



Novel serotonin 5-HT_{2A} receptor antagonists derived from 4-phenylcyclohexane-5-spiro-and 5-methyl-5-phenyl-hydantoin, for use as potential antiplatelet agents

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Abstract

Background Antiplatelet drugs have been used in the treatment of acute coronary syndromes and for the prevention of recurrent events. Unfortunately, many patients remain resistant to the available antiplatelet treatment. Therefore, there is a clinical need to synthesize novel antiplatelet agents, which would be associated with different pathways of platelet aggregation, to develop an alternative or additional treatment for resistant patients. Recent studies have revealed that 5-HT_{2A} receptor antagonists could constitute alternative antiplatelet therapy.

Methods Based on the structures of the conventional 5-HT_{2A} receptor ligands, two series of compounds with 4-phenylcyclohexane-5-spiro- or 5-methyl-5-phenyl-hydantoin core linked to various arylpiperazine moieties were synthesized and their affinity for 5-HT_{2A} receptor was assessed. Further, we evaluated their antagonistic potency at 5-HT_{2A} receptors using isolated rat aorta and cells expressing human 5-HT_{2A} receptors. Finally, we studied their anti-aggregation effect and compared it with ketanserin and sarpogrelate, the reference 5-HT_{2A} receptor antagonists. Moreover, the structure–activity relationships were studied following molecular docking to the 5-HT_{2A} receptor model.

Results Functional bioassays revealed some of the synthesized compounds to be moderate antagonists of 5-HT_{2A} receptors. Among them, **13**, 8-phenyl-3-(3-(4-phenylpiperazin-1-yl)propyl)-1,3-diazaspiro[4.5]decane-2,4-dione, inhibited collagen stimulated aggregation (IC₅₀ = 27.3 μM) being more active than sarpogrelate (IC₅₀ = 66.8 μM) and comparable with ketanserin (IC₅₀ = 32.1 μM). Moreover, compounds 2–5, 9–11, 13, 14 inhibited 5-HT amplified, ADP- or collagen-induced aggregation.

Conclusions Our study confirmed that the 5-HT_{2A} antagonists effectively suppress platelet aggregation and remain an interesting option for the development of novel antiplatelet agents with an alternative mechanism of action.

Keywords Imidazolidinone-2,4-dione · 5-HT_{2A} receptors · Hydantoin · Antiplatelet · Aggregation

Introduction

Antiplatelet drugs have long been used in the treatment of acute coronary syndromes and for the prevention of recurrent events. Large clinical trials have confirmed that antiplatelet agents such as clopidogrel and aspirin are capable of reducing the risk of myocardial infarction, stroke, or death [1, 2]. However, many patients treated with these agents still experience recurrent atherothrombotic events [3]. Moreover, some are resistant to aspirin [cyclooxygenase-1 (COX-1) inhibitor] or clopidogrel (P2Y₁₂ receptor antagonist), even when they are used in combination, which increases their risk of further cardiovascular events [4–6]. Therefore, there is still a clinical need to synthesize novel antiplatelet

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agents involving different pathways of platelet aggregation, as an alternative or additional treatment for resistant patients.

Serotonin (5-HT) is a platelet activator, stored in platelet-dense granules [5]. Among the seven classes of 5-HT receptors, only 5-HT_{2A} has been found in platelets. Upon platelet activation, 5-HT is released to plasma and, in an autocrine manner, promotes further platelet activation, acting at 5-HT_{2A} receptors. Released serotonin is also a potent vasoconstrictor of coronary arteries with a damaged endothelium [5, 7, 8]. Furthermore, coronary artery disease exerts a stimulating effect on platelets, triggering them to release serotonin [9]. Several studies on animal models and humans have shown that 5-HT_{2A} receptor antagonists can inhibit platelet aggregation [5, 8]. 5-HT may also be partially responsible for the increased residual platelet reactivity observed in patients who are on clopidogrel treatment after coronary stent placement, while 5-HT_{2A} antagonists reduce high on-treatment platelet reactivity. Therefore, in addition to the established therapies, serotonin antagonism at 5-HT_{2A} receptors might be a promising approach to improve the treatment outcomes of patients with coronary artery disease [5, 10].

Our previous studies [11–13] showed that a differently substituted hydantoin ring connected to arylpiperazine moiety being the amine part, may interact with 5-HT_{2A} receptors

and exert an antagonistic effect. Among these structures, two hydantoin derivatives with the highest affinity ($K_i = 50$ and 69 nM, respectively) and selectivity for 5-HT_{2A} receptors were selected, namely, 4-phenylcyclohexane-5-spirohydantoin (compound **A**) and 5-methyl-5-phenylhydantoin (compound **B**) linked with 4-phenylpiperazin-1-yl-propyl moiety, which displayed significant antagonistic activity for 5-HT_{2A} receptors (Fig. 1). The compounds **A** and **B** fit the pharmacophore models of 5-HT_{2A} receptor antagonists proposed over the past two decades, which typically share two aryl rings or hydrophobic fragments and basic amine moiety separated from each other by a distance of $5.2\text{--}8.4$ Å and $5.7\text{--}8.5$ Å [14] (Supplementary material Figure S1). These results were a prerequisite for designing a new series of 5-methyl-5-phenylhydantoin and 4-phenylcyclohexane-5-spirohydantoin derivatives, which can serve as 5-HT_{2A} serotonin receptor ligands. The compounds have been designed as analogs of the aforementioned hydantoin derivatives (compound **A** and **B**) by combining structural fragments present in the conventional 5-HT_{2A} receptor antagonists, namely ketanserin (5-HT_{2A}: $K_i = 2$ nM), ritanserin (5-HT_{2A}: $K_i = 0.45$ nM), and fananserin (5-HT_{2A}: $K_i = 0.37$ nM). The structural modifications carried out to develop novel 5-HT_{2A} receptor antagonists were (1) the extension of the alkyl spacer linking hydantoin with arylpiperazine moiety and (2) the

