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Identification of the first surrogate agonists for the G protein-coupled receptor GPR132†

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GPR132 is an orphan class A G protein-coupled receptor. It has been proposed to be activated by protons and to regulate apoptosis, atherosclerosis and inflammation, but these results are still preliminary. In the current work, we designed and screened a focused compound library using a β-arrestin recruitment assay, and thereby identified the first disclosed surrogate GPR132 agonist 1 with a potency of 3.4 μM. This constitutes the first available pharmacological tool for the in vitro characterization of the orphan receptor GPR132. The testing of 32 analogs furthermore identified a number of compounds with lower activity – of which six were agonists and two were antagonists – that were used to construct preliminary structure–activity relationships. Docking followed by a molecular dynamics simulation of compound 1 in a structural model of GPR132 displayed the putative interactions for the key ligand functionalities.

Introduction

G protein-coupled receptors (GPCRs) constitute the largest family of cell surface receptor proteins. They are involved in the regulation of a wide range of physiological processes including vision, metabolism, neurotransmission and inflammation, and approximately 30% of FDA-approved drugs are thought to target GPCRs. More than 800 different sequences in the human genome encode for GPCRs, of which approximately 350 sequences encode for non-olfactory receptors. Their involvement in various physiological processes has made GPCRs one of the most pursued target classes in academia as well as industry. GPR132 has been ascribed several putative endogenous ligands. In 2001, lysophosphatidylethanolamine was presented as an alternative endogenous agonist, but this could not be confirmed in another study. Independent analyses have instead provided evidence that GPR132 functions as a sensor for extracellular pH. This is in agreement with a phylogenetic analysis that grouped GPR132 with the proton-sensing receptors GPR4, GPR65, and GPR68.

Surrogate GPR132 ligands were reported by Invitrogen Discovery Sciences and Amgen in 2009, but without disclosing their structures. The physiological function of GPR132 is still unknown, although several hypotheses have been presented. Firstly, studies in this direction found that GPR132 triggers chemotaxis of monocyte phagocytic cells towards apoptotic cell remnants and promote anti-inflammatory responses in human skin epidermal cells and keratocytes by suppressing DNA synthesis and freezing specific cell cycle phases. Secondly, GPR132 is expressed on macrophages, and has been suggested to be involved in atherosclerotic lesion growth in the aortic tissue and the coronaries. However, later studies display ambiguous results as the formation of atherosclerosis lesions was suppressed and promoted by knockout and silencing of the GPR132 gene, respectively. Finally, over-expression of GPR132 has been found to inhibit tyrosine kinases involved in the activation of BCR/ABL fusion proteins, which play a key role in the oncogenesis and proliferation of some cancers.

In this article, we now disclose the first surrogate agonist for GPR132 with potency in the single micro-molar range (Fig. 1, Chart 1). Receptor structure modeling and ligand docking studies present the putative molecular mechanism of action, which could contribute to future structure-based optimization towards in vivo application. The description of the first publically available GPR132 surrogate agonist is an important first step towards the pharmacological and physiological characterization of the receptor.

Materials and methods

GPCR-focused compound libraries

We designed focused GPCR screening libraries based on identification of novel privileged structures, which are core scaffolds
Pharmacological assay
The G protein signaling pathways of many orphan GPCRs are still unknown, making second messenger-based assays unsuitable for screening for orphan receptor ligands. Thus, the β-arrestin2 recruitment assay (DiscoveRx) was employed in the screening and the subsequent hit validation. Since recruitment of β-arrestins constitutes an universal “off-switch” for GPCRs, assays based on β-arrestins binding to the activated receptors should offer the possibility of detecting GPCRs activation regardless of the routes of G protein signaling. DiscoveRx β-arrestin recruitment assays have been used previously for identification of endogenous and surrogate ligands of orphan GPCRs.

24 hours before performing the assay, the PathHunter β-arrestin recruitment assays, in CHO cell lines expressing human GPR132 (A) or β2-adrenergic receptor (B and C). All data represent mean ± SD from one representative out of five (A) or two (B and C) independent experiments performed in triplicates.

with known activity at multiple diverse receptor targets. We fetched all ~62 000 GPCR-active ligands from CheMBL ver. and identified ~50 frequent substructures using Gaston that form parts of existing ligands for several diverse GPCR targets, e.g. small molecule, peptide and lipid receptors. The design of focused screening libraries started by querying these “privileged structures” against the entire Enamine screening compound collection in Instant JChem. More than 50 000 compounds were exported to Schrödinger Maestro, prepared with LigPrep and physicochemical properties were calculated with QikProp. Filters were applied to remove reactive structural groups and to enrich the dataset for compounds with desirable solubility, size and interaction features. Specifically the applied filters were: molecular weight (3–500), rotatable bonds (2–10), hydrogen bond donors and acceptors (0–5 and 0–10, respectively, but requiring at least one of either), Q log P (≤ 6), Q log S (≥ 6), FISA (7–330) and number of rings (max 5). The remaining compounds were clustered with the Maestro script for Canvas similarity and clustering and 1–3 representative compounds were handpicked from each cluster. The final library consisted of ~1600 lead- and drug-like screening compounds specifically selected for likely GPCR-activity.

