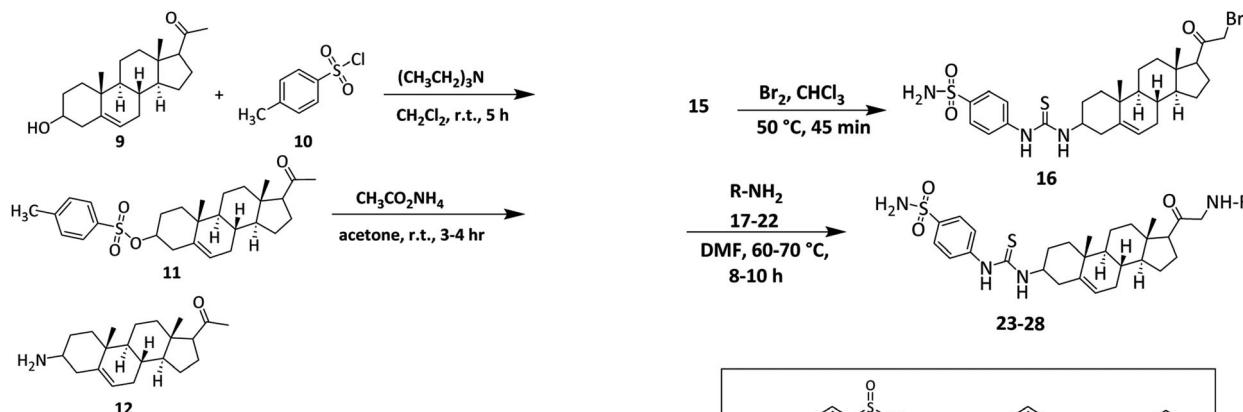
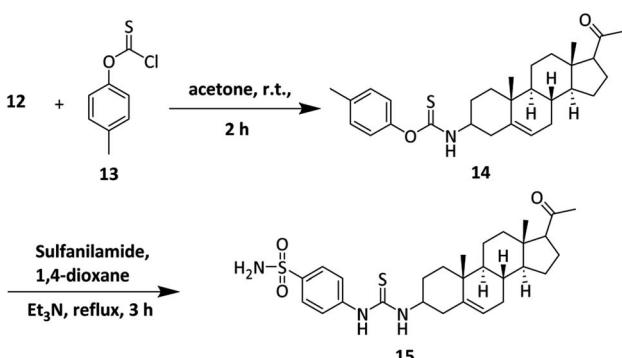


Fig. 2 Design rationale for current research.





Scheme 1 Synthesis of 3-amino derivative of pregnenolone.



Scheme 2 Synthesis of sulfonamide-thiourea adduct 15.

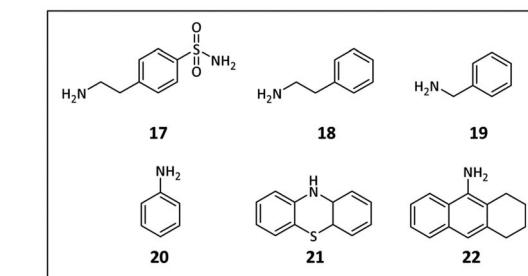
ammonium acetate was used. Matloobi *et al.* used ammonium acetate in DMSO for the introduction of amine. In the current study, we used acetone instead of DMSO.^{62,63} Tosylate 11 was further reacted with ammonium acetate in acetone to give the target compound 12.

The synthesis of sulfonamide derivative 15 is outlined in Scheme 2. Here, we used *o*-(*p*-tolyl)chlorothionoformate 13 for the synthesis of sulfonamide-thiourea adduct 15. Reaction of 3-amino derivative of pregnenolone 12 with *o*-(*p*-tolyl) chlorothionoformate 13 in acetone at room temperature stirring gave intermediate 14 in good yield. The synthesized intermediate finally reacted with sulfanilamide in 1,4-dioxane solvent under reflux conditions to give product 15.

Finally, the synthesis of target sulfonamide derivatives 23–28 is outlined in Scheme 3. The acetyl group was brominated by using bromine in chloroform.⁶⁴ The bromo-compound was used for the next step without any further purification. Compound 16 was reacted with various amine derivatives 17–22 in DMF to obtain target compounds 23–28.

In vitro pharmacology

In vitro human CA-II inhibition activities. The inhibition of pregnenolone derivatives 23–28 against CA-II was carried by using the spectrophotometric method described by Pocker and Meany.⁵⁵ The results are shown in Table 1. Acetazolamide was



Scheme 3 Synthesis of target compounds (23–28) of pregnenolone.

used as a reference standard drug. The inhibition potency of pregnenolone and its 3-amino derivative 12 was also accessed. Inhibition potency results revealed that the presence of amine at the 3-position of pregnenolone showed a little enhanced potency ($IC_{50} = 73.19 \mu M$) than 3-OH in pregnenolone (9, $IC_{50} = 89.38 \mu M$). While the thiourea derivative 15 displayed IC_{50} value of $48.91 \mu M$. Compound 23 containing di-sulfonamide groups emerged as a good inhibitor of hCA-II ($IC_{50} = 0.67 \pm 0.09 \mu M$). All the other synthesized derivatives showed IC_{50} values in the low micromole range (Table 1).

In vitro inhibition of cholinesterases. Ellman's method was employed to evaluate the potential inhibition of pregnenolone derivatives against both cholinesterases (AChE and BChE). The results of inhibition potential in terms of IC_{50} values in micromole (μM) and their selectivity results are summarized in Table 1. Pregnenolone 9, its amine derivative 12 and 4-(sulfamoylphenyl)thiourea derivative 15 was unable to show the inhibition of both cholinesterases. 4-(sulfamoylphenyl)thiourea derivative 23 with substitution at C-3 position showed excellent inhibition in nanomolar concentration with $IC_{50} = 0.04 \mu M$ (40 nM). The derivative also showed good inhibition of BChE having an IC_{50} value of $0.58 \mu M$ (SI = 14.5).