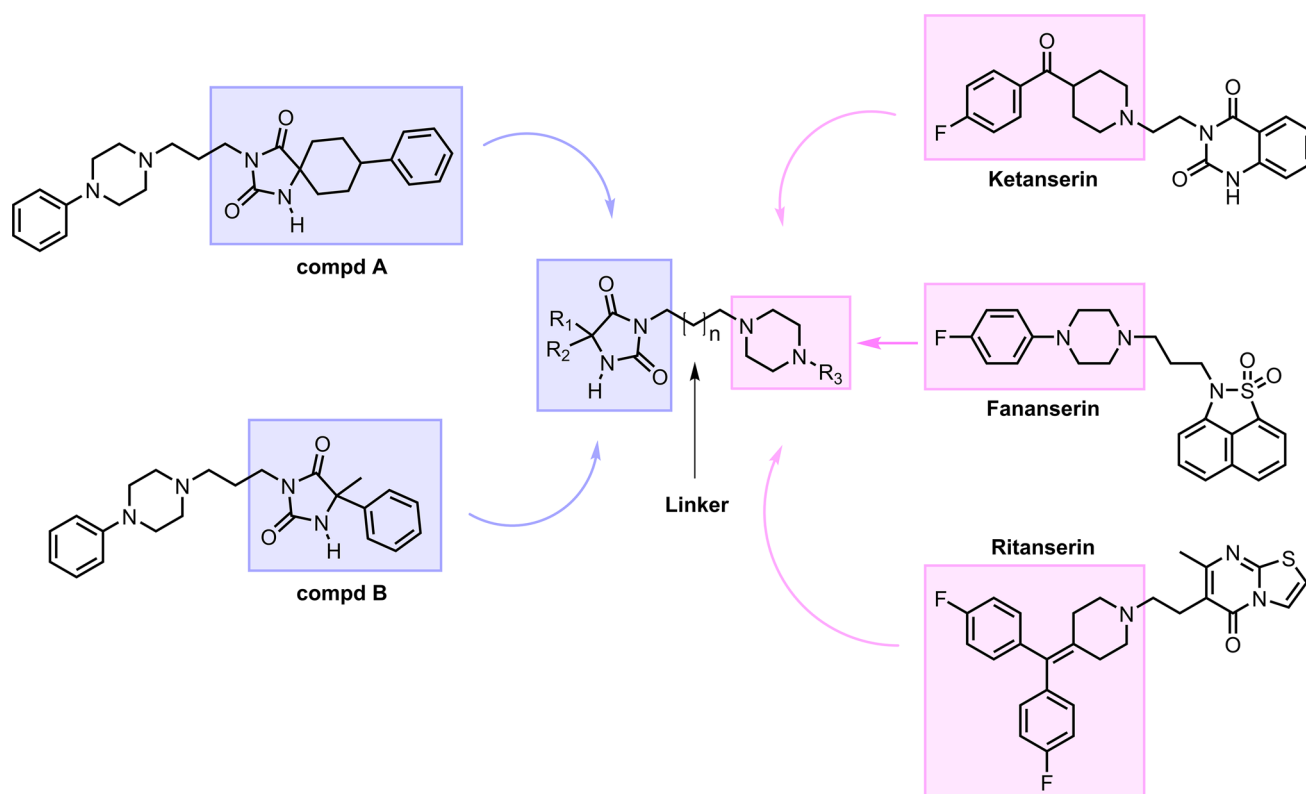


Fig. 1 The general structures of designed compounds based on structure compounds **9** and **12** as well as conventional 5-HT_{2A} antagonists: ketanserin, fananserin and ritanserin

introduction of a differently substituted arylpiperazine fragment in the amine part.

This manuscript describes the synthesis of the designed compounds and their affinity for the 5-HT_{2A} receptors. The intrinsic activity of the most promising compounds was evaluated, and molecular modeling studies were undertaken to identify the structural fragments responsible for their in vitro affinity. Finally, the anti-aggregation effect was tested after inducing the aggregation of whole rat blood with collagen and with 5-HT and ADP or collagen at the sub-threshold concentration.

Materials and methods

Chemistry

All chemicals and solvents were purchased from commercial suppliers (Aldrich and Chempur) and were used without further purification. Melting points were measured in open capillaries on an Electrothermal 9300 apparatus. Thin-layer chromatography (TLC) was run on Merck silica gel 60 F₂₅₄ aluminium sheets (Merck; Germany), using the following solvents (v:v): (S₁) dichloromethane (9)/methanol (0.3), (S₂) ethyl acetate (9)/methanol (1), (S₃) chloroform (7)/n-hexane (2)/acetone (2). Analytical HPLC was conducted on a Waters HPLC instrument with Waters 485 Tunable Absorbance Detector UV, equipped with a Symmetry column (C18, 3.5 µm, 4.6×30 mm) using water/acetonitrile gradient with 0.1% TFA as mobile phase at a flow rate of 5 ml/min. Additionally, the liquid chromatography/mass spectrometry (LC/MS) analysis was performed on the Waters Acquity TQD system, with a Waters TQD quadrupole mass spectrometer with detection by UV (DAD) using an Acquity UPLC BEH C18 column (1.7 µm, 2.1×100 mm). Water/acetonitrile gradient with 0.1% TFA was used as a mobile phase at a flow rate of 0.3 ml/min. The UPLC/MS purity of the investigated compounds (**1–12**) ranged to be over 95%. Elemental analyses (C, H, and N) for final compounds were carried out by a micro method using the elemental Vario EI III Elemental analyzer (Hanau, Germany). NMR spectra were recorded on Varian Mercury 300 MHz spectrometer (Varian Inc., Palo Alto, CA, USA) using the solvent (CDCl₃) signal as an internal standard; chemical shifts are expressed in parts per million (ppm). Signal multiplets are represented by the following abbreviations: s (singlet), brs (broad singlet), d (doublet), t (triplet), m (multiplet). The Discover model, CEM microwave reactor was used to carry out the pressurized reaction.

5-Methyl-5-phenylimidazolidine-2,4-dione (A, scheme 1), 8-phenyl-1,3-diazaspiro[4.5]decane-2,4-dione (C, scheme 1) and intermediate products (N3 halogealkyl derivatives of substituted imidazolidyno-2,4-diones) (B

and D, scheme 1) [11, 12, 15] as well as (4-fluorophenyl) (piperazin-1-yl)methanone [16] were described earlier and the analytical results are consistent with those previously published.

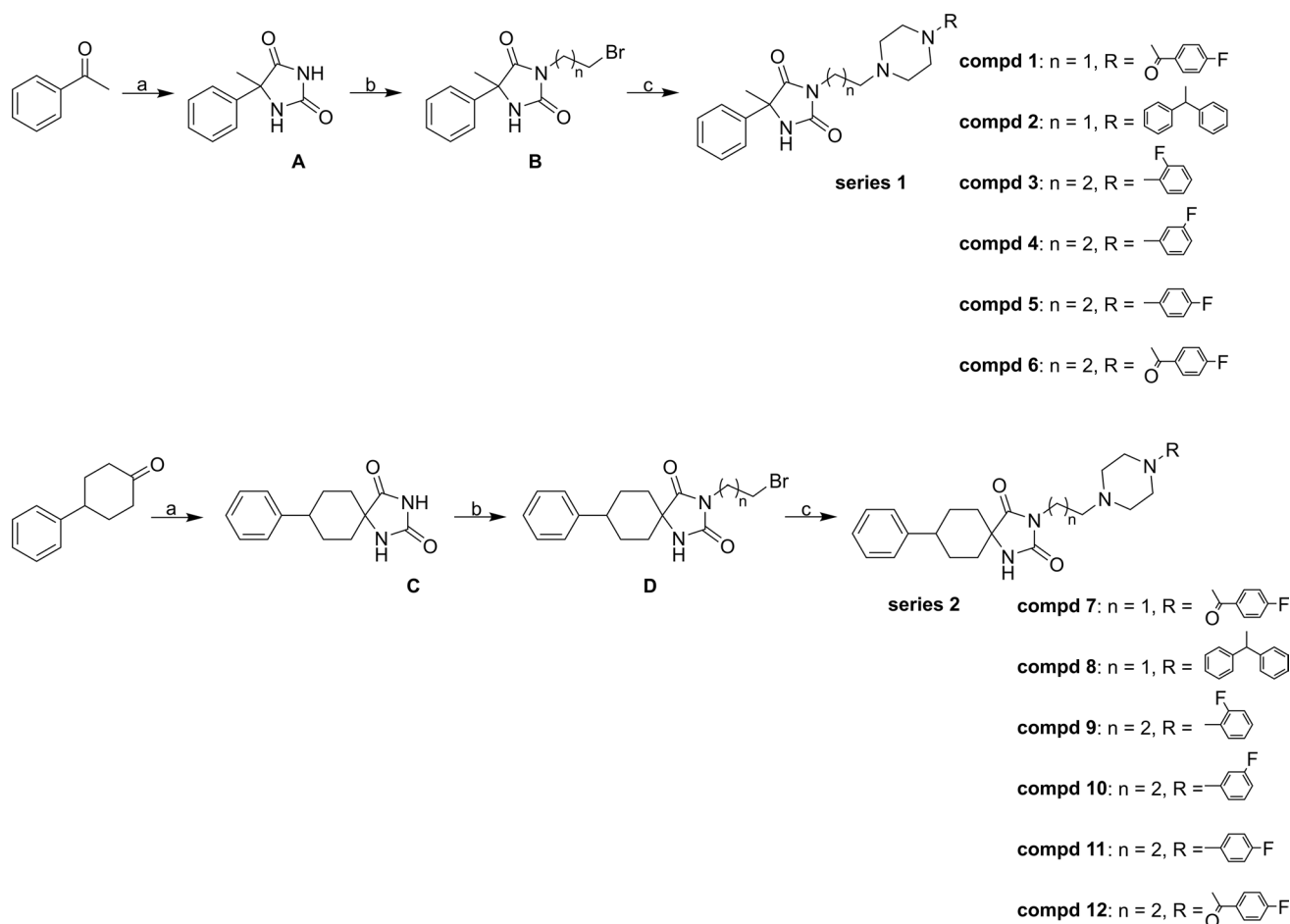
General procedure for the synthesis of final compounds (1–12)

The final compounds (**1–12**) were obtained within two different synthetic pathways, described in procedure A and B. Procedure A (**1–6**, **8**, **9**, **11**): the final compounds were synthesized in a CEM microwave reactor (Discover model). The intermediate compound (0.3 mmol), trimethylamine (0.6 mmol) and appropriate piperazine derivatives (0.36 mmol) in acetonitrile (4 mL) was added to the microwave flask, and stirred for 65–100 min, at 100 °C, 200 W. Upon cooling to room temperature, the reaction mixture was concentrated in vacuum and purified by column chromatography. Procedure B (**7**, **10**, **12**): the reaction was carried out in a round bottom flask on a magnetic stirrer. The intermediate compound (0.33 mmol), appropriate piperazine derivatives (0.4 mmol), and potassium carbonate (0.66 mmol) in acetonitrile (10 mL) were heated for 24–48 h at 80 °C, with continuous stirring. After concentration in a vacuum, extraction was carried out with methylene chloride, the organic layer was dried, concentrated, and purified using column chromatography.

