Bitopic fluorescent antagonists of the A₂A adenosine receptor based on pyrazolo[4,3-e][1,2,4]triazolo[1,5-c]pyrimidin-5-amine functionalized congeners‡

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A pyrazolo[4,3-e][1,2,4]triazolo[1,5-c]pyrimidin-5-amine antagonist of the A₂A adenosine receptor (AR) was functionalized as amine congeners, fluorescent conjugates and a sulfonate, and the A₂A-AR binding modes were predicted computationally. The optimal n-butyl spacer was incorporated into the following A₂A-AR-selective (Kᵢ, nM) conjugates: BODIPY630/650 derivative 11 (MRS7396, 24.6) and AlexaFluor488 derivative 12 (MRS7416, 30.3). Flow cytometry of 12 in hA₂A-AR-expressing HEK-293 cells displayed saturable binding (low nonspecific) and inhibition by known A₂A-AR antagonists. Water-soluble sulfonate 13 was a highly potent (Kᵢ = 6.2 nM) and selective A₂A-AR antagonist based on binding and functional assays. Docking and molecular dynamics simulations predicted the regions of interaction of the distal portions of these chain-extended ligands with the A₂A-AR. The BODIPY630/650 fluorophore of 11 was buried in a hydrophobic interhelical (TM1/TM7) region, while AlexaFluor488 of 12 associated with the hydrophilic extracellular loops. In conclusion, we have identified novel high affinity antagonist probes for A₂A-AR drug discovery and characterization.

Introduction

The development of selective agonists and antagonists of the four subtypes of adenosine receptors (ARs) has been extensively explored. Antagonists of the Gₛ protein-coupled A₂A-AR are sought for as agents for treating neurodegenerative conditions such as Parkinson’s disease (PD) and Alzheimer’s disease (AD), and for coadministration with cancer immunotherapy. Caffeine, the most consumed psychostimulant in the world, acts as a nonselective AR antagonist and readily enters the brain to antagonize the A₂A-AR at doses generally consumed. Epidemiological evidence showing a lower occurrence of AD and PD with modest caffeine intake points to the possibility that caffeine consumption is neuroprotective. Indeed, A₂A-AR antagonists can exert a neuroprotective effect on excitotoxicity in animal models; one mechanism appears to enhance the activity of an A₁AR, which forms a heterodimer with the A₂A-AR, to inhibit glutamate release. A₂A-AR antagonists also control microglia-mediated neuroinflammation. Therefore, it is possible that A₂A-AR antagonists may also delay neurodegenerative disease progression. In the striatum, A₂A-AR antagonists act to boost dopaminergic signaling, and thus provide symptomatic relief to reduce the motor deficits in PD without inducing dyskinesia.

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Certain selective A2A AR antagonists, e.g. 1–4 (Chart 1), have been shown to enter the brain at sufficient levels for imaging and efficacy in PD models.10–12 A caffeine-like 1,3,7-trialkylxanthine, istradefylline 1, is approved in Japan for treating PD, and its safety and efficacy in reducing off-time (when PD symptoms return) in levodopa-treated patients were established in a 52 week trial.10 A tricyclic pyrazolo-triazolopyrimidine derivative 4, which binds potently and selectively to the A2AAR, was in clinical trials for PD.12

In the periphery, elevated adenosine levels present in the microenvironment of tumors lead to a suppressed immune response, to shift the T cell response away from an aggressive state capable of attacking tumors.6 Increasing evidence supports the use of AR antagonists in cancer treatment, either with selectivity for the A2AAR alone or with dual selectivity for the A2AAR and A2BAR. Cytokine production in CD8+ chimeric antigen receptor (CAR) T cells was increased, and both CD8+ and CD4+ CAR T cells were activated upon blocking the A2AAR.7 Thus, there is great interest in the discovery of novel A2AAR antagonists to act either centrally or peripherally, and new tools for ligand discovery are needed.