Self-mediated A β _{1–42} aggregation inhibition

The selected synthesized compounds were used in the Thioflavin-T (ThT) fluorescence assay test to evaluate the effect on A β aggregation.¹⁰ For that purpose, each compound with one and half equivalent concentrations (1 : 0.5 and 1 : 1) was employed with A β _{1–42} peptide. Using ThT, a fluorescent dye that attaches specifically to fibrous structures, we were able to track the development of A β aggregation over time in the presence of certain compounds (15, 23, 26 and 27). Control



Table 1 Results of human carbonic anhydrase II, eeAChE, eqBChE and amyloid β_{1-42} peptide aggregation inhibitions

Compd no.	Structures	IC ₅₀ ^a (μM) ± SEM			Amyloid β_{1-42} peptide aggregation inhibition ^c
		hCA-II	eeAChE	eqBChE	
9		89.38 ± 1.21	73.18 ± 0.83	<100	— NA ^d
12		73.19 ± 1.14	67.67 ± 1.09	<100	— NA
15		48.91 ± 1.37	12.98 ± 0.18	63.02 ± 1.23	4.8 9.25 ± 0.17
23		0.67 ± 0.04	0.043 ± 0.001	0.58 ± 0.03	14.5 13.08 ± 0.16
24		5.44 ± 0.14	1.61 ± 0.10	12.84 ± 0.45	8.0 29.64 ± 1.21
25		1.02 ± 0.01	0.98 ± 0.10	11.27 ± 0.11	11.5 12.84 ± 0.15
26		1.56 ± 0.02	0.12 ± 0.01	2.84 ± 0.04	23.6 1.04 ± 0.01



Table 1 (Cont'd.)

Compd no.	Structures	IC ₅₀ ^a (μM) ± SEM				Amyloid β_{1-42} peptide aggregation inhibition ^c
		hCA-II	eeAChE	eqBChE	SI ^b	
27		5.18 ± 0.12	0.094 ± 0.002	15.01 ± 0.03	166.7	5.78 ± 0.07
28		31.83 ± 1.08	0.32 ± 0.01	13.50 ± 0.09	42.2	26.37 ± 0.25
Acetazolamide		0.028 ± 0.001	—	—	—	—
Donepezil		—	0.054 ± 0.001	—	—	—
Curcumin		—	—	—	—	7.29 ± 0.03

^a Values represent mean ± SEM; *n* = 3. ^b Selectivity index = IC₅₀ of BChE/IC₅₀ of AChE. ^c The A β_{1-42} was incubated in the absence or presence of different concentrations of the inhibitors (pregnenolone derivatives). ^d NA = no activity found in tested concentrations; yellow highlighted boxes represent the compounds that have balanced (moderate to excellent) biological activities against all tested target.

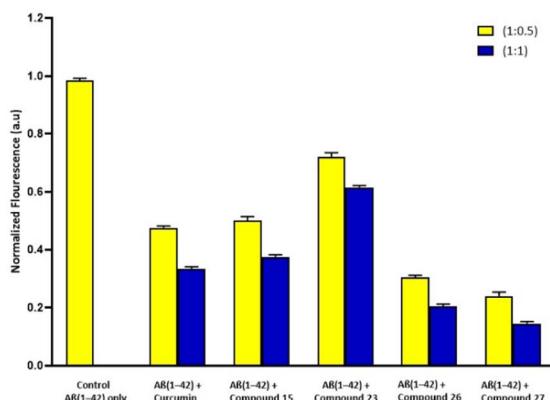


Fig. 3 Bar diagram illustration of the effect of curcumin and compounds (15, 23, 26 and 27) on ThT fluorescence and aggregation of $\text{A}\beta_{1-42}$ peptide after incubated for 24 hours.

experiment $\text{A}\beta_{1-42}$ alone in the ThT research demonstrated an increase in fluorescence intensity, but reduced fluorescence intensity was observed when $\text{A}\beta_{1-42}$ was incubated with synthesized derivatives and curcumin (standard drug/positive control). All of the synthesized derivatives have the potential to interact with the $\text{A}\beta$ peptide and prevent self-aggregation, based on the findings of the experiments. Initially, the fluorescence intensity of $\text{A}\beta_{1-42}$ was evaluated independently. The percentage of aggregates for the samples treated with synthesized compounds was evaluated by taking the $\text{A}\beta_{1-42}$ aggregation alone as 100%. The samples treated with curcumin, compound 15, compound 23, compound 26 and compound 27 fabricated minor amyloid fibrils. Furthermore, the sample treated with compound 27 exhibited less aggregation than compound 26 to some extent. After 24 hours of equimolar treatment, curcumin and compound 15 showed the inhibition of approximately 60% while compound 27 and compound 26 showed 85% and >80% $\text{A}\beta$ inhibition of fibrils respectively (Fig. 3).

Concentration-dependent self-mediated $\text{A}\beta_{1-42}$ aggregation inhibition

Following that, we explored the $\text{A}\beta_{1-42}$ aggregation (self-mediated) inhibition, with a focus on concentration dependence. Firstly $\text{A}\beta_{1-42}$ was incubated in both conditions (absence or presence) of different concentrations of the inhibitors (pregnenolone derivatives). The calculated IC_{50} values in μM of the compounds are presented in Table 1. Pregnenolone derivatives 15, 26 and 27 showed good inhibition potential towards self-mediated $\text{A}\beta_{1-42}$ aggregation having IC_{50} values of $9.25\ \mu\text{M}$, $1.04\ \mu\text{M}$ and $5.78\ \mu\text{M}$, respectively (Table 1). Furthermore, curcumin (standard drug) exhibited an IC_{50} value of $7.29\ \mu\text{M}$. The dose-response curve is shown in Fig. 4. This curve displays the inhibitory compound's (15, 23–28 and curcumin) concentration on the x -axis and the percentage inhibition on the y -axis. Results have shown that the amount of inhibition rises along with the concentration of the inhibitory compound.

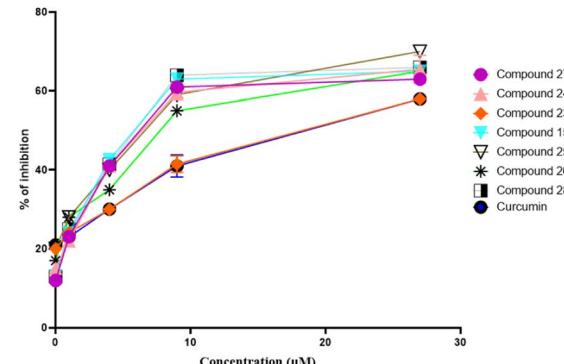


Fig. 4 Dose–response curve of compounds 15, 23–28, and curcumin against self-induced $\text{A}\beta_{1-42}$ aggregation.