To allow for library sub-selection based on target binding site charge, the overall library was divided into 3 sub-libraries containing negatively, neutral and positively charged compounds, respectively. All compounds were purchased from Enamine (Kiev, Ukraine) as 10 µL of 10 mM solution in 100% frozen DMSO plated on 384 well plates. The structure of compound 1 was confirmed by 1H NMR and HPLC-MS (Chart SI2†).

Pharmacological assay
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GraphPad Prism 6.0 (GraphPad Software, San Diego, C.A, USA).

**GPR132 homology modeling**

The crystal structure of the inactive human protease-activated 1 receptor (PAR1; PDB: 3W77) was selected as the main template, using the GPCRdb template selection tool. The resolution of the PAR1 structure is 2.0 Å. The protein sequence similarity between the target and template receptor is 64% in the seven trans-membrane helical (7 TM) region. Based on sequence length and similarity, a second template, the human chemokine receptor 4 (CXCR4) structure (PDB: 3ODU), was used for the part from the second extracellular loop (ECL2) spanning between TM5 and the cysteine forming the conserved disulfide bridge to TM3. To improve the GPR132 structure model for subsequent ligand docking, non-conserved amino acids within the binding pocket had their rotamers defined based on an in-house GPCR position-specific rotamer library that contains rotamers extracted from all published GPCR crystal structures. In this manner, the rotamers could be fine-tuned for S**131**, L**, Y**, W**2460**, T**128**, Y**61**, Y**635**, H**652**, V**654**, K**658**, L**7530**, L**7538**, and T**7542**. The protein sequences of the chimeric template and target were aligned with MEGA5.2, using the GPCRdb alignments as a reference for the seven helices.

Modeller9v13 (ref. 39) was applied to build 500 homology models and the best model was selected based on DOPE score. The model backbone was assayed by Ramachandran plots (Fig. S11†) within the PROCHECK webserver. The GPR132 model was prepared with the Schrödinger Protein Preparation Wizard, including an hydrogen optimization at pH 7 of the ionizable polar groups using Maestro PROPKA.

**Ligand docking and molecular visualization**

Compound 1 was docked with Glide in the extra precision mode, and flexible ligand sampling was applied. The partial charges of the ligands were assigned by Epik using the OPLS-2005 force field. The van der Waals radii scaling factor was set to 0.8 and the partial charge cutoff to 0.15. Further options were set to allow the rotation of hydroxyl hydrogen atoms in the binding site, enhance the planarity of conjugated π-systems, and to include the Epik state penalties to the scoring calculations. Subsequently, QM-Polarized Ligand Docking (QPLD) was applied. Jaguar performed the partial charge calculations with the QM estimation level set to accurate (6-31G*-LACVP* basis set, B3LYP density functional, and “Ultrafine” self-consistent field accuracy level). SiteMap was used to map the GPR132 binding cavity. Residue positions were indexed with the GPCRdb scheme, which corrects for helical bulges and constrictions. 2D structures were drawn in MarvinSketch and 3D structures were visualized using PyMOL.

**Molecular dynamics simulation**

The receptor–ligand complex was minimized using the energy minimization tool in MacroModel Schrödinger. The TNCG (Truncated Newton Conjugate Gradient) minimization method was used with maximum iteration steps set to 5000, and the convergence gradient to 0.001. Desmond was employed to conduct a molecular dynamics (MD) simulation for 20 ns on the complex in the predefined POPC membrane model. Explicit water molecules were handled using the simple point charge model. Constant temperature and pressure were applied at 300 K and 1.01325 bar, respectively. The system was coupled to an isotropic Berendsen thermo- and barostat, with relaxation set to 1 and 2 ps respectively.

**Results and discussion**

**Identification of the first surrogate GPR132 agonist**

Screening of the GPCR-focused libraries against the human GPR132 receptor yielded a hit (1, Chart 1) with 3.4 μM potency (Fig. 1A). Compound 1 is racemic and thus one of the enantiomers may have a higher potency than observed for the racemate. To evaluate whether the observed increase in luminescence is GPR132-mediated, 1 was also tested at another class A GPCR, the β2-adrenergic receptor. Ligand 1 did not induce β-arrestin recruitment in CHO cells expressing the β2-adrenergic receptor, while the positive control isoproterenol (ISO) produced a concentration-dependent response with an EC50 of approximately 100 nM (Fig. 1B and C). Hence, we concluded that the activity of 1 is not unspecific, but mediated via GPR132.