The analytical results of 8-phenyl-3-(3-(4-phenylpiperazin-1-yl)propyl)-1,3-diazaspiro[4.5]decane-2,4-dione (**13**) and 3-(3-(4-(3-chlorophenyl)piperazin-1-yl)propyl)-8-phenyl-1,3-diazaspiro[4.5]decane-2,4-dione (**14**) were previously described [11]. Detailed descriptions of compounds **1**, **6**, **7**, **8**, and **12** are in Supplementary materials.

3-(2-(4-Benzhydrylpiperazin-1-yl)ethyl)-5-methyl-5-phenylimidazolidine-2,4-dione (2) White powdery crystals. Yield: 72%; mp 212–215 °C; TLC: R_f =0.31 (S₁); HPLC: t_R =1.491; MS calcd for [M+H]⁺: C₂₉H₃₂N₄O₂ m/z : 468.25, found: 469.27; ¹H NMR (300 MHz, CDCl₃-d) δ ppm 1.81 (s, 3 H, -CH₃) 2.28 (br. s., 4 H, Pip) 2.47 (br. s., 4 H, Pip) 2.57 (t, J =6.45 Hz, 2 H, -CH₂-Pip) 3.61 (t, J =6.45 Hz, 2 H, Hyd-CH₂-) 4.13 [s, 1 H, -CH-(Ph)₂] 7.13–7.20 (m, 2 H, Ph, Hyd) 7.22–7.30 (m, 5 H, Ph) 7.31–7.42 (m, 7 H, Ph) 7.47–7.54 (m, 2 H, Ph). Anal. calcd for C₂₉H₃₂N₄O₂ (468.60): C: 74.33, H: 6.88, N: 11.96; Found C: 74.08, H: 7.02, N: 11.50.

3-(3-(4-(2-Fluorophenyl)piperazin-1-yl)propyl)-5-methyl-5-phenylimidazolidine-2,4-dione (3) White powdery crystals. Yield: 83%; mp 126–129 °C; TLC: R_f =0.25 (S₁); HPLC: t_R =1.217; MS calcd for [M+H]⁺: C₂₃H₂₇FN₄O₂ m/z : 410.21, found: 411.38; ¹H NMR (300 MHz, CDCl₃-d) δ ppm 1.77–1.88 (m, 5 H, -CH₃, -CH₂-CH₂-CH₂-) 2.39 (t, J =7.03 Hz, 2 H, -CH₂-Pip) 2.56 (t, J =4.7 Hz, 4 H, Pip)



Scheme 1 The synthetic routes of novel 5-methyl-5-phenylhydantoin (series 1) and 4-phenylcyclohexane-5-spirohydantoin (series 2) derivatives. Reagents and conditions: **A** KCN, $(\text{NH}_2)_2\text{CO}_3$, 50% EtOH, 70 °C, 3 h; **B** $\text{Br}(\text{CH}_2)_n\text{Br}$, K_2CO_3 , DMF, 70 °C, 3 h; **C** arylpiperazine, TEA, MeCN, MW: 200 W, 100 °C, 65–100 min (compd **1–6**, **8**, **9**, **11**)/arylpiperazine, K_2CO_3 , 80 °C, MeCN, 24–48 h (compd **7**, **10**, **12**).

3.07 (t, $J=4.7$, 5.3 Hz, 4 H, Pip) 3.54 (t, $J=7$, 7.1 Hz, 2 H, Hyd- CH_2 -) 6.86–7.08 (m, 5 H, Ph) 7.28–7.42 (m, 3 H, Hyd, Ph) 7.47–7.54 (m, 2 H, Ph). Anal. calcd for $\text{C}_{23}\text{H}_{27}\text{FN}_4\text{O}_2$ (410.49): C: 67.30, H: 6.63, N: 13.65; Found C: 66.95, H: 6.72, N: 13.19.

3-(3-(4-(3-Fluorophenyl)piperazin-1-yl)propyl)-5-methyl-5-phenylimidazolidine-2,4-dione (4) White powdery crystals. Yield: 33%; mp 125–126 °C; TLC: $R_f=0.26$ (S_1); HPLC: $t_R=1.253$; MS calcd for $[\text{M}+\text{H}]^+$: $\text{C}_{23}\text{H}_{27}\text{FN}_4\text{O}_2$ m/z : 410.21, found: 411.18; ^1H NMR (300 MHz, CDCl_3 - d) δ ppm 1.77–1.89 (m, 5 H, $-\text{CH}_2-\text{CH}_2-\text{CH}_2-$, $-\text{CH}_3$) 2.38 (t, $J=7$, 7.1 Hz, 2 H, $-\text{CH}_2$ -Pip) 2.47–2.55 (t, $J=4.7$, 5.3 Hz, 4 H, Pip) 3.08–3.16 (t, $J=4.7$, 5.3 Hz, 4 H, Pip) 3.58 (t, $J=7.33$ Hz, 2 H, Hyd- CH_2 -) 6.46–6.67 (m, 4 H, Hyd, Ph) 7.11–7.22 (m, 1 H, Ph) 7.29–7.42 (m, 3 H, Ph) 7.47–7.53 (m, 2 H, Ph). Anal. calcd for $\text{C}_{23}\text{H}_{27}\text{FN}_4\text{O}_2 \times \text{H}_2\text{O}$ (428.51): C: 64.47, H: 6.82, N: 13.08; Found C: 64.66, H: 6.55, N: 12.63.

zine, TEA, MeCN, MW: 200 W, 100 °C, 65–100 min (compd **1–6**, **8**, **9**, **11**)/arylpiperazine, K_2CO_3 , 80 °C, MeCN, 24–48 h (compd **7**, **10**, **12**).

3-(3-(4-(4-Fluorophenyl)piperazin-1-yl)propyl)-5-methyl-5-phenylimidazolidine-2,4-dione (5) White powdery crystals. Yield: 71%; mp 141–143 °C; TLC: $R_f=0.54$ (S_2); HPLC: $t_R=1.235$; MS calcd for $[\text{M}+\text{H}]^+$: $\text{C}_{23}\text{H}_{27}\text{FN}_4\text{O}_2$ m/z : 410.21, found: 411.25; ^1H NMR (300 MHz, CDCl_3 - d) δ ppm 1.79–1.90 (m, 5 H, $-\text{CH}_3$, $\text{CH}_2-\text{CH}_2-\text{CH}_2-$) 2.41 (t, $J=7.33$ Hz, 2 H, $-\text{CH}_2$ -Pip) 2.56 (t, $J=4.1$, 5.3 Hz, 4 H, Pip) 3.07 (t, $J=4.7$, 5.3 Hz, 4 H, Pip) 3.58 (t, $J=7.03$ Hz, 2 H, Hyd- CH_2 -) 6.35 (s, 1 H, Hyd) 6.79–6.88 (m, 2 H, Ph) 6.90–6.99 (m, 2 H, Ph) 7.29–7.42 (m, 3 H, Ph) 7.45–7.53 (m, 2 H, Ph). Anal. calcd for $\text{C}_{23}\text{H}_{27}\text{FN}_4\text{O}_2$ (410.49): C: 67.30, H: 6.63, N: 13.65; Found C: 66.82, H: 6.70, N: 13.36.

