

## RESEARCH ARTICLE

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# Bitopic fluorescent antagonists of the A<sub>2A</sub> 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 A<sub>2A</sub> adenosine receptor (AR) was functionalized as amine congeners, fluorescent conjugates and a sulfonate, and the A<sub>2A</sub>AR binding modes were predicted computationally. The optimal *n*-butyl spacer was incorporated into the following A<sub>2A</sub>AR-selective (*K<sub>i</sub>*, nM) conjugates: BODIPY630/650 derivative **11** (MRS7396, 24.6) and AlexaFluor488 derivative **12** (MRS7416, 30.3). Flow cytometry of **12** in hA<sub>2A</sub>AR-expressing HEK-293 cells displayed saturable binding (low nonspecific) and inhibition by known A<sub>2A</sub>AR antagonists. Water-soluble sulfonate **13** was a highly potent (*K<sub>i</sub>* = 6.2 nM) and selective A<sub>2A</sub>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 A<sub>2A</sub>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 A<sub>2A</sub>AR drug discovery and characterization.

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† Electronic supplementary information (ESI) available: Chemical synthesis, characterization data, HPLC analysis, pharmacological studies, off-target screening and additional molecular modeling procedures/results. 3D coordinates of the hA<sub>2A</sub>AR in complex with **11** (docking pose). 3D coordinates of the hA<sub>2A</sub>AR in complex with **12** (docking pose, BM2). Video of 30 ns of MD simulations of the hA<sub>2A</sub>AR in complex with **3**. Video S1. Trajectory visualization (left panel) and ligand-protein interaction energy profile (right panel) of 30 ns of membrane MD simulation of the 3-hA<sub>2A</sub>AR complex. Video of 30 ns of MD simulations of hA<sub>2A</sub>AR in complex with **12**. Video S2. Trajectory visualization of 30 ns of membrane MD simulation of the 12-hA<sub>2A</sub>AR complex starting from the docking binding mode denoted "BM2": side view facing TM5, TM6, and TM7 (left panel), and side view facing TM4, TM5 and TM6 (right panel). See DOI: 10.1039/c7md00247e

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## Introduction

The development of selective agonists and antagonists of the four subtypes of adenosine receptors (ARs) has been extensively explored.<sup>1–3</sup> Antagonists of the G<sub>s</sub> protein-coupled A<sub>2A</sub>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.<sup>4–7</sup> Caffeine, the most consumed psychostimulant in the world, acts as a nonselective AR antagonist and readily enters the brain to antagonize the A<sub>2A</sub>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, A<sub>2A</sub>AR antagonists can exert a neuroprotective effect on excitotoxicity in animal models; one mechanism appears to enhance the activity of an A<sub>1</sub>AR, which forms a heterodimer with the A<sub>2A</sub>AR, to inhibit glutamate release.<sup>8</sup> A<sub>2A</sub>AR antagonists also control microglia-mediated neuroinflammation.<sup>9</sup> Therefore, it is possible that A<sub>2A</sub>AR antagonists may also delay neurodegenerative disease progression. In the striatum, A<sub>2A</sub>AR antagonists act to boost dopaminergic signaling, and thus provide symptomatic relief to reduce the motor deficits in PD without inducing dyskinesia.