**GPR132 structure model and compound 1 binding mode**

To achieve a GPR132 model of sufficient quality for ligand docking, we found that it was necessary to locally replace the main template (the protease-activated 1 receptor) with alternative templates. The part of the second extracellular loop (ECL2) from the conserved cysteine (C**45x50**) to TM5 is crucial as it acts as a lid of the binding pocket in many class A GPCRs, and it often forms ligand interactions. This segment was longer in the protease-activated 1 receptor (10 instead of 7 residues); thus the human chemokine receptor 4 (also has 7 residues) was the most suitable template for modeling this segment. Furthermore, for binding pocket residues not conserved between the protease-activated 1 receptor and GPR132, we applied rotamers from the other most homologous receptors. In this way, 29 out of 36 (~81%) side chains within the GPR132 binding cavity could be defined according to a conserved crystal structure template. Validation of the GPR132 model in PROCHECK showed that 93.5% of the residues are within the optimal backbone torsion angles and 6.5% are near-optimal (Fig. S11†).

The docking pose of ligand 1 displayed close shape complementarity with the GPR132 binding pocket and good agreement with the receptor hydrophobic and hydrogen bonding sites (Fig. S12A–C). The R- and S-enantiomers exhibited very similar binding mode. However, the hydrophobic bonding site matched the methyl substituent better for the R-enantiomer, which was therefore chosen for the further analyses. Five residues displayed hydrogen bond interactions with the ligand namely V**131**, D**34x32**, H**637**, Y**63x31**, and K**6x38x**. Both Glide and QPLD resulted in identical outcome. To evaluate the
accuracy of our docking procedure, we re-docked the ligands in the three most homologous GPCR crystal structures; the protease-activated 1 receptor (PDB: 3VW7), the \( \mu \)-opioid receptor (PDB: 4DKL), and the chemokine receptor type 5 (PDB: 4MBS) respectively. The docked ligands showed very similar poses as the crystallized with RMSD values of 0.643, 1.306, and 0.270 Å, respectively (Fig. SI3A–C†).

In our molecular dynamics simulation of the 1 – GPR132 complex, in a membrane and solvent environment, the ligand remained in the same binding cavity throughout the entire 20

**Chart 1**  Chemical structures of the first disclosed GPR132 surrogate agonist 1 and its analogs 2–33. \( ^{Ag} \) denotes the six analogs 2, 9, 18, 31, 32, and 33 with weak agonistic activity. \( ^{Ant} \) denotes the two analogs 11 and 12 with weak antagonistic activity. \( ^{1} \) marks the three 1-substituted anthraquinone analogs 12, 16, and 29. 2D structures were drawn in MarvinSketch.49
ns simulation (Video SI1†), confirming that the complex is stable. The observed interactions and their durations are summarized in Fig. 2. The anthraquinone of 1 appeared to be occupying a hydrophobic site between TM3-5 that provided π–π aromatic (residues F333 and Y539) and hydrophobic (residues F335 and P439) contacts (Fig. 2C). The pyridinyl ring formed a π–π interaction with W266 and, together with the methylated linker; favorable hydrophobic contacts with several aliphatic residues could be seen (Fig. 2C).

The linker amide group displayed hydrogen bonding to both Y651 and a bridging water molecule to D4552, a residue position in the second extracellular loop that often forms contact with class A receptor ligands. In shorter periods of the simulation, the bond with Y651 was also indirectly formed via a water molecule (Fig. 2). All active analogs have this amide, indicating that its interactions could contribute to the stabilization of an active receptor conformation. The carbonyl oxygen in the amide linker also displayed a potential indirect interaction to Y540 (Fig. 2C). On the other hand, the carbonyl oxygen in the propionate group presented indirect hydrogen bonding to Y651 (Fig. 2C).

The oxygen atom in the 10-position of the anthraquinone ring displayed three putative water-mediated hydrogen bond interactions bridging K658, D532, and the backbone of L4554, respectively. The atom also presented a potential direct interaction with Y540. The simulation showed that the most frequent hydrogen bonds were the indirect ones to L4554 and D532 via bridging water molecules (Fig. 2C). The oxygen in the 9-position presented a direct hydrogen bond interaction with H457 or an indirect one with D4552 (Fig. 2C). The pyridinyl ring nitrogen interacted firmly with Y139 (Fig. 2). Altogether, six of the seven polar ligand atoms displayed hydrogen bond interactions, whereof four mainly indirectly via water molecules, and two directly to the receptor side chains.