3-(3-(4-(2-Fluorophenyl)piperazin-1-yl)propyl)-8-phenyl-1,3-diazaspiro[4.5]decane-2,4-dione (9) White powdery crystals. Yield: 53%; mp 179–181 °C; TLC: $R_f=0.34$ (S_1); HPLC: $t_R=1.523$; MS calcd for $[\text{M}+\text{H}]^+$: $\text{C}_{27}\text{H}_{33}\text{FN}_4\text{O}_2$ m/z : 464.26, found: 465.28; ^1H NMR (300 MHz, CDCl_3 - d) δ ppm 1.66–1.82 (m, 4 H, Cyclohex-

ane) 1.75 (q, $J=2.9, 11.7, 11.8$ Hz, 4 H, Cyclohexane) 1.85–2.14 (m, 7 H, Cyclohexane, $-\text{CH}_2-\text{CH}_2-\text{CH}_2-$, Ph-CH-) 2.47 (t, $J=7.33$ Hz, 2 H, $-\text{CH}_2$ -Pip) 2.61 (t, $J=4.1$ Hz, 4 H, Pip) 3.08 (t, $J=4.1, 4.6$ Hz, 4 H, Pip) 3.65 (t, $J=7.03$ Hz, 2 H, Hyd- CH_2 -) 6.87–7.07 (m, 4 H, Ph) 7.19–7.25 (m, 1 H, Ph) 7.30–7.36 (m, 4 H, Ph) 8.06 (s, 1 H, Hyd). Anal. calcd for C₂₃H₂₅N₄O₃F (424.48): C: 69.80, H: 7.16, N: 12.06; Found C: 69.58, H: 7.22, N: 12.18.

3-(3-(4-(3-Fluorophenyl)piperazin-1-yl)propyl)-8-phenyl-1,3-diazaspiro[4.5]decane-2,4-dione (10) White powdery crystals. Yield: 52%; mp 193–195 °C; TLC: $R_f=0.37$ (S₁); HPLC: $t_R=1.546$; MS calcd for $[\text{M}+\text{H}]^+$: C₂₇H₃₃FN₄O₂ m/z : 464.26, found: 465.27; ¹H NMR (300 MHz, CDCl₃-*d*) δ ppm 1.74 (q, $J=3.5, 11.7, 11.8$ Hz, 4 H, Cyclohexane) 1.86–2.09 (m, 7 H, Cyclohexane, $-\text{CH}_2-\text{CH}_2-\text{CH}_2-$, Ph-CH-) 2.44 (t, $J=7.33$ Hz, 2 H, $-\text{CH}_2$ -Pip) 2.55 (t, $J=4.7, 5.3$ Hz, 4 H, Pip) 3.16 (t, $J=4.7, 5.3$ Hz, 4 H, Pip) 3.65 (t, $J=7.03$ Hz, 2 H, Hyd- CH_2 -) 6.47–6.66 (m, 3 H, Ph) 7.12–7.25 (m, 2 H, Ph) 7.30–7.35 (m, 4 H, Ph) 8.05 (s, 1 H, Hyd). Anal. calcd for C₂₃H₂₅N₄O₃F (424.48): C: 69.80, H: 7.16, N: 12.06; Found C: 69.92, H: 7.34, N: 11.86.

3-(3-(4-(4-Fluorophenyl)piperazin-1-yl)propyl)-8-phenyl-1,3-diazaspiro[4.5]decane-2,4-dione (11) White powdery crystals. Yield: 60%; mp 216–217 °C; TLC: $R_f=0.31$ (S₁); HPLC: $t_R=1.526$; MS calcd for $[\text{M}+\text{H}]^+$: C₂₇H₃₃FN₄O₂ m/z : 464.26, found: 465.28; ¹H NMR (300 MHz, CDCl₃-*d*) δ ppm 1.70–1.84 (q, $J=9.4, 12.9, 14.6$, 4 H, Cyclohexane) 1.85–2.13 (m, 7 H, Cyclohexane, $-\text{CH}_2-\text{CH}_2-\text{CH}_2-$, Ph-CH-) 2.45 (t, $J=7.33$ Hz, 2 H, $-\text{CH}_2$ -Pip) 2.56 (t, $J=4.7$ Hz, 4 H, Pip) 3.07 (t, $J=4.7$ Hz, 4 H, Pip) 3.65 (t, $J=7.03$ Hz, 2 H, Hyd- CH_2 -) 6.78–6.87 (m, 2 H, Ph) 6.88–6.99 (m, 2 H, Ph) 7.18–7.25 (m, 1 H, Ph) 7.28–7.39 (m, 4 H, Ph) 8.38 (s, 1 H, Hyd). Anal. calcd for C₂₃H₂₅N₄O₃F (424.48): C: 69.80, H: 7.16, N: 12.06; Found C: 69.87, H: 7.32, N: 11.81.

Radioligand binding assay

Preparation of solutions of test and reference compounds

Stock solutions of tested compounds (10 mM) were prepared in DMSO. Serial dilutions of compounds were prepared in 96-well microplate in assay buffers using an automated pipetting system epMotion 5070 (Eppendorf). Each compound was tested in eight concentrations from 10⁻⁵ to 10⁻¹² M (final concentration).

5-HT_{2A} receptor binding assay

Radioligand binding was performed using membranes from CHO-K1 cells stably transfected with the human 5-HT_{2A} receptor (PerkinElmer). All assays were carried

out in duplicates. Data were fitted to a one-site curve-fitting equation with Prism 6 (GraphPad Software) and K_i values were estimated from the Cheng – Prusoff equation. The full description of this assay is in Supplementary material.

Functional bioassays

In vitro functional bioassays at cells transfected with human 5HT_{2A} receptor (aequorin and luminescence-based intracellular calcium assay)

Tested and reference compounds were dissolved in DMSO at a concentration of 10⁻² M. Serial dilutions were prepared in 96-well microplate in assay buffer and eight to ten concentrations were tested.

A cellular aequorin-based functional assay was performed with recombinant Chinese hamster ovary cells expressing mitochondrially targeted aequorin, human G-protein-coupled receptors, and the promiscuous G protein $\alpha 16$ for 5-HT_{2A}. The full description of this assay is in Supplementary materials.

Functional bioassays at 5HT_{2A}-receptors in isolated rat aorta

Isolated rat aorta was used to determine the antagonistic activity of studied compounds for 5HT_{2A}-receptors. The male Wistar rats weighting 200–350 g were anesthetized with thiopental sodium (75 mg/kg i.p.) and the aorta was dissected and placed in a Krebs–Henseleit solution (NaCl 118 mM, KCl 4.7 mM, CaCl₂ 2.25 mM, MgSO₄ 1.64 mM, KH₂PO₄ 1.18 mM, NaHCO₃ 24.88 mM, glucose 10 mM, C₃H₃O₃Na 2.2 mM, EDTA 0.05 mM), denuded of the endothelium, cleaned of surrounding fat tissues and cut into 4-mm-long rings. The aortic rings were incubated in 30 ml chambers filled with a Krebs–Henseleit solution at 37 °C and pH 7.4 with constant oxygenation (O₂/CO₂, 19:1). Two stainless steel stirrups were inserted through the lumen of each aortic segment: one stirrup was attached to the bottom of the chamber and the other to an isometric FDT10-A force–displacement transducer (BIOPAC Systems, Inc., COMMAT Ltd., Turkey). The aortic rings were stretched and maintained at optimal tension of 2 g and allowed to equilibrate for 2 h.

After the equilibration period, the aortic rings were contracted to maximal tension with KCl (60 mM). The depolarizing solution KCl (60 mM) was obtained by equimolar substitution of NaCl for KCl. Then the cumulative concentration–response curves to 5-HT were determined in the absence and presence of antagonist. Tissues were incubated with antagonists for 30 min.