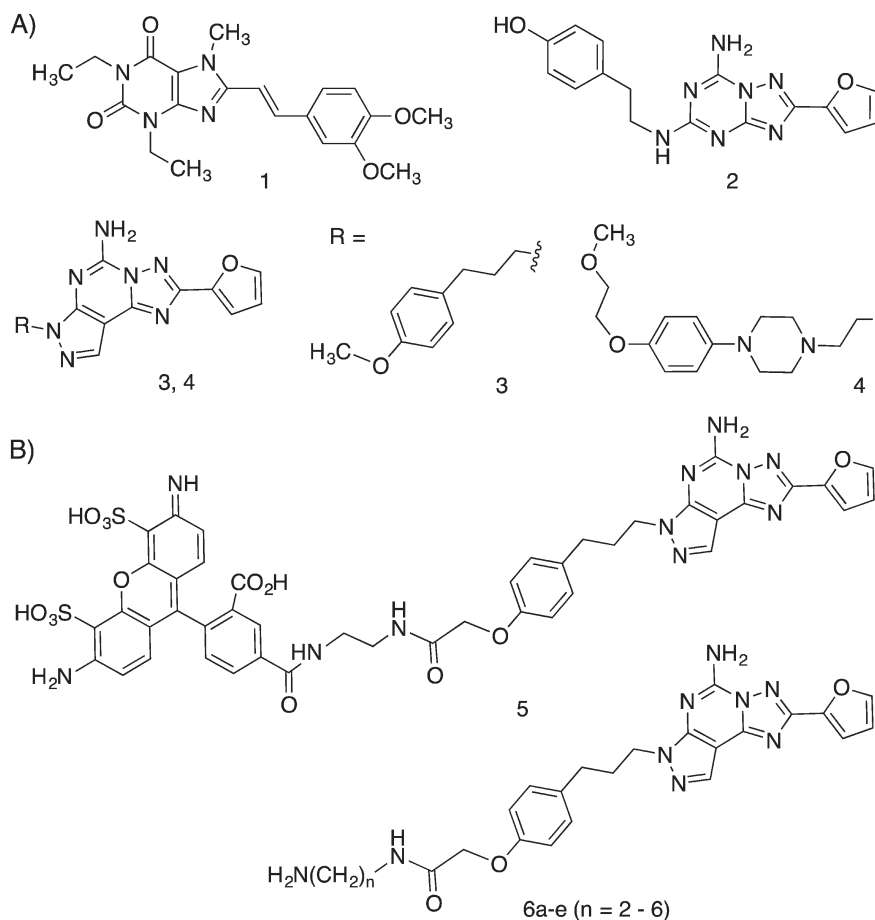
Certain selective A<sub>2A</sub>AR antagonists, *e.g.* 1–4 (Chart 1), have been shown to enter the brain at sufficient levels for imaging and efficacy in PD models.<sup>10–12</sup> A caffeine-like 1,3,7-trialkylxanthine, istradefylline 1, is approved in Japan for treating PD, and its safety and efficacy in reducing in off-time (when PD symptoms return) in levodopa-treated patients were established in a 52 week trial.<sup>10</sup> A tricyclic pyrazolo-triazolopyrimidine derivative 4, which binds potently and selectively to the A<sub>2A</sub>AR, was in clinical trials for PD.<sup>12</sup>

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.<sup>6</sup> Increasing evidence supports the use of AR antagonists in cancer treatment, either with selectivity for the A<sub>2A</sub>AR alone or with dual selectivity for the A<sub>2A</sub>AR and A<sub>2B</sub>AR. Cytokine production in CD8<sup>+</sup> chimeric antigen receptor (CAR) T cells was increased, and both CD8<sup>+</sup> and CD4<sup>+</sup> CAR T cells were activated upon blocking the A<sub>2A</sub>AR.<sup>7</sup> Thus, there is great interest in the discovery of novel A<sub>2A</sub>AR 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,<sup>13–15</sup> 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 A<sub>2A</sub>AR antagonists used 5-amino-7-(3-(4-methoxyphenyl)propyl)-2-(2-furyl)pyrazolo[4,3-*e*]-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) A<sub>2A</sub>AR.<sup>16–20</sup> More than two dozen high-resolution X-ray crystallographic A<sub>2A</sub>AR structures with bound agonists or antagonists have been determined and used for the *in silico* screening of chemical libraries. A<sub>2A</sub>AR 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.<sup>16,17</sup> Although no X-ray structure of the 3-A<sub>2A</sub>AR complex has been determined, we used the 2-A<sub>2A</sub>AR structure as the structural template for molecular modeling. The hypothetical docking pose of 3 and its AlexaFluor488 derivative 5



**Chart 1** Structures of selective A<sub>2A</sub>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.<sup>13,14</sup> The modeling also predicted stabilizing interactions of the tethered fluorophore with specific charged and H-bonding residues of the second extracellular loop (EL2).