Cell viability/neurotoxicity

The cell lines of human neuroblastoma SH-SY5Y were employed to assess the profile cytotoxicity of $\text{A}\beta_{1-42}$ aggregates with the selected synthesized compounds.^{65,66} Experiments for cell viability were conducted to figure out whether these compounds can abandon the neuronal cells from the toxicity of $\text{A}\beta$ -species. MTT assay was accomplished to compare induced $\text{A}\beta_{1-42}$ death of neurons in the incidence and absence of distinct concentrations. Cell viability was assessed at 1, 5, and $10\ \mu\text{M}$ doses along with the standard drug donepezil. Most potent compounds 15, 26 and 27 were selected for this MTT assay. Results are summarized in Fig. 5. The results showed that the viability of cells treated with $\text{A}\beta_{1-42}$ was reduced in contrast to control cells which were without any compound or peptide. Compound 27 at $5\ \mu\text{M}$ was completely non-cytotoxic. Significant toxicity was observed on incubating $5\ \mu\text{M}$ $\text{A}\beta_{1-42}$ with cells of SH-SY5Y, as the death of cells was about 65% enhanced than in the control. On incubation of cells with $\text{A}\beta_{1-42}$ having the selected compounds, an obvious dose–response neuroprotection was perceived. Compound 27, cells of human neuronal SH-SY5Y were effectively protected from the toxicity of the $\text{A}\beta$ peptide. Donepezil

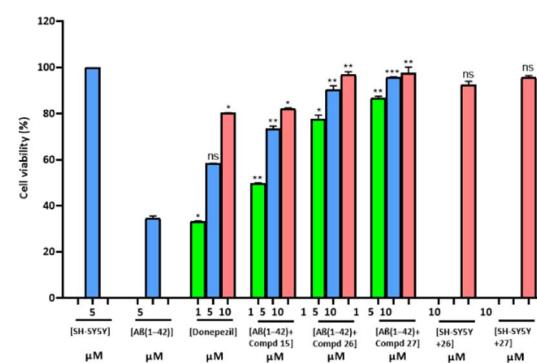


Fig. 5 Viability of the SH-SY5Y cells with $\text{A}\beta_{1-42}$ in the presence and absence of designated concentrations of donepezil and compounds (15, 26, and 27) after 48 h, as determined using the MTS assay. The control group was observed as 100% cell viability, and the assays were performed in triplicate. $*P < 0.05$, $**P < 0.01$, $***P < 0.001$ and ns = non-significant.



(standard drug), compound **15** and compound **26** along with A β ₁₋₄₂ peptide increased cell viability up to approximately 60%, 75% and 90% respectively at the equimolar concentrations.

Hence, A β ₁₋₄₂ peptide in the presence of these compounds, especially **26** and **27** exhibited low toxicity at high concentrations.

The SH-SY5Y cells were also contemplated with compound **26** and compound **27** at 10 μ M concentrations to analyze the cytotoxicity of compounds. Compound **27** showed a more noticeable effect than compound **26** on comparing the average viability. So, the results have shown that both compounds **26** and **27** had a reduced level of A β aggregates and also had reduced cellular cytotoxicity (Fig. 5).

PAMPA BBB assay

In AD therapies the blood–brain barrier (BBB) penetration is a necessary concern. Parallel artificial membrane permeation assay (PAMPA) was employed by utilizing a previously reported procedure.^{59,61,67} Pregnenolone **9** and its 3-amino derivative **12** showed high BBB penetration. Selected all three compounds *i.e.* **23**, **26** and **27** also found to be BBB penetrants in PAMPA assay

Table 2 *In vitro* PAMPA-BBB penetration results

Compounds number	(PAMPA-BBB) ^a P_e (tested) (10^{-6} cm s ⁻¹)	Prediction of CNS penetration ^{b,c}
<i>P_e</i> (tested) evaluation (10^{-6} cm s ⁻¹) for the standard and compounds		
23	16.4	High
26	13.8	High
27	18.1	High
Pregnenolone 9	20.4	High
C-3 amino-pregnenolone (12)	19.6	High
Donepezil	16.7	High
Validation of the model by four commercial drugs		
Diazepam	15.30	High
Atenolol	0.75	No
Alprazolam	5.60	High
Lomefloxacin	1.12	No

^a Data represent are the assay mean for the marketed drugs ($n = 3$).

^b 'CNS+' (prediction of high BBB permeation); P_e (10^{-6} cm s⁻¹) > 4.39 .

^c 'CNS-' (prediction of low BBB permeation); P_e (10^{-6} cm s⁻¹) < 1.78 .

(Table 2). The presence of a hydrophobic steroid nucleus could be the reason for this permeation.

Docking studies

Docking simulations were employed *via* molecular operating environment software (MOE, 2016.0208) on human carbonic anhydrase II complexed with acetazolamide (PDB accession: 3HS4). The reliability of the docking procedure to reproduce the correct binding orientation was determined by the redocking of the extracted ligand from the crystal structure. The protocol with an RMSD value less than 2 \AA was used for docking runs.

Docking studies on hCA-II

Molecular docking runs showed favourable binding modes and hydrophilic/hydrophobic binding affinities with the important residues (amino acid) present in the binding site. In all cases, the sulfonamide nucleus was found to coordinate with the Zn centre. A two-dimensional (2D) binding affinity plot of potent compound **23** is illustrated in Fig. 6. 4-(Sulfamoylphenyl)thiourea moiety was found in contact with the Zn centre *via* coordination bond with one of the nitrogen atoms of sulfonamide at a distance of 2.07 \AA . The computed binding energy for compound **23** is -9.087 kcal mol⁻¹. The binding pose showed that the 4-(sulfamoylphenyl)ethylamino group oriented towards the zinc centre and penetrated deep into the catalytic site. While all the other compounds contacted the zinc centre *via* (4-sulfamoylphenyl)thiourea moiety (Fig. 6a).

Overall, the steroid derivative **23** established three hydrogen bond binding affinity and five hydrophobic interactions (π - π stacking, π -sulfur and π -alkyl). Phe70, Asp71 and Thr200 established hydrophilic (hydrogen bond) binding affinities, while Phe131 forms π - π stacking interactions and amino acid residues His94, His96 and Trp209 are implied in the π -sulfur type of hydrophobic interactions (Fig. 6b).