Concentration–response curves were analyzed using GraphPad Prism 6.0 software (GraphPad Software Inc.,

San Diego, CA, USA). Contractile responses to 5-HT (in the presence or absence of tested compounds) are expressed as a percentage of the maximal KCl effect reached in the concentration–response curves obtained before incubation with the tested compounds ($E_{\max} = 100\%$). Data are the mean \pm SEM of three separate experiments. The affinity was estimated with the equation $pK_B = \log(\text{concentration ratio} - 1) - \log(\text{molar antagonist concentration})$, where the concentration ratio is the ratio of equi-effective agonist concentrations in the absence and in the presence of the antagonist.

In vitro whole blood aggregation tests

In vitro aggregation assays were performed using freshly drawn whole rat blood with a Multiplate platelet function analyzer (Roche Diagnostic), the five-channel aggregometer based on measurements of electric impedance. Blood was collected from carotid arteries with a hirudin blood tube (Roche Diagnostic). 300 μ l of hirudin anticoagulated blood was mixed with 300 μ l prewarmed isotonic saline solution containing studied compound or vehicle (deionized water or DMSO 0.1%) and preincubated for 3 min at 37 °C with continuous stirring. Aggregation was induced by adding collagen (Hyphen-Biomed, France) (final concentration 1.6 μ g/ml), 5-HT (30 μ M) and sub-threshold concentration of collagen (0.8 μ g/ml), 5-HT (6 μ M) and sub-threshold concentration of ADP (ADP test, Roche Diagnostic), (1.6 μ M). The aggregation process was recorded for 6 min. Ketanserin (ketanserin (+) tartrate salt, Sigma-Aldrich, Germany), sarpogrelate (sarpogrelate hydrochloride, Sigma-Aldrich, Germany), and aspirin (Tocris, UK) were used as reference compounds. The Multiplate software analyzed the area under the curve (AUC) of the clotting process of each measurement and calculated the mean values. Each concentration of studied compounds was tested at least three times. The exact sample size (n value) is presented on Figs. 2–4. Concentration-inhibition curves were constructed and analyzed by non-linear curve fitting using GraphPad Prism 6.0 (GraphPad Software Inc., San Diego, CA, USA).

Statistical analysis

Data were presented as mean \pm standard error the mean (SEM). Statistical comparisons were made by the one-way analysis of variance (one-way ANOVA) and the significance of the differences between the control group and treated groups was determined by the Dunnett post hoc test. $p < 0.05$ was considered significant.

Molecular modeling studies

Molecular docking was performed to the previously developed homology models of the 5-HT_{2A} receptor. The

procedure for obtaining ligand-optimized models of high predictive value was characterized in detail previously [17, 18]. The 5-HT_{2A} receptor model was based on the 5-HT_{2B} receptor crystal structure 4IB4 [19]. Glide XP flexible docking was carried out using the OPLS3 force field and default parameters. H-bond constraint, as well as centroid of a grid box for docking to the receptor models were located on Asp86^{3.32}. The selection of the poses was based on the prevalence of well-scored (glide gscore) complexes and visual evaluation of binding interactions. Ligand structures were optimized using LigPrep. The presented computational tools were implemented in Small-Molecule Drug Discovery Suite (Schrodinger, Inc.), which was licensed for Jagiellonian University Medical College.

In silico toxicity prediction

In silico toxicity was assessed with a novel approach (pkCSM) which uses graph-based signatures to develop predictive models of ADMET properties or toxicity of introduced structures [20].

Results and discussion

Chemistry

The final compounds (series 1 and 2) were synthesized as outlined in Scheme 1. The substituted imidazolidine-2,4-dione derivatives (A and C) used as starting compounds were prepared from 1-phenylethanone or 4-phenylcyclohexanone, respectively, by the Bucherer–Bergs reaction followed by N3-alkylation with dihalogenoalkanes in the hydantoin ring. The resulting intermediates (B and D) were coupled with appropriate arylpiperazines in the presence of potassium carbonate or triethylamine to obtain the final compounds. The crude final products were purified using column chromatography. The structure and purity of the synthesized compounds were confirmed by spectral and chromatographic analyses, and the reaction progress was monitored by thin-layer chromatography. The detailed physicochemical and spectral data are summarized in the Experimental Part.

In vitro 5-HT_{2A} receptor activity of compound 1–14

The novel series of compounds targeting serotonin 5-HT_{2A} receptors were designed by coupling the amide part of 4-phenylcyclohexane-5-spiro- or 5-methyl-5-phenylimidazolidine-2,4-dione with various arylpiperazine moieties, using linkers of different lengths. The heterocyclic imide ring, or substituted imidazolidine-2,4-dione in this case, seemed to be a good starting point for designing 5-HT_{2A} receptor ligands because it is also present in the structures

of the conventional 5-HT_{2A} receptor antagonists such as ketanserin and fananserin (Fig. 1). In the amine part, the piperazine analogs (4-fluorophenyl)(piperidin-4-yl)methanone and 4-(bis(4-fluorophenyl)methylene)-piperidine, found in ketanserin and ritanserin, were introduced. Furthermore, a key structural fragment of fananserin, 4-fluorophenyl-1-piperazine, was chosen as the amine part, as well as its analogs with a fluorine atom at positions 2 and 3.

In the synthesized series, ten compounds (**2–6**, **9–14**) displayed low-to-high binding affinity for the 5-HT_{2A} receptors ($K_i = 21$ –2584 nM, Table 1). Compounds containing phenyl(piperazin-1-yl)methanone (**1**, **6**, **7**, **12**) or 1-(diphenylmethyl)piperazine (**2**) fragment (except compound **2**) showed low or no affinity, while compound **2**, which is a 1-(diphenylmethyl)piperazine derivative with

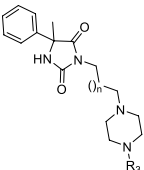
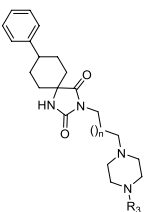
5-methyl-5-phenylimidazolidine-2,4-dione moiety, showed a moderate affinity for the tested serotonin receptors ($K_i = 450$ nM, Table 1). However, the replacement of the aforementioned piperazines by fluorophenylpiperazine led to an increase in their affinity. Interestingly, compounds **5** and **11** containing a 4-fluorophenyl-1-piperazine fragment, which is present in fananserin, as well as compounds **13** and **14** having 1-phenylpiperazine and 3-chlorophenyl-1-piperazine moiety, respectively, displayed the highest affinity for the 5-HT_{2A} receptors in the prepared series ($K_i = 21$ –50 nM). Already published compounds **13** and **14** showed selectivity for serotonin 5-HT_{1A} and dopamine D₂ receptors [11] (for details see Supplementary materials Table S1). Moreover, the position of the fluorine atom in the 1-phenylpiperazine fragment seemed to determine the affinity of compounds for 5-HT_{2A} receptors as shifting the fluorine atom of this fragment from position 4 to position 3 or 2 caused a significant (6- to 10-fold) decrease in affinity (Table 1). The influence of the linker length on the compounds' affinity for 5-HT_{2A} receptors was also investigated. It was observed that compounds with a linker having two methylene groups showed no affinity for the receptors (except compound **2**), but the elongation of the linker to three methylene groups resulted in an increased binding affinity.

For further characterization of the compounds, K_i (5-HT_{2A}) < 1000 nM was set as selection criteria. Based on this, compounds **2–5**, **9–11** were chosen for further determination of functional activity for the 5-HT_{2A} receptors. The binding affinity of compounds **13** and **14** has been studied and published in our previous work [11].

Biofunctional assays

The antagonistic activity of compounds **2–5**, **9–11** for 5-HT_{2A} receptors was analyzed by evaluating the inhibition of serotonin-induced contraction in rat aorta. The tested compounds shifted the 5-HT concentration–response curve to the right without lowering the maximal response, which indicated that they are competitive antagonists of the rat 5-HT_{2A} receptors (Figure S2). The affinities of these compounds were expressed as pK_B values, estimated from the 5-HT concentration–response curves and a single concentration of the compounds. The pK_B values ranged from 6.161 (compound **2**) to 7.430 (compound **11**) (Table 2) and were in line with the radioligand binding data. Additionally, compounds **2–5**, **9**, and **11** were studied in cells transfected with human 5-HT_{2A} receptors. The results of this bioassay revealed that compounds **3** and **11** are moderate antagonists of the human 5-HT_{2A} receptors ($pK_B \approx 6.758$, Table 2). The antagonistic activity of compounds **13** and **14** has been studied and published in our previous work [11].