Preliminary structure–activity relationships of compound 1 analogs

The structures of compound 1 and the 32 tested analogs are shown in Chart 1. All active analogs have lower potency than 1, and because of their limited solubility, it was not possible to produce full concentration–response curves and derive EC50/IC50 values. Instead, their relatively weak activities are reported in Table 1 at their maximum soluble concentration, which was 10 μM or 20 μM, respectively. Six of the analogs, namely 2, 9, 18, 31, 32 and 33, display weak agonistic activity.

Fig. 2 (A) Top and (B) side views of a representative binding pose of compound 1 (green), obtained from the molecular dynamics simulation. Hydrogen bonds are shown in dashed yellow lines representing the pairings between the ligand polar groups and the different receptor residues (grey). The simulation ligand interactions diagram (C) represents a summary of the interactions observed in the simulation obtained from Desmond. All detected hydrogen bonds (grey lines) are represented with their corresponding percentages. Residue positions are indexed with the GPCRDb scheme48 to correct for bulges and constrictions.
These two compounds have higher similarity to other non-

Table 1 Maximal effects* of the active analogs† of 1 at PathHunter

<table>
<thead>
<tr>
<th>Compound</th>
<th>Rel. $E_{\text{max}}$ (%)</th>
<th>SEM</th>
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<td>0.3</td>
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* The relative $E_{\text{max}}$ of tested compounds was calculated by dividing the span of compound response by the span of 1 response in the same experiment and on the same plate. † Compounds with a relative $E_{\text{max}}$ higher than 5% were considered as agonists.

with relative $E_{\text{max}}$ between 8 and 18% at the highest soluble concentration (10 or 20 µM, Table 1).

Compound 2 is the most similar analog to 1 and also has the highest activity of the analogs (relative $E_{\text{max}}$ of 18% at 10 µM). It differs by having a terminal pyrimidine on a trional group instead of a pyridine on a tetrahedral carbon, which is also the chiral center in compound 1. Glide docking of compound 2 showed that the hydrogen bond to Y1359 is lost as this compound lacks the pyridine nitrogen in the 4-position (Fig. SI4†). Compounds 3–5 have a terminal phenyl moiety and in turn lack the methyl group, indicating that either a hydrogen bond acceptor in the aryl 4-position or a methyl function could be beneficial for activity. Compounds 9 and 10 contain a terminal meta- and para-methoxy-phenyl, respectively, on a linker that is extended by an extra nitrogen or oxygen atom, respectively, compared to 1. In our docking, the nitrogen in 9 formed a hydrogen bond to the backbone carbonyl of C35300 (forms a disulfide bridge from the second extracellular loop to the top of TM3), whereas the oxygen in 10 displayed no interaction (Fig. SI4B and C†). This provides a hypothesis for the activity of 9, as opposed to 10. In the final analogs with agonistic activity, 31–33, the 1,1-diphenyl group docked into the same site as the pyridine moiety, not the anthraquinone core, of compound 1 (Fig. SI4D–F†). In each compound, the propionate linker showed a hydrogen bond to Y6951. Only compound 32 was still able to display hydrogen bonding to K9258 via the acetamide terminal group (Fig. SI4D–F†).

Two of the analogs, 11 and 12, display antagonistic activity: the inhibition of compound 1 is 39–46% at 20 µM (Table 2). These two compounds have higher similarity to other non-

Table 2 Inhibition* of 1-mediated β-arrestin recruitment in CHO

<table>
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<tr>
<th>Compound</th>
<th>Inhibition (%)</th>
<th>SEM</th>
<th>Conc. (µM)</th>
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<td>16</td>
<td>20</td>
<td>2</td>
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<tr>
<td>12</td>
<td>39</td>
<td>14</td>
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* The responses have been normalized to ligand buffer (100% inhibition) and 8.5 µM (EC50) of 1 (0% inhibition). † Compounds with more than 25% inhibition were considered as antagonists.

antagonist analogs than to each other. Compound 11 is one of five analogs, 7–11, which have a linker that is one atom longer than in compound 1. Moreover, 11 is unique in that it has the largest terminal substituent, an indole, and lacks a preceding polar functionality. 12 is one of three 1-substituted anthraquinones (12, 16 and 29). Compared to 16 the ring of 12 is non-aromatic and positioned on a one atom longer linker. Furthermore, varying linker lengths are seen among both agonistic and antagonistic analogs; compounds 9, 11, 12, and 18 have linkers that are between one atom longer and two atoms shorter.

Taken together, the testing of commercially available analogs has allowed for the derivation of preliminary structure–activity relationships, that although limited, can serve as a guide for further lead optimization by custom synthesis.

Conclusions

The association of GPR132 with several alternative proposed physiological ligands and functions warrants the development of tool compounds that can be used to characterize this receptor. In this work, we have disclosed the first surrogate agonist 1. The analogs tested, and the binding modes and structure–activity relationships proposed herein, will be valuable for future medicinal chemistry optimization of compound 1, as well as mutagenesis of interacting receptor residues.

Acknowledgements

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