The binding orientation of native acetazolamide is shown in (Fig. 7a). It showed a strong contact of acetazolamide with zinc ion at a distance of 1.83 \AA . Pregnenolone (**9**), C-3 amine derivative **12** and 4-(sulfamoylphenyl)thiourea analogue **15** experimentally showed poor CA-II inhibition having IC₅₀ values of 89.38 μM , 73.19 μM , and 48.91 μM respectively. The binding orientation in the binding site of hCA-II showed that these

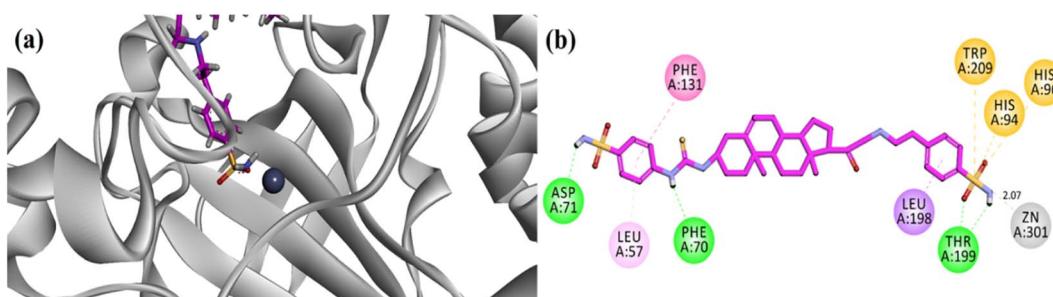


Fig. 6 (a) Three-dimensional ribbon diagram of most potent compound **23** into the binding region of hCA-II. (b) Two-dimensional (2-D) interaction plots of potent compound **23** into the binding site of hCA-II (PDB accession: 3HS4).



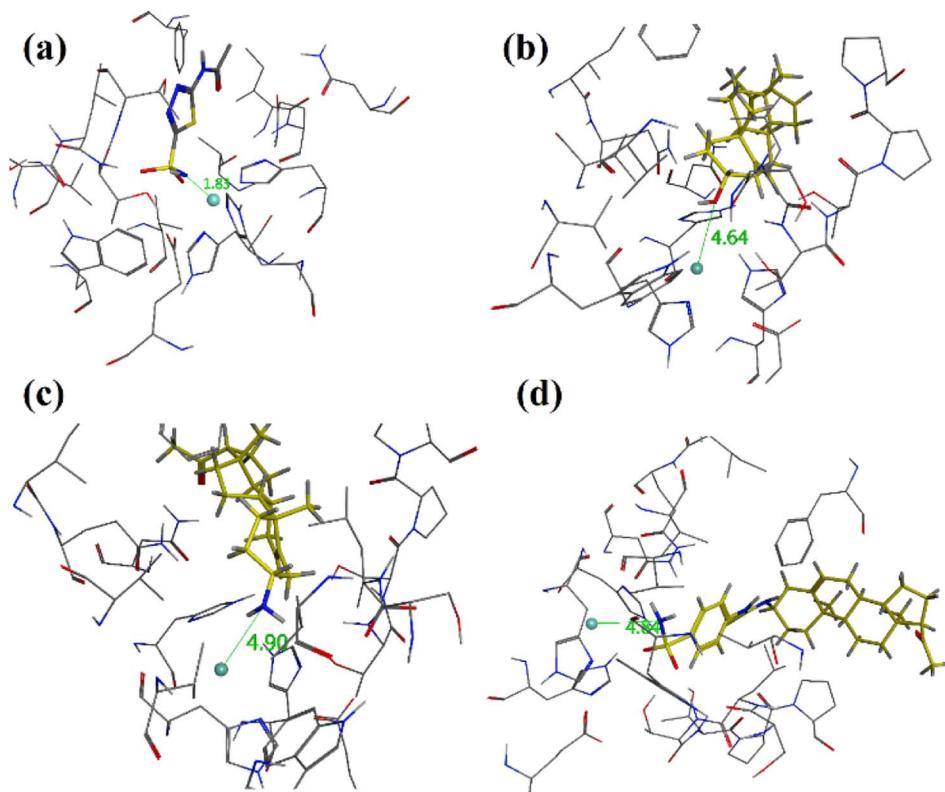


Fig. 7 Three dimensional diagrams of (a) native acetazolamide, (b) pregnenolone (9), (c) 3-amino-pregnenolone derivative **12** and (d) derivative **15** into the binding site of hCA-II. The distance from zinc centre is shown as dotted green lines and distance is in Å (PDB accession: 3HS4).

compounds bind far from the Zn centre at a distance of 4.64 Å, 4.90 Å and 4.83 Å respectively (Fig. 7a-d).

Docking studies on AChE

Docking studies on AChE was carried out on *Torpedo californica* acetylcholinesterase (*TcAChE*) complexed with

alkylene-linked bis-tacrine (PDB ID = 2CKM). After validation of the docking protocol, the docking simulation studies were employed for synthesized pregnenolone derivatives into the binding site of 2CKM. Docking studies support dual binding site (PAS and CAS) inhibition of AChE which ultimately results in $\text{A}\beta_{1-42}$ aggregation and AChE inhibition. All the active compounds showed interactions with amino acid residues

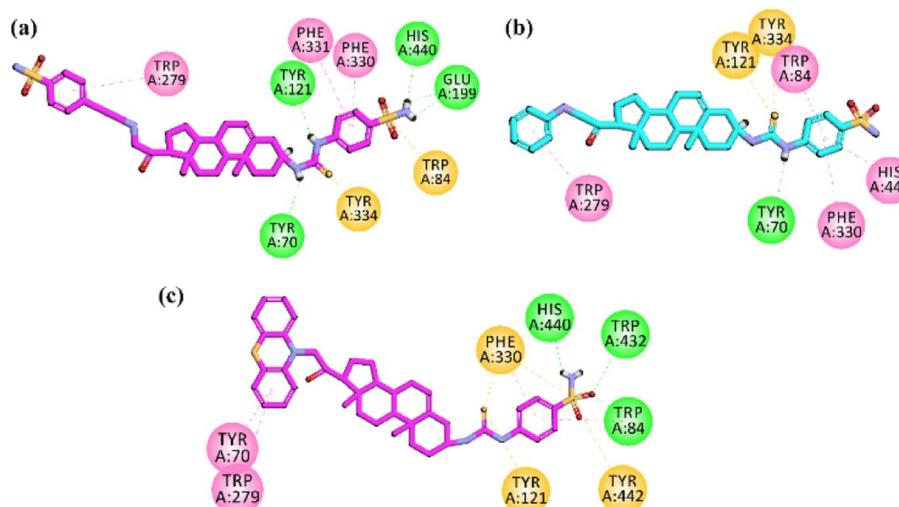


Fig. 8 Two-dimensional (2-D) interaction plots of potent compound (a) **23** (b) **26** and (c) **27** into the binding region of AChE (PDB accession: 2CKM).

present in PAS (Tyr334, Trp279, Tyr121 and Tyr70) and CAS (Phe331, Phe330 and Trp84). Two-dimensional (2-D) binding affinity plots of active compounds **23**, **26** and **27** into the binding site of TcAChE are shown in Fig. 8a–c. It is important to note from 2D interaction plots that all three compounds under study interact with His440 (catalytic triad residue) *via* hydrogen (compounds **23** and **27**) or π – π stacking interactions (compound **26**). The high inhibition potential of these compounds towards AChE may be due to these interactions.