Various fluorescent probes have been developed over the past few years for characterizing ARs,13–15 and their use in fluorescence polarization (FP), fluorescence resonance energy transfer (FRET) and flow cytometry (FCM) has been proven to be feasible. Moreover, these ligands can provide a better understanding of receptor location, function and regulation. Most of the reports on fluorescent A2AAR antagonists used 5-amino-7-(3-[4-methoxy]phenylpropyl)-2-[2-furyl]pyrazolo[4,3-c]-1,2,4-triazolo[1,5-c]pyrimidine (SCH442416, 3) as the pharmacophore due to the p-methoxyphenylpropyl side chain moiety of the antagonist, which served as the site for attachment of functionalized chains through an ether linkage.

Much of the ligand development for ARs now benefits from a detailed structural knowledge of the human (h) A2AAR.16–20 More than two dozen high-resolution X-ray crystallographic A2AAR structures with bound agonists or antagonists have been determined and used for the in silico screening of chemical libraries. A2AAR complexes with antagonist 4-(2-[7-amino-2-[2-furyl]-1,2,4]triazolo[2,3-a][1,3,5]triazin-5-ylamino)ethyl)phenol (ZM241385, 2) show that the heterocyclic pharmacophore binds to residues in the orthosteric binding pocket that also coordinates with the adenine moiety of AR agonists.16,17 Although no X-ray structure of the 3-A2AAR complex has been determined, we used the 2-A2AAR structure as the structural template for molecular modeling. The hypothetical docking pose of 3 and its AlexaFluor488 derivative 5

**Chart 1** Structures of selective A2A AR antagonists and the target series: (A) widely used pharmacological probes and clinical candidates (1–4); (B) reported fluorescent probe 5 derived from 3 and functionalized amine congeners of varied chain length 6 explored in this study.
established most of the conserved interactions in the orthosteric binding site.\textsuperscript{13,14} The modeling also predicted stabilizing interactions of the tethered fluorophore with specific charged and H-bonding residues of the second extracellular loop (EL2).

Conjugate 5 was not optimal for fluorescent binding due to its moderate hA2AAR affinity ($K_i = 111 \text{ nM}$).\textsuperscript{13} Also, its green light emission presents difficulties in fluorescence microscopy due to cell autofluorescence. Thus, there remains a need for A2AAR antagonist fluorescent probes of higher affinity and compatibility with microscopy. A BODIPY650/655 conjugate containing a secondary amine in the linking chain was reported to have a $K_i$ value of 15 nM for the A2AAR, but its utility was not established.\textsuperscript{13} With this aim, we explored further the SAR of the distal region of this chemical series by varying the chain length of the spacer group and the terminal fluorophore in order to enhance the affinity, selectivity and photophysical properties.

**Results and discussion**

We initially prepared a series of primary amine congeners of the A2AAR antagonist 3 by extending the $p$-methoxyphenylpropyl chain, which is predicted to lie outside the orthosteric binding site of the pharmacophore and be able to reach accessory sites.\textsuperscript{13} The congener 6a ($n = 2$) was reported earlier,\textsuperscript{13} and we hypothesized that extension of the alkyl spacer could enhance the affinity or selectivity for this receptor, perhaps by establishing H-bond or electrostatic interactions with the polar residues in the EL region of the A2AAR. The synthesis of the pyrazolo[4,3-$c$][1,2,4]triazolo[1,5-$c$]pyrimidin-5-amine derivatives is shown in Scheme 1.