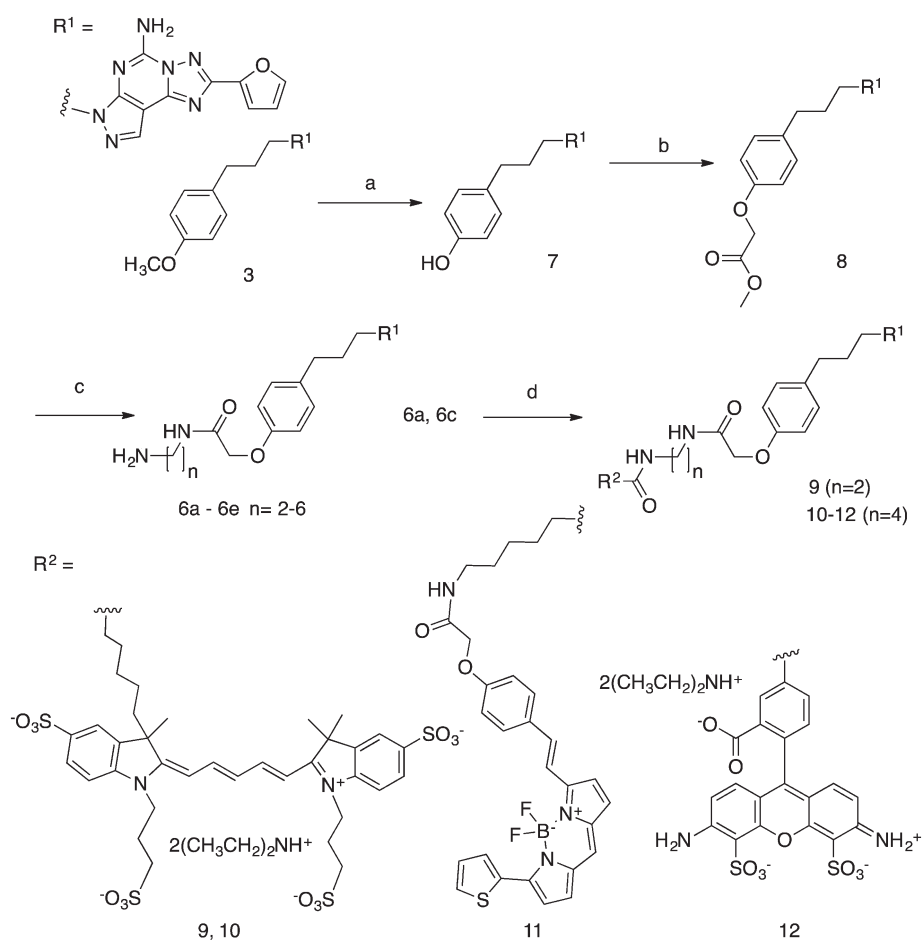
Conjugate 5 was not optimal for fluorescent binding due to its moderate hA<sub>2A</sub>AR affinity ( $K_i = 111$  nM).<sup>13</sup> Also, its green light emission presents difficulties in fluorescence microscopy due to cell autofluorescence. Thus, there remains a need for A<sub>2A</sub>AR antagonist 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 A<sub>2A</sub>AR, but its utility was not established.<sup>13</sup> 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 A<sub>2A</sub>AR 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.<sup>13</sup> The congener 6a ( $n = 2$ ) was reported earlier,<sup>13</sup> 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 A<sub>2A</sub>AR. The synthesis of the pyrazolo[4,3-*e*][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<sub>3</sub> and then treated with an excess of methyl-2-bromoacetate (12 eq.) and Cs<sub>2</sub>CO<sub>3</sub> 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<sub>3</sub>N to



**Scheme 1** Synthesis of A<sub>2A</sub>AR antagonist functionalized congeners of 6 and their fluorescent derivatives 9–12. (a) BBr<sub>3</sub>, CH<sub>2</sub>Cl<sub>2</sub>, rt, 4 h; (b) methyl 2-bromoacetate, Cs<sub>2</sub>CO<sub>3</sub>, MeOH, 40 °C, overnight; (c) diaminoalkane, 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<sub>3</sub>N, DMF, rt, overnight.