Docking studies on amyloid- β_{42} peptide

We applied molecular modelling studies to explore how the synthesized compounds interact with amyloid- β . The analysis was performed on two different forms of amyloid- β . We consider helical monomer (PDB ID = 1IYT) and β -sheet pentamer or $\text{A}\beta_{42}$ protofibril oligomer (PDB ID = 2BEG). Both the structures have been determined by NMR spectroscopy. The β -sheet pentamer or $\text{A}\beta_{42}$ protofibril oligomer (PDB ID = 2BEG) has been used as a suitable model in several recent computational studies. The pentameric model, 2BEG, has been used as a suitable model in several recent computational studies.

Docking studies on helical monomer

To prevent the $\text{A}\beta$ (self-aggregation), the small molecule is needed to impede the formation of structures of β -sheet.^{68,69} The stability of these β -sheet structures relies on the creation of salt bridges involving Asp23 and Glu22, as well as interactions with hydrophobic residues such as Phe20, Glu22, Asp23, Gly29, and Ala30. We conducted molecular docking experiments with newly synthesized compounds, targeting the binding region of $\text{A}\beta_{1-42}$, utilizing the protein data bank code 1IYT. There were significant hydrophobic interactions observed with the amino acids Asp23, Glu22, Phe20, and Phe19, effectively impeding the formation of β -sheet structures. Additionally, when analysing the docking results for compound **15**, hydrogen bond interactions with residues Asp23, and Glu22 were found.

Moreover, Asn27, Lys28 and Gly25 also established hydrogen bond interactions. A π -sulphur interaction with important residue Asn27 was also observed. Docking results of compound **26** exhibited binding affinities with the binding region of $\text{A}\beta$ through hydrophilic (hydrogen bond) formation through the oxygens of $-\text{SO}_2\text{OH}$ with Leu34 and Lys28, A π – π interaction from benzene ring with Phe19 was observed. Moreover, $-\text{NH}$ also formed hydrogen bond interactions with Asp23, Glu22, and π -anion interaction with residue Phe20. This array of binding affinities might contribute to α -helical stabilization and hindrance of the formation of β -sheet. The binding affinities in docking results of the compound **27** explained its extraordinary affinity as it shows the important interactions with the amino acid residues Lys28, Asp23, Glu22, Phe20 and Phe19, and through π -sulphur, π -anion and conventional hydrogen bonding. NH_2 group showed binding through hydrogen bonding through the residues Asp23, and Glu22, while sulphur showed π -sulphur binding affinity with residue Phe20 (Fig. 9). The binding modes of compounds **26** and **27** are shown in Fig. 10.

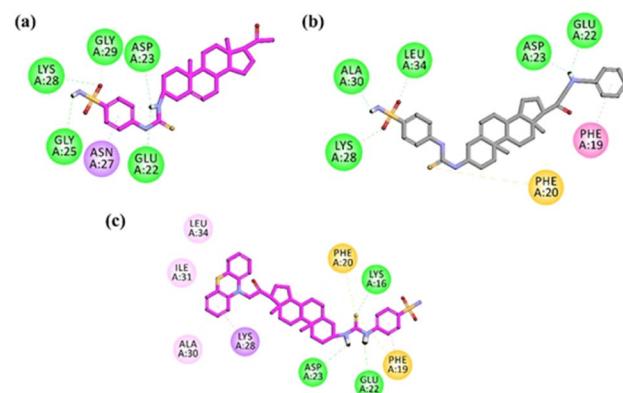


Fig. 9 Two-dimensional (2-D) interaction plots of active compound (a) **15**, (b) **26** and (c) **27** into the 3D crystallographic structure of amyloid- β_{42} peptide (PDB ID = 1IYT).

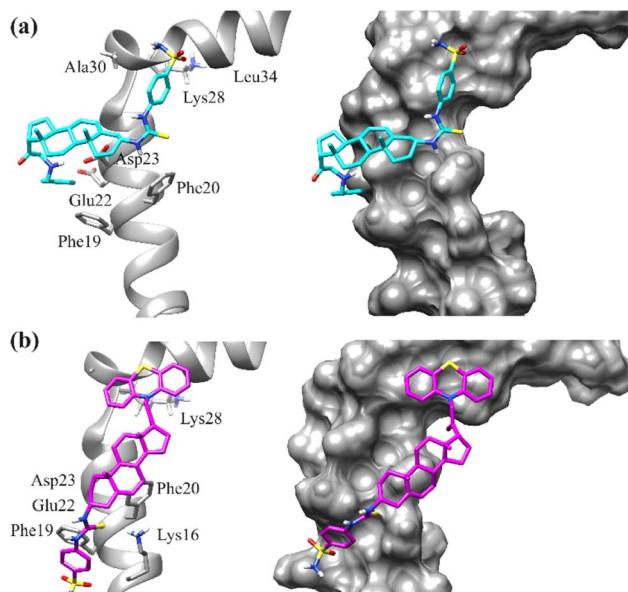


Fig. 10 Binding mode of compound (a) **26** (b) **27** into the binding site of amyloid- β_{42} peptide (PDB ID = 1IYT). Left: ribbon diagrams; right: surface diagrams.

Docking studies on β -sheet pentamer (oligomer)

Next, we docked selected compounds **15**, **23**, **26** and **27** with β -sheet pentamer or $\text{A}\beta_{42}$ protofibril (PDB ID = 2BEG) to determine the preferred binding region. Two possible binding modes of selected compounds were observed (Fig. 11). Compounds **15** and **23** were docked at the C-terminal of the pentamer.⁷⁰ The 2-D interaction plots are shown in Fig. S1 (ESI†). While compounds **26** and **27** were docked outside the amyloid fibrils.