The synthesis of fluorescent ligands 9–12 was accomplished in 4 steps from commercially available 3. The latter was demethylated to 7 by the action of BBr\textsubscript{3} and then treated with an excess of methyl-2-bromoacetate (12 eq.) and Cs\textsubscript{2}CO\textsubscript{3} as a base to provide ester 8 in 90% yield. Other methods, such as using NaH as the base with 1 eq. of methyl-2-bromoacetate in DMF, were attempted but led to the formation of di and tri-alkylated compounds. Amide synthesis was then performed by treatment of ester 8 with a series of alkyl diamines to give the amine congeners 6a–e. Two of these amines, 6a ($n = 2$) and 6c ($n = 4$), were reacted with activated fluorophore moieties in DMF in the presence of Et\textsubscript{3}N to

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**Scheme 1** Synthesis of A2AAR antagonist functionalized congeners of 6 and their fluorescent derivatives 9–12. (a) BBr\textsubscript{3}, CH\textsubscript{2}Cl\textsubscript{2}, rt, 4 h; (b) methyl-2-bromoacetate, Cs\textsubscript{2}CO\textsubscript{3}, MeOH, 40 °C, overnight; (c) diaminooalkane, MeOH (9:1 v/v), rt, overnight; (d) activated fluorophore (AlexaFluor647 N-hydroxysuccinimidyl ester for 9 and 10, BODIPY 630/650 N-hydroxysuccinimidyl ester for 11, AlexaFluor488 carboxylic acid, 2,3,5,6-tetrafluorophenyl ester for 12), Et\textsubscript{3}N, DMF, rt, overnight.
afford, after semi-preparative HPLC, compounds 9–12 in good yield (40–60%): cyanine 5 red fluorescent AlexaFluor647 \((n = 2, 9; n = 4, 10)\), red fluorescent BODIPY630/650 \((n = 4, 11)\) and green fluorescent AlexaFluor488 \((n = 4, 12)\). Both fluorophores are commonly incorporated into ligand tools for chemical biology.21

The affinity of the amine congeners and other antagonists was measured using standard radioligand binding assays for hA1, A2A and A3ARs (Table 1).14,15,22 Membranes of human embryonic kidney (HEK)293 cells expressing the AR of interest were used in the assay. Primary amine congeners 6a–6d containing spacers of 2–5 methylenes were similar in terms of hA3AR affinity \((K_i = 6–9 \text{nM})\), with the butylamino congener 6c displaying the highest A3AR selectivity in the series. The affinity of this compound for the hA3AR was determined to be 6.0 \muM, with only slightly higher hA1AR affinity. Homogeneity in 6 benzylmethylenes in compound 6c lowered A3AR affinity. Among the fluorescent conjugates of butylamino congener 6c, i.e. 10, 11, and 12, the highest A3AR affinity and selectivity were observed with the BODIPY630/650 fluorophore 11 and AlexaFluor488 12 (Fig. 1). However, fluorescent 11 and 12 exhibited only a 4-fold weaker A3AR affinity with respect to the parent amino derivative 6c. Interestingly, AlexaFluor488 conjugate 12 was at least as potent in terms of A3AR affinity as its shorter homologue 5. No dependence of the affinity on the chain length was evident when comparing AlexaFluor647 conjugates 9 and 10. Affinity for the mouse (m)ARs was also measured for the selected compounds by methods described,22 and 11 was particularly potent and selective for the mA3AR \((K_i = 2.1 \text{nM})\), in contrast to 12 \((K_i = 585 \text{nM})\).

We performed molecular modeling analysis to identify possible binding modes of fluorescent conjugates 11 and 12 using the high resolution 2-A2AR X-ray structure (PDB ID: 4EIY).17 We first docked reference compound 3 at the hA2AR by retaining several water molecules observed in the X-ray structure as described (ESI†). In the corresponding docking pose (Fig. 2A), the pyrazolotriazolopyrimidine core of 3 docked in the orthosteric binding site similar to the triazolopyrimidine core of 2, with a \(\pi–\pi\) stacking interaction of the aromatic core with F168 (EL2), and a H-bonding network with N253 (6.55, using standard GPCR notation23) and E169 (EL2). The methoxy-phenyl substituent of 2 pointed toward the extracellular (EC) side of the receptor and interacted with the side chain of Glu169 (EL2) through a water molecule. The binding mode of 3 described above was validated