Table 1 The radioligand binding data of 4-phenylcyclohexane-5-spirohydantoin and 5-methyl-5-phenylhydantoin derivatives and reference drug (mianserin) for serotonin 5-HT_{2A} receptors

Compd	<i>n</i>	R	K_i [nM] ± SEM
	1	0	–CO ₄ FPh nd
	2	0	–CH(Ph) ₂ 450.0 ± 43.9
	3	1	–2FPh 516.0 ± 52.0
	4	1	–3FPh 340.0 ± 57.0
	5	1	–4FPh 48.0 ± 4.9
	6	1	–CO ₄ FPh nd
	7	0	–CO ₄ FPh nd
	8	0	–CH(Ph) ₂ nd
	9	1	–2FPh 220.0 ± 19.6
	10	1	–3FPh 297.0 ± 19.9
	11	1	–4FPh 21.0 ± 1.7
	12	1	–CO ₄ FPh 2584.0 ± 302.0
	13 (Compd A)^{a,b}	1	–Ph 20.0 ± 4.8 ^a
	14^a	1	–3ClPh 24.0 ± 2.5 ^a
Mianserin			1.2 ± 0.2

The data for **13**, **14**, were already published in [11] and re-printed in the current manuscript for comparison, with permission from Acta Poloniae Pharmaceutica—Drug Research

nd no affinity detected

^aThe data for **13**, **14**, were already published in [11] and re-printed in the current manuscript for comparison, with permission from Acta Poloniae Pharmaceutica—Drug Research

^bCompd A Fig. 1

Table 2 In vitro functional antagonist data of tested 4-phenylcyclohexane-5-spirohydantoin and 5-methyl-5-phenylhydantoin derivatives and reference drugs (ketanserin, sarpogrelate) for 5-HT_{2A} receptors [a] in rat aorta and [b] in cells expressing human 5-HT_{2A} receptors

Compd	pK _B ± SEM	
	a	b
2	6.161 ± 0.06	5.706 ± 0.21
3	6.666 ± 0.12	6.758 ± 0.15
4	6.357 ± 0.18	5.035 ± 0.23
5	6.957 ± 0.23	5.032 ± 0.30
9	7.018 ± 0.08	5.702 ± 0.16
10	6.300 ± 0.36	nt
11	7.430 ± 0.32	6.757 ± 0.06
13 (Compd A)	7.665 ± 0.03 ^a	nt
14	7.110 ± 0.05 ^a	nt
Ketanserin	9.439 ± 0.09 ^b	7.839 ± 0.03 ^b
Sarpogrelate	8.220 ± 0.08 ^b	7.296 ± 0.04 ^b

nt not tested

^{a,b}The data for ^a**13**, **14**, as well as for ^bketanserin and sarpogrelate were already published in [7, 11], respectively, and re-printed in the current manuscript for comparison, with permission from European Journal of Pharmacology and Acta Poloniae Pharmaceutica—Drug Research

In vitro whole blood aggregation tests

Compounds with confirmed 5-HT_{2A} antagonistic activity (**2–5**, **9–11**, **13**, **14**) were subjected to further studies. First, to evaluate the antiplatelet effect, freshly isolated rat whole blood was incubated with the selected compounds (3–200 μM) or vehicle (DMSO), and the aggregation responses were recorded. Platelet aggregation was triggered by collagen or using collagen or ADP at the subthreshold concentration and 5-HT. Ketanserin and sarpogrelate, which

exert antiplatelet activity due to 5-HT_{2A} antagonism as well as aspirin (COX-1 inhibitor) were used as reference compounds for this analysis.

When studying aggregation, various agonists can be used individually to identify a pathway that may be potentially affected by a test compound. Collagen is one of the most important, natural platelet agonist. The final effect of collagen stimulation is increased intracellular Ca²⁺ concentration, which leads to morphological changes, ADP and 5-HT secretion, and the synthesis of thromboxane A2 (TXA2) [21]. Therefore using collagen as an aggregation agonist, provides a general means of assessing platelet function and anti-aggregation effect of test compounds in vitro [21]. Figure 2 shows the effect of compound **13**, ketanserin, sarpogrelate, and aspirin on aggregation induced by collagen. In our study, only aspirin and the most potent 5-HT_{2A} receptor antagonists were able to inhibit collagen-induced aggregation. These compounds significantly decreased platelet aggregation ($F_{12,38} = 9.873$, $p < 0.0001$). Among the obtained compounds, only compound **13**, which was identified as the most potent 5-HT_{2A} antagonist, was able to inhibit collagen-stimulated aggregation of whole rat blood ($IC_{50} = 27.3 \pm 3.8$ μM). Its anti-aggregation effect was greater than that of sarpogrelate ($IC_{50} = 66.8 \pm 12.9$ μM) and was comparable to ketanserin ($IC_{50} = 32.1 \pm 3.0$ μM). For comparison, the IC_{50} value for aspirin was equal to 14.5 ± 1.1 μM (Table 3, Fig. 2).

Further, the antiplatelet activity was studied using 5-HT as a co-agonist of aggregation induced by collagen or ADP at the sub-threshold concentration. Physiologically, many factors that induce platelet aggregation act synergistically. 5-HT itself is only a weak platelet activator, but it can enhance the aggregation process induced by collagen or ADP, thus activating platelet 5-HT_{2A} receptors [5, 22, 23]. Collagen used at a concentration of 0.8 μg/ml did not induce aggregation of whole rat blood, while 5-HT used

Fig. 2 Effects of compound **13**, sarpogrelate, ketanserin and aspirin on in vitro whole rat blood aggregation induced by collagen (1.6 μg/ml). Results are expressed as mean ± SEM, *** $p < 0.001$, **** $p < 0.0001$ versus control group (0.1% DMSO in saline), one-way ANOVA, post hoc Dunnet test. AUC area under curve

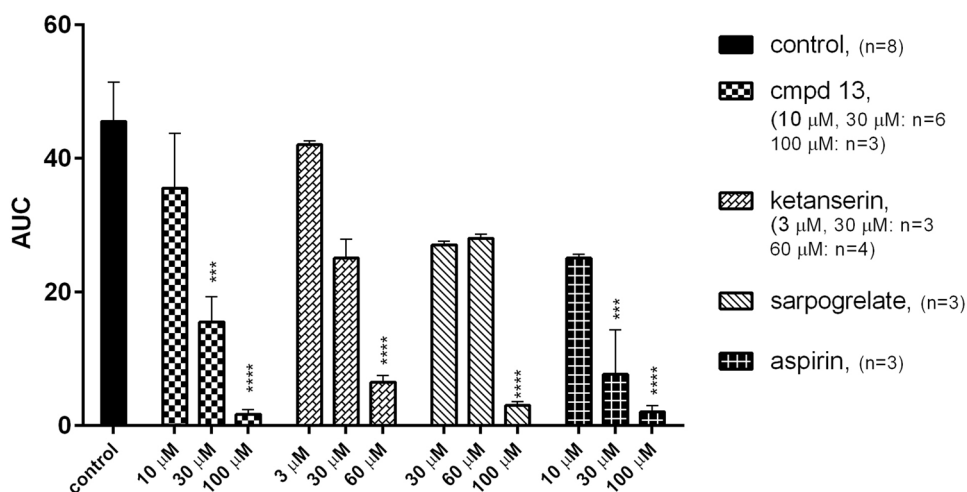


Table 3 Potencies of studied and reference compounds in inhibiting whole rat blood aggregation in vitro induced by [a] collagen (1.6 µg/ml), [b] 5-HT (30 µM) and collagen (0.8 µg/ml), [c] 5-HT (6 µM) and ADP (1.6 µM)