The 2-D interaction plot of **26** showed three hydrogen bond interactions with Glu22, Asp23 and Gly25. While weak hydrophobic interactions were also observed with Val24 and Val36 (Fig. 12).^{26,70}



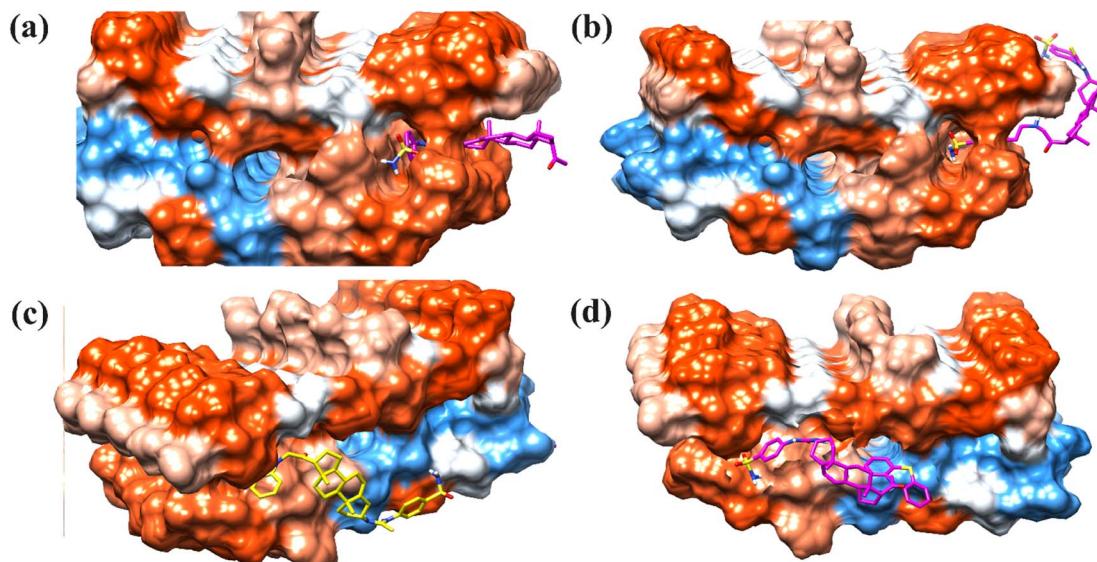


Fig. 11 T3-D docked pose of compounds (a) 15, (b), 23, (c) 26 and (d) 27 into the A_β protofibrils (PDB ID: 2BEG).

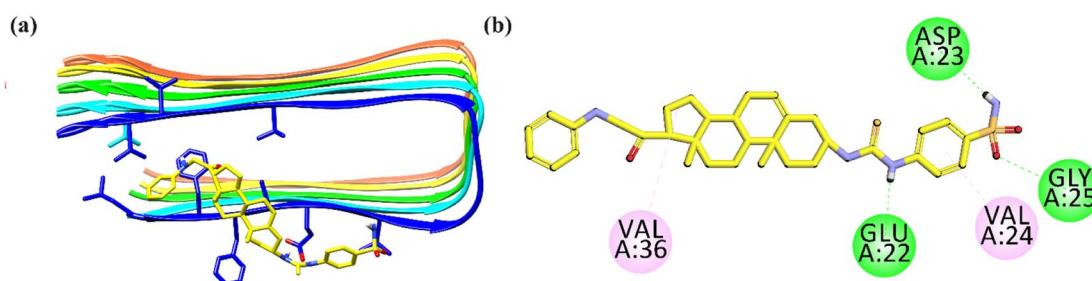


Fig. 12 Binding mode of most active A_β-aggregation inhibitor 26 with A_β protofibrils (a) ribbon diagram; (b) 2-D interaction plot.

Conclusions

Current work is focused on identifying the compounds that have balanced even mild biological activities against multiple targets instead of finding one-target compound with high potency. Structural modification of the steroid nucleus has got considerable attention in recent years. This is because steroid-based therapeutics have several advantages and possess interesting structural features for the treatment of various diseases. Alzheimer's disease, a neurodegenerative condition, is characterized by the accumulation of a peptide called amyloid- β_{1-42} peptide. Drugs that are expected to reduce A β_{1-42} production, prevent A β_{1-42} aggregation, and promote A β_{1-42} clearance are promising approaches for treating AD. Moreover, small molecules having the ability to bind with the peripheral anionic site (PAS) will inhibit A β_{1-42} aggregation and AChE. Additionally, studies have revealed that some carbonic anhydrase inhibitors can diminish the neuroinflammation and deposition of A β_{1-42} aggregates at the walls of vessels of the brain *in vivo*. Keeping these points in mind, our current study involved the structural modification of pregnenolone. Compound 23 with two sulfonamide moieties present on two different phenyl rings emerged as potent hCA-II inhibitors with

an IC₅₀ value of $0.67 \pm 0.09 \mu\text{M}$. Self-mediated A β_{1-42} aggregation inhibition assay showed that compounds 26 and 27 have shown >80% and 85% inhibition of A β fibrils respectively. In concentration-dependent self-mediated A β_{1-42} aggregation inhibition assay compounds 15, 26 and 27 showed good inhibition potential with IC₅₀ values of $9.25 \mu\text{M}$, $1.04 \mu\text{M}$ and $5.78 \mu\text{M}$ respectively. In cholinesterase inhibition assay 4-(sulfonylphenyl)thiourea derivative 23 emerged as a nanomolar inhibitor of AChE and BChE with IC₅₀ = $0.04 \mu\text{M}$ and $0.58 \mu\text{M}$ respectively with SI = 14.5.

Analysis of docked complexes showed that all the compounds confined in the active site coordinated with zinc centre and formed hydrogen bond interactions with key amino acid residues. Docking studies support dual binding site (PAS and CAS) inhibition of AChE which showed A β_{1-42} aggregation and AChE inhibition. Moreover, docking studies carried out on 3D crystallographic structure of A β_{1-42} peptide (PDB ID = 1IYT) showed significant hydrophobic interactions with the amino acids Phe19, Phe20, Glu22, and Asp23, effectively impeding the formation of β -sheet structures.

In conclusion, our current study is the first-ever report on the structural modification of pregnenolone at C-3 (-OH) and C-17 (acetyl group). Our synthesized derivatives 23, 25-27 exhibited



concomitant inhibition of all the tested macromolecular targets and hence emerged as multitarget hybrid compounds among all synthesized pregnenolone derivatives.

Author contributions

UR conceived, designed, and supervised this study. He was involved in all the phases (from designing to synthesis/pharmacological evaluation and manuscript writing/editing) that led to the completion of the manuscript. AT and BM synthesized the compounds. *In vitro* experiments were performed by FH and AS. Docking studies were performed and written by AT, BM and UR. The manuscript was drafted by AT, BM, and FH. UR reviewed and edited the drafts.

Conflicts of interest

The authors declare no conflict of interest.