**Table 1.** AR binding affinity determined for a series of pyrazolo[4,3-e][1,2,4]triazolo[1,5-c]pyrimidin-5-amine derivatives \((R^1\)), as in Scheme 1. Human ARs, unless otherwise noted (m indicates mouse).

| Compd | Structure | Affinity, \(K_i\), nM (or % inhibit) |
|-------|-----------|----------------------------------|
| 3b    | \(R = \text{OCH}_3\) | \((35 \pm 5\%)\) |
| 3f    | \(R = \text{AlexaFluor488,} \ n = 2\) | \(4.1\) |
| 6a    | \(n = 2\) | \(1270 \pm 140\) |
| 6b    | \(n = 3\) | \(1300 \pm 350\) |
| 6c    | \(n = 4\) | \(2390 \pm 100\) |
| 6d    | \(n = 5\) | \(1910 \pm 120\) |
| 6e    | \(n = 6\) | \(6500 \pm 2830\) |
| 7b    | \(R = \text{OH}\) | \((66 \pm 2\%)\) |
| 9     | \(R = \text{AlexaFluor647,} \ n = 2\) | \((13 \pm 6\%)\) |
| 10    | \(R = \text{AlexaFluor647,} \ n = 4\) | \((20 \pm 3\%)\) |
| 12    | \(R = \text{AlexaFluor488,} \ n = 4\) | \((40 \pm 3\%)\) |
| 13    | \(R = \text{AlexaFluor488,} \ n = 4\) | \((40 \pm 3\%)\) |

\(\sim\) Competition radioligand binding assays were conducted with membranes prepared from HEK-293 cells expressing recombinant A1, A2A, or A3ARs (human) unless otherwise noted. Their incubation was performed for 1 h at 25 °C. The radioligands used were: A1AR, \(^3\text{H}\)8-cyclopentyl-1,3-dipropylxanthine \((\text{H})\text{DPCPX,} 0.5 \text{nM}) 14; A2AAR, \(^3\text{H}\)ZM241385 2 (1.0 nM) or from published data13 (as noted) \(^3\text{H}\)2-[2-carboxyaminoethyl]-2-carboxyaminoethyl]-N\(^\sim\)N\(^\sim\)-dimethyluronamidin \((\text{H})\text{CGS21680,} 10 \text{nM}) 15; A3AR, \(^[^3\text{H}]\)N\(^\sim\)N\(^\sim\)-[4-amino-3-iodobenzyl]adenosine-5'N-methyluronamide \((\text{H})\text{DEA-MAECA,} 0.2 \text{nM}) 16. Nonspecific binding was determined using 10 \muM 8-B[4-[4-[[4-[[(2-aminoethyl)amino]carbonyl][methyl]oxy]phenyl]-1,3-dipropylxanthine] (XAC) A1AR and A3AR or 10 \muM adenosine-5'-N-ethyluronamide (NECA) A1AR. HEP-233 cells expressing recombinant mA1, mA2A, or mA3ARs were used. Values are expressed as the mean ± SEM from 3 independent experiments. The cell lines were from the American Type Culture Collection (ATCC, Manassas, VA), and the cDNA for the ARs was obtained from cdna.org. Data from Kumar et al. and Keskés et al.13,14: Percent inhibition at 10 \muM. Fluorophore moiety, as shown in Scheme 1 \((R^1)\). Using \(^[^3\text{H}]\)I6 as the radioligand.11, MRS7396; 12, MRS7416; 13, MRS7352.
using 30 ns molecular dynamics (MD) simulation. During the simulation (Video S1, replica analysis reported in Table S1†), the methoxy-phenyl moiety folded toward transmembrane domain 2 (TM2) and established a π–π stacking interaction with Y271 (7.36), while the aromatic core maintained the H-bond network observed in the initial docking pose. Notably, the most energetically favored ligand–protein complex featured the same interaction pattern described above (Fig. S1†).