We performed molecular modeling analysis to identify possible binding modes of fluorescent conjugates **11** and **12** using the high resolution 2-A<sub>2A</sub>AR X-ray structure (PDB ID: 4E1Y).<sup>17</sup> We first docked reference compound **3** at the hA<sub>2A</sub>AR 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 notation<sup>23</sup>) 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

Chemical structures of compounds 3, 7, 13, 6, and 5, 9, 12 are shown. Structure 3, 7, 13 is a triazole-fused pyrazole derivative with a 4-R-benzyl group and a furan-2-yl group. Structure 6 is a triazole-fused pyrazole derivative with a 4-(3-R<sub>1</sub>-propoxy)benzyl group and a furan-2-yl group, linked via a carbamate to a polyamine chain. Structure 5, 9, 12 is a triazole-fused pyrazole derivative with a 4-(3-R<sub>2</sub>-propoxy)benzyl group and a furan-2-yl group, linked via a carbamate to a polyamine chain.

<sup>a</sup> Competition radioligand binding assays were conducted with membranes prepared from HEK-293 cells expressing recombinant A<sub>1</sub>, A<sub>2A</sub>, or A<sub>3</sub>ARs (human) unless otherwise noted. Their incubation was performed for 1 h at 25 °C. The radioligands used were: A<sub>1</sub>AR, [<sup>3</sup>H]8-cyclopentyl-1,3-dipropylxanthine ([<sup>3</sup>H]DPCPX, 0.5 nM) 14; A<sub>2A</sub>AR, [<sup>3</sup>H]ZM241385 2 (1.0 nM) or from published data<sup>13</sup> (as noted) [<sup>3</sup>H]2-[p-(2-carboxyethyl)phenyl-ethylamino]-5'-N-ethylcarboxamidoadenosine [<sup>3</sup>H]CGS21680, 10 nM) 15; A<sub>3</sub>AR, [<sup>125</sup>I]N<sup>6</sup>-(4-amino-3-iodobenzyl)adenosine-5'-N-methyluronamide ([<sup>125</sup>I]-AB-MECA, 0.2 nM) 16. Nonspecific binding was determined using 10 μM 8-[4-[[[2-aminoethyl]amino]carbonyl]methyl]-oxy]phenyl]-1,3-dipropylxanthine (XAC) 17 (A<sub>1</sub>AR and A<sub>2A</sub>AR) or 10 μM adenosine-5'-N-ethyluronamide (NECA) 18 (A<sub>3</sub>AR). HEK-293 cells expressing recombinant mA<sub>1</sub>, mA<sub>2A</sub>, or mA<sub>3</sub>ARs 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.<sup>b</sup> Data from Kumar *et al.* and Kecskés *et al.*<sup>13,14</sup> <sup>c</sup> Percent inhibition at 10 μM. <sup>d</sup> Fluorophore moiety, as shown in Scheme 1 (R<sup>2</sup>). <sup>e</sup> Using [<sup>3</sup>H]16 as the radioligand.<sup>f</sup> 11, MRS7396; 12, MRS7416; 13, MRS7352.