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References

- U. Rashid and F. L. Ansari, in *Drug Design and Discovery in Alzheimer's Disease*, Elsevier, 2014, pp. 40–141.
- E. Cerf, R. Sarroukh, S. Tamamizu-Kato, L. Breydo, S. Derclaye, Y. F. Dufrêne, V. Narayanaswami, E. Goormaghtigh, J.-M. Ruysschaert and V. Raussens, *Biochem. J.*, 2009, **421**, 415–423.
- B. S. Gadad, G. B. Britton and K. Rao, *J. Alzheimer's Dis.*, 2011, **24**, 223–232.
- D. A. Butterfield, A. M. Swomley and R. Sultana, *Antioxid. Redox Signaling*, 2013, **19**, 823–835.
- H. Hampel, F. Caraci, A. C. Cuello, G. Caruso, R. Nisticò, M. Corbo, F. Baldacci, N. Toschi, F. Garaci and P. A. Chiesa, *Front. Immunol.*, 2020, **11**, 456.
- V. Tiwari and S. Shukla, *Front. Genet.*, 2023, **14**, 1057068.
- G. Dixit and A. Prabhu, *J. Mol. Struct.*, 2023, **1294**, 136498.
- H. Hampel, J. Hardy, K. Blennow, C. Chen, G. Perry, S. H. Kim, V. L. Villemagne, P. Aisen, M. Vendruscolo and T. Iwatsubo, *Mol. Psychiatry*, 2021, **26**, 5481–5503.
- S. S. Hirschbeck, E. T. Lindberg, J. H. Jang, M. R. Jacob, K. L. Lazar Cantrell and T. D. Do, *ACS Chem. Neurosci.*, 2024, **15**(7), 1523–1532.
- F. Hussain, A. Tahir, M. S. Jan, N. Fatima, A. Sadiq and U. Rashid, *RSC Adv.*, 2024, **14**, 10304–10321.
- V. Kumar and K. Roy, *Computational Modeling of Drugs against Alzheimer's Disease*, 2023, pp. 3–47.
- J. Bieschke, *Neurotherapeutics*, 2013, **10**, 429–439.
- Z. Du, M. Li, J. Ren and X. Qu, *Acc. Chem. Res.*, 2021, **54**, 2172–2184.
- X. Han and G. He, *ACS Chem. Neurosci.*, 2018, **9**, 198–210.
- S. Sharari, N. N. Vaikath, M. Tsakou, S. S. Ghanem and K. Vekrellis, *Int. J. Mol. Sci.*, 2023, **24**, 11326.
- G. E. Vitek, B. Decourt and M. N. Sabbagh, *Expert Opin. Invest. Drugs*, 2023, **32**, 89–94.
- A. V. Atoki, P. M. Aja, E. N. Ondari and T. S. Shinkafi, *Int. J. Food Prop.*, 2023, **26**, 2091–2127.
- N. Stern, A. Gacs, E. Tátrai, B. Flachner, I. Hajdú, K. Dobi, I. Bágyi, G. Dormán, Z. Lőrincz and S. Cseh, *Int. J. Mol. Sci.*, 2022, **23**, 13098.
- A. Alvarez, C. Opazo, R. Alarcón, J. Garrido and N. C. Inestrosa, *J. Mol. Biol.*, 1997, **272**, 348–361.
- N. C. Inestrosa, A. Alvarez, C. A. Perez, R. D. Moreno, M. Vicente, C. Linker, O. I. Casanueva, C. Soto and J. Garrido, *Neuron*, 1996, **16**, 881–891.
- A. Alvarez, R. Alarcon, C. Opazo, E. O. Campos, F. J. Munoz, F. H. Calderon, F. Dajas, M. K. Gentry, B. P. Doctor and F. G. De Mello, *J. Neurosci.*, 1998, **18**, 3213–3223.
- F. J. Carvajal and N. C. Inestrosa, *Front. Mol. Neurosci.*, 2011, **4**, 19.
- J. L. Sussman, M. Harel, F. Frolov, C. Oefner, A. Goldman, L. Toker and I. Silman, *Science*, 1991, **253**, 872–879.
- M. Bartolini, C. Bertucci, V. Cavrini and V. Andrisano, *Biochem. Pharmacol.*, 2003, **65**, 407–416.
- N. C. Inestrosa, M. C. Dinamarca and A. Alvarez, *FEBS J.*, 2008, **275**, 625–632.
- O. M. Waly, K. M. Saad, H. I. El-Subbagh, S. M. Bayomi and M. A. Ghaly, *Eur. J. Med. Chem.*, 2022, **231**, 114152.
- P. Agostinho, A. Pliassova, C. R. Oliveira and R. A. Cunha, *J. Alzheimer's Dis.*, 2015, **45**, 329–347.
- H. M. Al-Kuraishi, M. S. Jabir, A. I. Al-Gareeb, A. K. Albuhadily, S. Albukhaty, G. M. Sulaiman and G. E.-S. Batiha, *Ageing Res. Rev.*, 2023, 102119.
- M. Vallée, *J. Steroid Biochem. Mol. Biol.*, 2016, **160**, 78–87.
- Y. Xia, Z.-H. Wang, Z. Zhang, X. Liu, S. P. Yu, J.-Z. Wang, X.-C. Wang and K. Ye, *Prog. Neurobiol.*, 2021, **204**, 102113.
- A. D. Corbett, *Pseudo-Dynamic Combinatorial Libraries*, 2003.
- T. K. Hurst, D. Wang, R. B. Thompson and C. A. Fierke, *Biochim. Biophys. Acta*, 2010, **1804**, 393–403.
- C. T. Supuran, *Expert Opin. Ther. Targets*, 2023, **27**, 897–910.
- C. T. Supuran and A. Scozzafava, *Expert Opin. Ther. Pat.*, 2000, **10**, 575–600.
- A. Nocentini, A. Angelini, F. Carta, J.-Y. Winum, R. Zalubovskis, S. Carradori, C. Capasso, W. A. Donald and C. T. Supuran, *J. Enzyme Inhib. Med. Chem.*, 2021, **36**, 561–580.
- S. Kumar, S. Rulhania, S. Jaswal and V. Monga, *Eur. J. Med. Chem.*, 2021, **209**, 112923.
- C. T. Supuran, *Curr. Pharm. Des.*, 2010, **16**, 3233–3245.
- S. Ghorai, S. Pulya, K. Ghosh, P. Panda, B. Ghosh and S. Gayen, *Bioorg. Chem.*, 2020, **95**, 103557.
- A. Angelini, A.-M. Alaa, A. Nocentini, A. S. El-Azab, P. Gratteri and C. T. Supuran, *Bioorg. Med. Chem.*, 2017, **25**, 5373–5379.
- M. A. Mohamed, A.-M. Alaa, H. M. Sakr, A. S. El-Azab, S. Bua and C. T. Supuran, *Bioorg. Med. Chem.*, 2017, **25**, 2524–2529.
- C. T. Supuran, *Clin. Sci.*, 2021, **135**, 1233–1249.
- D. Tarkowská, *Molecules*, 2019, **24**, 2585.