The potent fluorescent conjugate 11 was then docked in the A2AAR structure (Fig. 2B) with the same water molecules retained as in the 3–A2AAR complex.17 The pyrazolotriazolopyrimidine core established the same interactions observed for 3 in the orthosteric binding site. The fluorophore linker pointed toward the EC side and folded back toward the TM bundles by directing the fluorophore group to an aromatic pocket at the interface between TM1 and TM7 (transparent surface in Fig. 2B). To explore other orientations of the linker at the receptor’s EC side, we docked compound 6c, the amino precursor of 11, at the A2AAR. To sample all the possible H-bond acceptor/donor partners on the EC side of the receptor, we removed water molecules interacting with E169 during the docking. We then clustered the binding modes obtained according to the linker orientation. This selection resulted in two alternative binding modes (Fig. S2†) that were subsequently subjected to MD validation (30 ns of simulation run in triplicate for each binding mode, see Table S1†). In the most energetically favored docking mode (hereby referred to as “BM1”), orange carbon sticks in Fig. S2† docking score = −12.077 kcal mol$^{-1}$), the tail was oriented toward TM4 and TM5 with the amide moiety establishing a H-bond with the side chain of E169, while the terminal amine group H-bonded with the backbone of E169 (EL2) and the side chain of K150 (EL2). This latter H-bond was expected to be unstable in a dynamic environment due to competition in the formation of a salt-bridge between K150 and D170 (EL2). In the alternative binding mode (BM2, green carbon sticks in Fig. S2† docking score = −10.994 kcal mol$^{-1}$), the tail was oriented toward TM1 and TM2 and did not establish additional interactions. MD trajectory analysis revealed that BM1 achieved ligand–protein complexes that were more energetically favored (data not shown). Fig. S3† depicts the two 11–A2AAR complexes with the most favorable ligand–protein interaction energy (IE, values differed by less than 2 kcal mol$^{-1}$ and were considered equivalent) obtained for BM1 that features different orientations of the tail. We therefore searched for hydrophobic and aromatic regions in the proximity of the terminal amine moiety to investigate the compatibility of these orientations with the fluorophore insertion. Specifically, we searched for receptor regions rich in hydrophobic and aromatic residues at 5, 13, and 14 Å from the nitrogen atom of the terminal alkylamino group of 6c (Fig. S3B†). The choice of distances reflected the analysis of the N-group distance to aromatic fluorophores in both the docked and the energy minimized three-dimensional structures of 11 (Fig. S3A†). Of the two possible orientations of 11, only one displayed hydrophobic and aromatic residues at distances compatible with the insertion of the aromatic fluorophore. Notably, the aromatic/hydrophobic region is located at the interface between TM1 and TM7 (Fig. S3C†), thus suggesting the same orientation of the ligand as observed from the docking analysis. Nonetheless, we cannot exclude that 11 might explore different regions on the receptor’s EC side.