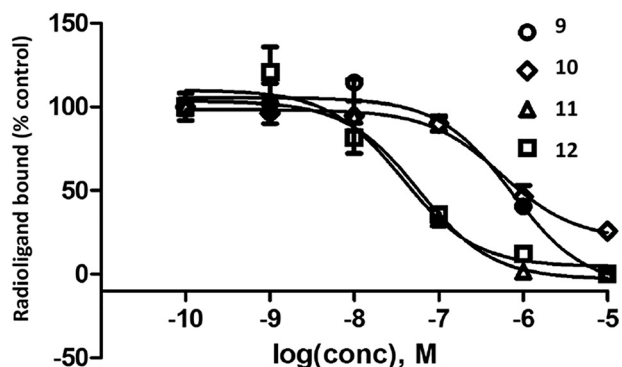


Fig. 1 Antagonist radioligand ( $[^3\text{H}]2$ , 1.0 nM) binding inhibition curves for the  $\text{hA}_{2\text{A}}\text{AR}$  for four antagonist fluorescent conjugates. Compound numbers: MRS7322 **9**, MRS7395 **10**, MRS7396 **11** and MRS7416 **12**. Membranes from HEK-293 cells expressing the  $\text{hA}_{2\text{A}}\text{AR}$  were used, and their incubation was performed for 1 h at 25 °C. Results are expressed as the mean  $\pm$  SEM. The  $K_i$  values from three independent experiments are listed in Table 1.

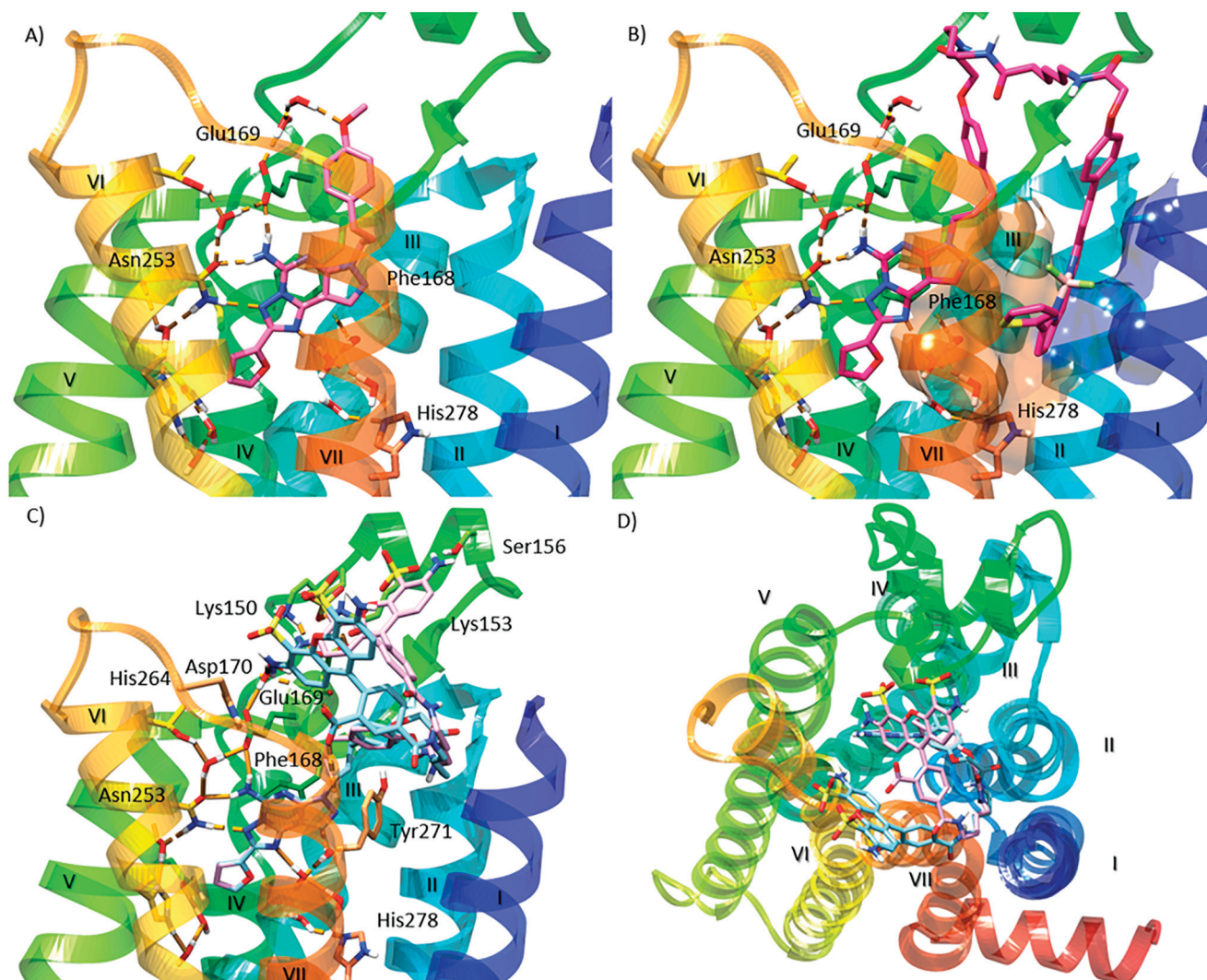
using 30 ns molecular dynamics (MD) simulation. During the simulation (Video S1, replica analysis reported in Table S1†), the methoxy-phenyl moiety folded toward transmembrane domain 2 (TM2) and established a  $\pi$ - $\pi$  stacking interaction with Y271 (7.36), while the aromatic core maintained the 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  $\text{A}_{2\text{A}}\text{AR}$  structure (Fig. 2B) with the same water molecules retained as in the  $3\text{-A}_{2\text{A}}\text{AR}$  complex.<sup>17</sup> 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  $\text{hA}_{2\text{A}}\text{AR}$ . 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 =  $-12.077 \text{ kcal mol}^{-1}$ ), 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 =  $-10.994 \text{ kcal mol}^{-1}$ ), 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**- $\text{A}_{2\text{A}}\text{AR}$  complexes with the most favorable ligand-protein interaction energy (IE, values differed by less than  $2 \text{ kcal mol}^{-1}$  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 rings 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  $\text{A}_{2\text{A}}\text{AR}$  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 =  $-11.490 \text{ kcal mol}^{-1}$ ), the fluorophore group was projected toward EL3. In the alternative binding mode (BM2, purple carbon sticks, docking score =  $-11.279 \text{ kcal mol}^{-1}$ ), 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);  $\pi$ - $\pi$  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