Compound	a (collagen) IC ₅₀ [µM]	b (5-HT + collagen) IC ₅₀ [µM]	c (5-HT + ADP) IC ₅₀ [µM]
2	na	139.8 ± 18.9	na
3	na	17.4 ± 10.9	37.9 ± 6.5
4	na	153.1 ± 26.4	83.1 ± 8.1
5	na	127.2 ± 32.7	123.7 ± 36.2
9	na	138.6 ± 16.1	na
10	na	94.3 ± 53.5	118.6 ± 32.1
11	na	91.3 ± 7.1	140.5 ± 44.6
13 (Compd A)	27.3 ± 3.8	53.3 ± 4.6	9.9 ± 2.5
14	na	75.8 ± 16.6	93.8 ± 9.4
Sarpogrelate	66.8 ± 12.9	nt	22.7 ± 2.1 ^a
Ketanserin	32.1 ± 3.0	1.3 ± 0.0	9.7 ± 2.1 ^a
Aspirin	14.5 ± 1.1	nt	nt

na not active, nt not tested

^aThe data for ketanserin and sarpogrelate were already published in [7] and re-printed in the current manuscript for comparison, with permission from European Journal of Pharmacology

alone could not cause aggregation at any concentration tested. However, the combination of 5-HT at a concentration of 30 µM with collagen at the subliminal concentration resulted in the maximal aggregation response (Fig. 3). This aggregation effect was attenuated by the studied compounds (2–5, 9–11, 13, 14) and ketanserin [($F_{31,84} = 7.808$, $p < 0.0001$)], (Fig. 3). The obtained IC₅₀ values ranged from 17.4 ± 10.9 µM (compound 3) to 153.1 ± 26.4 µM (compound 4), (Table 3). The reference compound ketanserin inhibited blood aggregation with an IC₅₀ value

of 1.3 ± 0.01 µM, which confirmed its high antagonistic activity for 5-HT_{2A} receptors found in functional studies.

The tested compounds also inhibited 5-HT-amplified and ADP-induced platelet aggregation. At a concentration of 1.6 µM, ADP used alone could cause only partial and transient aggregation of rat blood in vitro, while 5-HT used alone did not induce aggregation at any concentration tested. However, combining 5-HT at a concentration of 6 µM with ADP at the sub-threshold concentration resulted in a maximal aggregation response. The serotonin-mediated amplification of ADP-stimulated aggregation was attenuated when rat blood was preincubated with compounds 3, 4, 5, 10, 13, and 14 ($F_{24,54} = 20.60$, $p < 0.0001$), with the IC₅₀ values ranging from 9.9 ± 2.5 µM (compound 13) to 140.5 ± 44.6 µM (compound 11). For comparison, the IC₅₀ value of ketanserin was similar to that of compound 13 and was equal to 9.7 ± 2.1 µM [7]. As observed for induction with collagen alone, compound 13 was more potent than sarpogrelate, which IC₅₀ value, found in our previous work was equal to 22.7 ± 2.1 [7]. On the other hand, even at a concentration of 200 µM, compound 2 did not exhibit any significant inhibitory effect on ADP- and 5-HT-induced blood aggregation, which proved its lowest intrinsic activity toward 5-HT_{2A} receptors found in the biofunctional study. Besides compound 13, compound 3 was observed to be the most potent in the entire series of newly synthesized compounds, with an IC₅₀ value of 17.4 ± 10.9 µM against collagen- and 5-HT-induced blood aggregation and a value of 37.9 ± 6.5 µM against ADP- and 5-HT induced blood aggregation. Thus, the effect of compound 3 on ADP- and 5-HT-induced aggregation was comparable to that of sarpogrelate. The results of this analysis are presented in Fig. 4 and Table 3.

It was found that the most potent 5-HT_{2A} receptor antagonists—compound 13, ketanserin, and

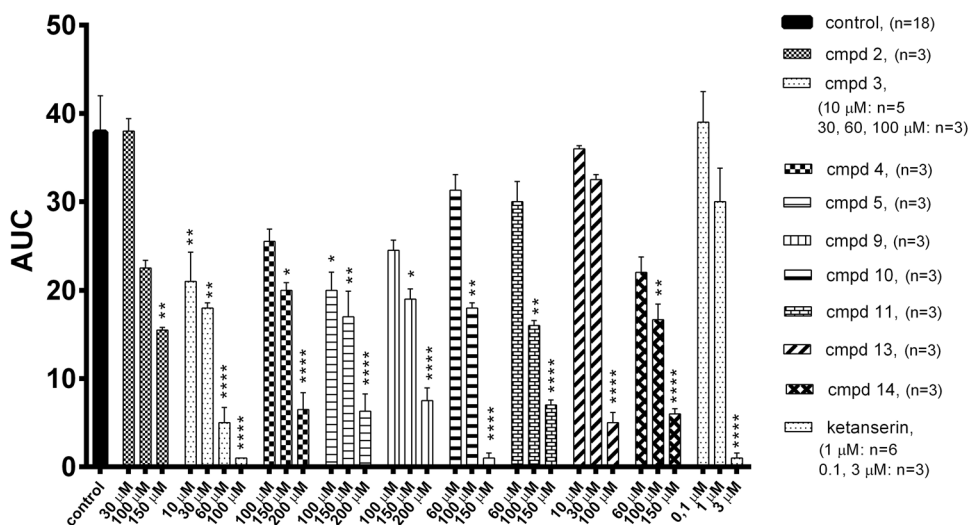
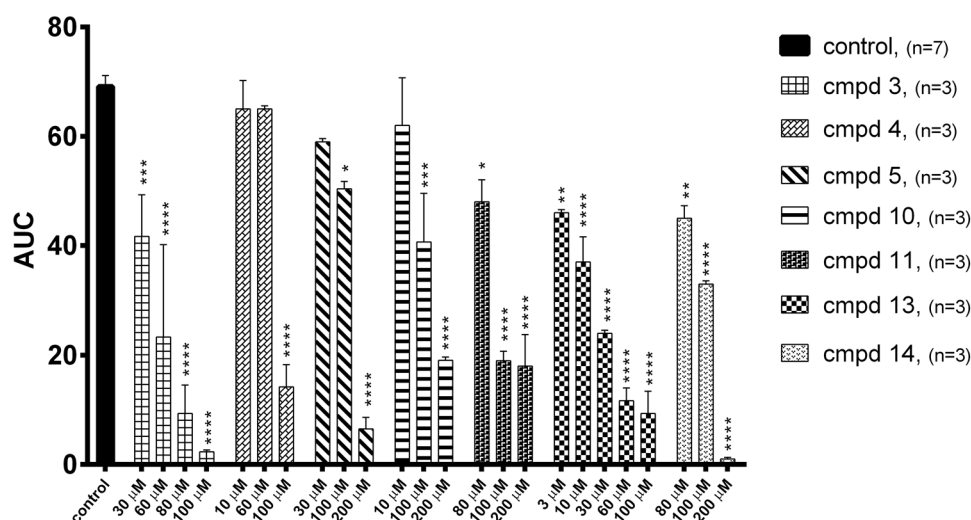
Fig. 3 Effects of compounds 2, 3, 4, 5, 9, 10, 11, 13, 14 and ketanserin on in vitro whole rat blood aggregation induced by simultaneous addition of 5-HT (30 µM) and collagen (0.8 µg/ml). Results are expressed as mean ± SEM, * $p < 0.05$, ** $p < 0.01$, *** $p < 0.0001$ versus control group (0.1% DMSO in saline), one-way ANOVA, post hoc Dunnett test. AUC area under curve

Fig. 4 Effects of compounds **3**, **4**, **5**, **10**, **11**, **13**, **14** on in vitro whole rat blood aggregation induced by simultaneous addition of 5-HT (6 μ M) and ADP (1.6 μ M). Results are expressed as mean \pm SEM, * p < 0.05, ** p < 0.01, *** p < 0.001, **** p < 0.0001 versus control group (0.1% DMSO in saline), one-way ANOVA, post hoc Dunnett test. AUC area under curve



sarpogrelate—inhibited collagen-induced platelet aggregation, whereas other 5-HT_{2A} antagonists—compounds **2–5**, **9–11**, and **14**—inhibited 5-HT-potentiated platelet aggregation. Among these, compound **3** was the most active and inhibited collagen- or ADP-induced and 5-HT-amplified aggregation with the lowest IC₅₀ values.