The fluorescent conjugate 12 (considered as the species carrying a −2 net charge) was docked in the A2AAR structure by following the same procedure described for compound 11 (ESI†). The docking output suggested two equally plausible orientations of the fluorophore (Fig. 2C and D). In one docking pose (BM1, cyan carbon sticks in Fig. 2C, docking score = −11.490 kcal mol$^{-1}$), the fluorophore group was projected toward EL3. In the alternative binding mode (BM2, purple carbon sticks, docking score = −11.279 kcal mol$^{-1}$), the fluorophore group interacted with the residues in EL2 and EL3. The interaction pattern of the core was the same as those for the other members of this chemical series: H-bonds with N253 and E169 (EL2); π–π stacking with H252 (6.52), F168 (EL2), and Y271 (7.36). In the MD simulation starting from BM1 (30 ns run in triplicate, see Table S1†) the fluorophore group and the linker fluctuated on the EC surface of the receptor without engaging in specific interactions except for a H-bond involving the fluorophore carboxylate moiety (data not shown). On the other hand, the simulations starting from BM2 (30 ns, replica analysis in Table S1†) achieved more energetically favored ligand–protein complexes. The simulations returned the ligand–protein complex characterized by the lowest IE value for all trajectories and converged in a unique binding mode (Fig. S4†) featuring the fluorophore group stacked between EL2 and EL3. In such a conformation, the ligand established a tight network of H-bonds and salt-bridges between charged residues in EL2 and EL3 and the polar/charged counterparts in the fluorophore moiety. In particular, during the MD simulation (visualization of run 3 trajectory selected as an example, Video
S2f) the sulfonic groups established salt bridges with K153 (persistent) and K150 (intermittent) in EL2, one of the amine moieties interacted with the sidechain of E169 (EL2), and the carboxylate moiety of the ligand replaced E169 (EL2) in the salt bridge with H264 (EL3). Moreover, for most of the total simulation time, the pyrazolotriazolopyrimidine core maintained its interaction pattern with N253 (6.55) and F168 (EL2), while the linker between the core and the fluorophore group was anchored to TM7, EL2 and TM2 through H-bond interactions with the backbone of S67 (2.65) and the sidechains of Q157 (EL2) and Y271 (7.36), respectively (Video S2f).

Thus, both 11 and 12 are bitopic in the sense that each bridged two separate domains of the A2AR, i.e. the orthosteric binding site that is well defined in X-ray structures16,17 and an additional domain. The BODIPY630/650 fluorophore of 11 is predicted to be buried in a hydrophobic region, while AlexaFluor488 of 12 associates with the hydrophilic ELs. The bitopic nature of these conjugates does not necessarily imply allosteric modulation of the orthosteric action,28 which is unexplored in this series.

In order to further explore the ligand interactions with the EC surface, we prepared an aryl sulfonate 13, which contained a terminal group capable of multiple polar interactions (Scheme S1, ESI†). This terminal phenylsulfonic acid moiety also would allow for π–π interactions with aromatic residues. A further benefit of incorporating a sulfonate group is that it carries a permanent negative charge at physiological pH and would prevent penetration into the blood brain barrier,22 a useful characteristic for a pharmacological probe for
in vivo studies.\textsuperscript{24} 13 was potent and selective for binding to the h and mA\textsubscript{2}ARs, with 671- and 426-fold selectivity compared to the hA\textsubscript{1}AR and hA\textsubscript{3}AR, respectively. 13 binding to the mA\textsubscript{2}AR and mA\textsubscript{3}AR was insignificant.

The binding of 13 was modeled using the same docking procedure and MD validation described for 6c. Fig. 3 depicts the three alternative binding modes of 13. In the most energetically favorable pose (BM1, blue carbon sticks) the sulfophenyl ring established a π–cation interaction with K153 (EL2) and the sulfonic group engaged in an H-bond interaction with the residue backbone. In the other two docking poses (BM2 and BM3, magenta and orange carbon sticks, respectively) the sulfophenyl ring established a π–π stacking interaction with H264 (EL3) and the sulfonic group engaged in an H-bond interaction with the residue backbone. In the other two docking poses (BM2 and BM3, magenta and orange carbon sticks, respectively) the sulfophenyl ring established a π–π stacking interaction with K153 (EL2) and the sulfonic group interacted with the side chain of either K153 (EL2; BM2, magenta) or S156 (EL2; BM3, orange). During the MD simulation, the ligand’s sulfophenyl tail fluctuated considerably (high averaged root mean square deviation values, Table S1\textsuperscript{†}), thus demonstrating the instability of the interaction pattern predicted in the initial docking poses. On the other hand, the three different initial poses converged in a unique binding mode, as the ligand–protein complexes with the lowest IE value returned by each different binding mode featured the same conformation (Fig. S5). In particular, the 7-phenylpropyl ring moved toward TM7 to establish a π–π stacking interaction with Y271 (7.36).