43 A. Gupta, B. S. Kumar and A. S. Negi, *J. Steroid Biochem. Mol. Biol.*, 2013, **137**, 242–270.

44 N. H. Mustafa, M. Sekar, S. Fuloria, M. Y. Begum, S. H. Gan, N. N. I. M. Rani, S. Ravi, K. Chidambaram, V. Subramaniyan and K. V. Sathasivam, *Molecules*, 2022, **27**, 2032.

45 J. Y. Winum, A. Scozzafava, J. L. Montero and C. T. Supuran, *Med. Res. Rev.*, 2005, **25**, 186–228.

46 A. Nocentini, A. Bonardi, P. Gratteri, B. Cerra, A. Gioiello and C. T. Supuran, *J. Enzyme Inhib. Med. Chem.*, 2018, **33**, 1453–1459.

47 A. Khalid, M. N. Ghayur, F. Feroz, A. Gilani and M. I. Choudhary, *J. Steroid Biochem. Mol. Biol.*, 2004, **92**, 477–484.

48 A. Cecchi, A. Hulikova, J. Pastorek, S. Pastoreková, A. Scozzafava, J.-Y. Winum, J.-L. Montero and C. T. Supuran, *J. Med. Chem.*, 2005, **48**, 4834–4841.

49 A. Peschiulli, D. Oehlrich, M. Van Gool, N. Austin, S. Van Brandt, M. Surkyn, M. De Cleyn, A. Vos, G. Tresadern and F. J. Rombouts, *ACS Med. Chem. Lett.*, 2021, **13**, 76–83.

50 Z. Y. Wang, M. R. Pergande, C. W. Ragsdale and S. M. Cologna, *Curr. Biol.*, 2022, **32**, 2572–2579.

51 D. S. Reddy, *Prog. Brain Res.*, 2010, **186**, 113–137.

52 V. Kolas, J. S. A. Bandonil, N. Wali, K.-C. Hsia, J.-J. Shie and B.-c. Chung, *Cell Biosci.*, 2022, **12**, 190.

53 M. B. Tufail, M. A. Javed, M. Ikram, M. H. Mahnashi, B. A. Alyami, Y. S. Alqahtani, A. Sadiq and U. Rashid, *Steroids*, 2021, **168**, 108801.

54 B. A. Alyami, I. Ejaz, M. H. Mahnashi, Y. S. Alqahtani, A. O. Alqarni, M. S. Jan, A. Sadiq and U. Rashid, *Steroids*, 2022, **185**, 109059.

55 Y. Pocker and J. Meany, *Biochemistry*, 1967, **6**, 239–246.

56 N. Ur Rehman, S. A. Halim, M. Khan, H. Hussain, H. Yar Khan, A. Khan, G. Abbas, K. Rafiq and A. Al-Harrasi, *Pharmaceuticals*, 2020, **13**, 94.

57 A. Khan, M. Khan, S. A. Halim, Z. A. Khan, Z. Shafiq and A. Al-Harrasi, *Front. Chem.*, 2020, **8**, 598095.

58 M. A. Javed, S. Bibi, M. S. Jan, M. Ikram, A. Zaidi, U. Farooq, A. Sadiq and U. Rashid, *RSC Adv.*, 2022, **12**, 22503–22517.

59 S. K. Bowroju, N. Mainali, S. Ayyadevara, N. R. Pentala, S. Krishnamachari, S. Kakraba, R. J. S. Reis and P. A. Crooks, *Molecules*, 2020, **25**, 3610.

60 C. B. Mishra, S. Shalini, S. Gusain, A. Prakash, J. Kumari, S. Kumari, A. K. Yadav, A. M. Lynn and M. Tiwari, *RSC Adv.*, 2020, **10**, 17602–17619.

61 B. Brus, U. Kosak, S. Turk, A. Pislar, N. Coquelle, J. Kos, J. Stojan, J.-P. Colletier and S. Gobec, *J. Med. Chem.*, 2014, **57**, 8167–8179.

62 M. Matloobi and C. O. Kappe, *J. Comb. Chem.*, 2007, **9**, 275–284.

63 F. Iftikhar, F. Yaqoob, N. Tabassum, M. S. Jan, A. Sadiq, S. Tahir, T. Batool, B. Niaz, F. L. Ansari and M. I. Choudhary, *Bioorg. Chem.*, 2018, **80**, 99–111.

64 F. Sonmez, B. Zengin Kurt, I. Gazioglu, L. Basile, A. Dag, V. Cappello, T. Ginex, M. Kucukislamoglu and S. Guccione, *J. Enzyme Inhib. Med. Chem.*, 2017, **32**, 285–297.

65 R. Saini, G. R. Navale, S. Singh, H. K. Singh, R. Chauhan, S. Agrawal, D. Sarkar, M. Sarma and K. Ghosh, *Int. J. Biol. Macromol.*, 2023, **248**, 125847.

66 M. A. Javed, N. Ashraf, M. Saeed Jan, M. H. Mahnashi, Y. S. Alqahtani, B. A. Alyami, A. O. Alqarni, Y. I. Asiri, M. Ikram and A. Sadiq, *ACS Chem. Neurosci.*, 2021, **12**, 4123–4143.

67 S. Manzoor, M. T. Gabr, B. Rasool, K. Pal and N. Hoda, *Bioorg. Chem.*, 2021, **116**, 105354.

68 T. Umar, S. Shalini, M. K. Raza, S. Gusain, J. Kumar, P. Seth, M. Tiwari and N. Hoda, *Eur. J. Med. Chem.*, 2019, **175**, 2–19.

69 A. Pasieka, D. Panek, N. Szałaj, A. Espargaró, A. Więckowska, B. Malawska, R. Sabaté and M. Bajda, *ACS Chem. Neurosci.*, 2021, **12**, 2057–2068.

70 A. A. Mohammed, S. S. Barale, S. A. Kamble, S. B. Paymal and K. D. Sonawane, *Int. J. Biol. Macromol.*, 2023, **242**, 124880.