Molecular modeling studies

The lead compounds **3** and **13**, although having different arylpiperazine substitutions, both showed a significant affinity for 5-HT_{2A} receptors in the radioligand binding assays. Nevertheless, compared to compound **3**, compound **13** showed over 25-fold higher affinity for the target protein. To analyze this difference at the molecular level, docking to the homology model was performed. In the case of both compounds, the arylpiperazine moiety bound in the orthosteric binding site, as expected for the monoaminergic ligands. This moiety was anchored through the salt bridge (charge-assisted hydrogen bond) between the basic nitrogen atom and Asp86^{3,32}, as well as through CH– π stacking between the aromatic ring and Phe228^{6,52} (Fig. 5A). The hydantoin-containing fragments of the compounds were found to be located in the accessory cavity of the receptor. In the case of compound **13** (but not compound **3**), the spiro-cyclohexane structure in this region provided a favorable conformation, allowing the formation of π – π interactions between the phenyl ring and Tyr70. Moreover, the hydantoin itself formed an H-bond with Ser157 and overlapped with 1,2,3,4-tetrahydroquinazoline-2,4-dione moiety of the reference 5-HT_{2A} receptor ligand ketanserin (Fig. 5B). The above relationships may help in the design of further 5-HT_{2A} receptor ligands of the proposed chemotype.

In silico toxicity prediction

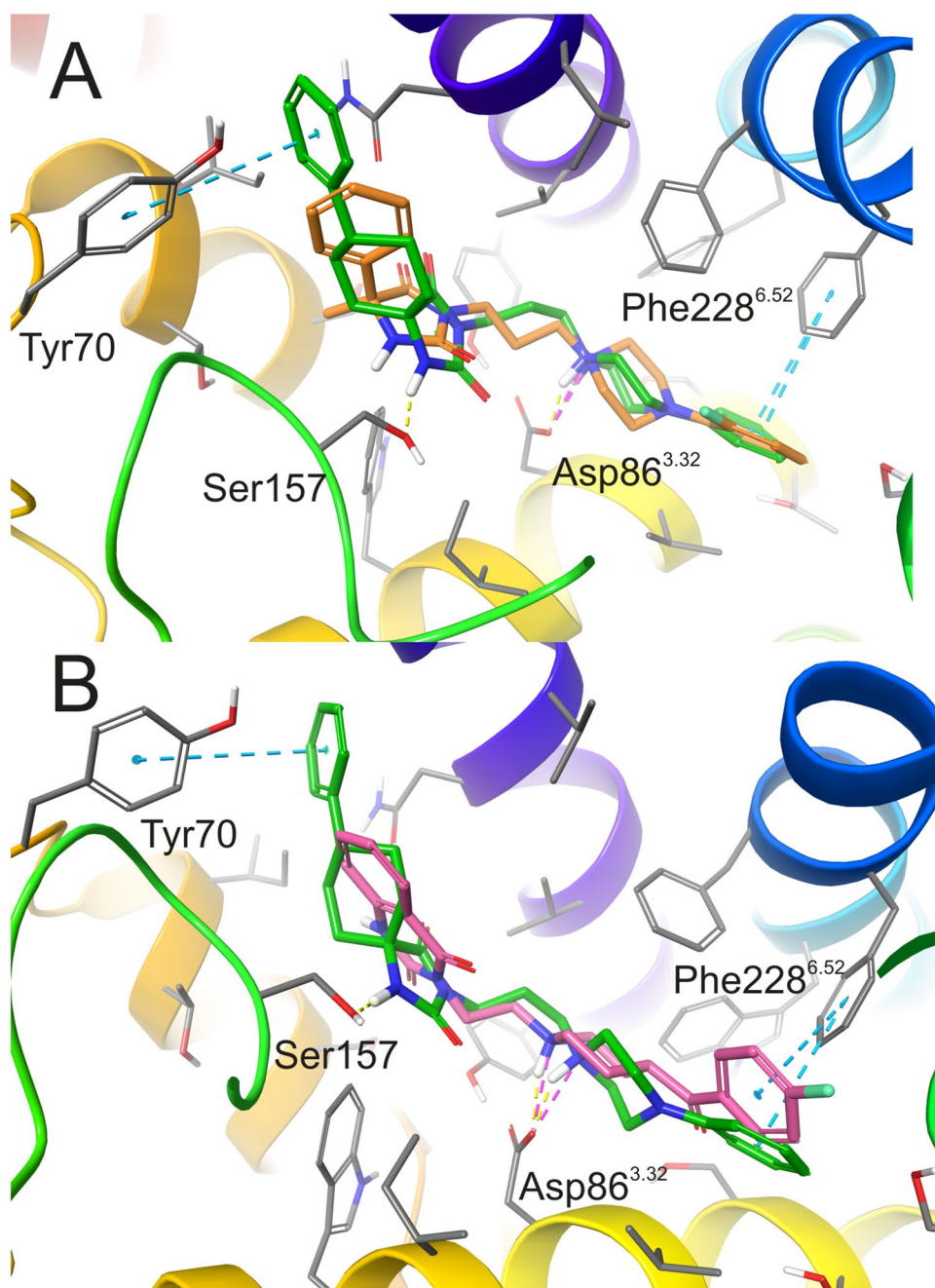
In preliminary studies, the balance between activity, potency, pharmacokinetics, and toxicity is crucial for effective drug candidates. Therefore, computational approaches to optimize pharmacokinetics and toxicity properties have been developed that allow the initial estimation of these parameters for the newly synthesized structures. One of these predictive models was used to assess the *in silico* toxicity of the most active compounds **2–5**, **9–11**, **13**, and **14** (for detailed information please see Table S2 in the Supplementary materials) with a hydantoin core.

Based on the prediction results, the tested compounds showed no toxicity in the AMES test, which measures the potential mutagenic activity. Moreover, one of the hERG predictive models indicated no cardiotoxicity of the tested compounds, while the other suggested a possible interaction with the potassium channel. Besides, this model predicted the possibility of hepatotoxicity of some compounds, but these results require further evaluation in *in vitro* tests.

Conclusion

A series of compounds with 5-methyl-5-phenylhydantoin and 4-phenylcyclohexane-5-spirohydantoin core connected to arylpiperazine moiety via a methylene linker were designed and synthesized (**1–14**) as potential 5-HT_{2A} receptor antagonists. The analysis of binding affinity and functional bioassays showed that a majority of these compounds exhibited moderate affinity and antagonistic activity for rat and human serotonin 5-HT_{2A} receptors, which further led to the evaluation of their anti-aggregation effect. In the entire series, compound **13** was found to be the most potent, as it inhibited collagen-stimulated aggregation of

Fig. 5 The predicted binding mode of compound **13** (green) displayed together with (A) compound **3** (orange) and (B) reference compound ketanserin (pink) in the 5-HT_{2A} receptor homology model based on 4IB4. The arylpiperazine moieties interact with Asp86^{3.32} (salt bridge/charge-reinforced hydrogen bond) and Phe228^{6.52} (CH- π stacking) while hydantoin-containing fragment of compound **13** interacts exclusively with Tyr70 (π - π stacking) and Ser157 (H-bond) from extracellular loops 1 and 2, respectively. Amino acid residues engaged in ligand binding (within 4 Å from the ligand atoms) are represented as sticks



whole rat blood. The anti-aggregation effect of compound **13** was comparable to that of ketanserin and greater than sarpogrelate. The tested compounds were also analyzed for their inhibitory effect on 5-HT-amplified, and ADP- or collagen-induced platelet aggregation. The 5-HT-dependent aggregation studies showed compounds **3** and **13** as the most active. From the results of this study, it can be concluded that 5-HT_{2A} antagonists may effectively inhibit platelet aggregation and are an interesting target for the development of novel antiplatelet agents with an alternative mechanism of action.

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Author contributions Substantial contributions to the conception or design of the work ACz MK AB; or the acquisition and funding ACz MK MP BF, analysis, or interpretation of data for the work ACz MK AB AS MKoI; drafting the work or revising it critically for important intellectual content ACz MK AB; final approval of the version to be published; and ACz MK, agreement to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved ACz MK AS AB.

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Declarations

Conflict of interest The authors declare no conflict of interest relevant to the ideas or the contents of this manuscript.

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