Two representative derivatives were shown to be A\textsubscript{2}AR antagonists in a cyclic AMP assay (Fig. S6). The activation curve for the known agonist 15 right-shifted in a parallel manner at fixed concentrations of 11 and 13. The EC\textsubscript{50} of 15 shifted from 0.89 ± 0.17 nM to 128 ± 35 nM in the presence of 11 (1000 nM) and to 10.2 ± 2.3 nM in the presence of 13 (100 nM). Off-target binding activities at 45 diverse receptors were determined by PDSP25 for selected compounds: 6c and 13 (ESI†). The only K\textsubscript{i} values below 10 μM were: 6c, 3.36 μM, 5HT\textsubscript{2}A serotonin receptor; 1.92 μM, 5HT\textsubscript{2}B receptor. Thus, the primary amine precursor of the fluorescent conjugates 10–12 was not promiscuous in its interaction with other proteins, i.e. this chemical series is not associated with pan-assay interference compounds.\textsuperscript{26}

Compounds 11 and 12 were tested as fluorescent tracers for flow cytometry of HEK-293 cells expressing the hA\textsubscript{2}AR (Fig. 4A). 11 proved to have high nonspecific binding, with a low level of specific binding amounting to <25% of the total and therefore was not optimal for flow cytometric analysis. Thus, 11 tended to bind non-specifically to hydrophobic membranes by undetermined mechanisms. However, the more hydrophilic 12 displayed low nonspecific binding by this method, and its binding was saturable with a K\textsubscript{d} value of 45.4 nM (Fig. 4B). For inhibition studies, the cells were co-incubated for one hour prior to flow cytometry with fluorescent probe 12 (10 nM) and a test antagonist. The binding of 12 was inhibited by two known A\textsubscript{2}AR antagonists, nonxanthine 3 and xanthine 17, with the expected range of potency.\textsuperscript{27} The K\textsubscript{i} values were 3.8 and 17.2 nM (Fig. 5), respectively, which corresponded closely to the reported values of 4.1 and 18 nM at hA\textsubscript{2}AR.\textsuperscript{1,13} Agonist inhibition of the competitive binding of 12 was complex (ESI†).

**Conclusions**

We succeeded in identifying useful A\textsubscript{2}AR probes by systematically varying the chain length of amine-functionalized congeners of a potent antagonist and coupling the primary amine having an optimal length to various fluorophores. We successfully enhanced their A\textsubscript{2}AR affinity compared to known fluorescent A\textsubscript{2}AR ligands.\textsuperscript{15} Conjugates 11 and 12 were potent and selective antagonist probes for the A\textsubscript{2}AR, but 12 was more promising for characterization of the hA\textsubscript{2}AR in whole cells by flow cytometry. Molecular modeling suggested that the fluorophore of 11 interacted with

![Fig. 3](image-url)
hydrophobic regions between TMs. However, the hydrophilic fluorophore of 12 was coordinated to the ELs, consistent with its lower overall hydrophobicity and more favorable whole cell binding characteristics compared to 11 under the present conditions. Evaluation of 11 and 12 by confocal microscopy studies remains to be performed. Thus, we have introduced antagonist ligands displaying high A2AAR affinity and selectivity that may serve as versatile tools to better study this receptor.

Conflict of interest
The authors declare no competing interests.

Abbreviations
AN Acetonitrile
AR Adenosine receptor
BODIPY 4,4-Difluoro-4-bora-3a,4a-diaza-s-indacene
CNS Central nervous system
DMF N,N-Dimethylformamide
EDC N-(3-Dimethylaminopropyl)-N'-ethylcarbodiimide
HEK Human embryonic kidney
IE Interaction energy
NECA 5’-N-Ethylcarboxamidoadenosine
PD Parkinson’s disease
TBAP Tetrabutylammonium dihydrogen phosphate
THF Tetrahydrofuran
TLC Thin layer chromatography
TM Transmembrane domain

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