**Fig. 2** Molecular modeling of antagonist binding to the hA<sub>2A</sub>AR. Details of the binding site of the X-ray structure of the receptor modeled with various ligands docked: (A) known antagonist **3**, with the retention of a subset of water molecules found in the high resolution A<sub>2A</sub>AR structure;<sup>17</sup> (B) the BODIPY630/650-labeled antagonist **11**, showing the most energetically favorable orientation of the terminal fluorophore chain; side (C) and top (D) views of the two possible orientations of the fluorophore group of derivative **12**. Residues establishing polar (dashed orange lines) and  $\pi$ - $\pi$  interactions with the docked ligands are represented as sticks. Aromatic residues establishing hydrophobic contacts with the terminal fluorophore of compound **11** are represented as transparent surfaces with colors matching the corresponding TM domain. Non-polar hydrogen atoms are omitted.

S2†) 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 S2†).

Thus, both **11** and **12** are bitopic in the sense that each bridged two separate domains of the A<sub>2A</sub>AR, *i.e.* the orthosteric binding site that is well defined in X-ray structures<sup>16,17</sup>

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,<sup>28</sup> 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  $\pi$ - $\pi$  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,<sup>22</sup> a useful characteristic for a pharmacological probe for





*in vivo* studies.<sup>24</sup> 13 was potent and selective for binding to the h and mA<sub>2A</sub>ARs, with 671- and 426-fold selectivity compared to the hA<sub>1</sub>AR and hA<sub>3</sub>AR, respectively. 13 binding to the mA<sub>1</sub>AR and mA<sub>3</sub>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  $\pi$ - $\pi$  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  $\pi$ -cation 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†), 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  $\pi$ - $\pi$  stacking interaction with Y271 (7.36).

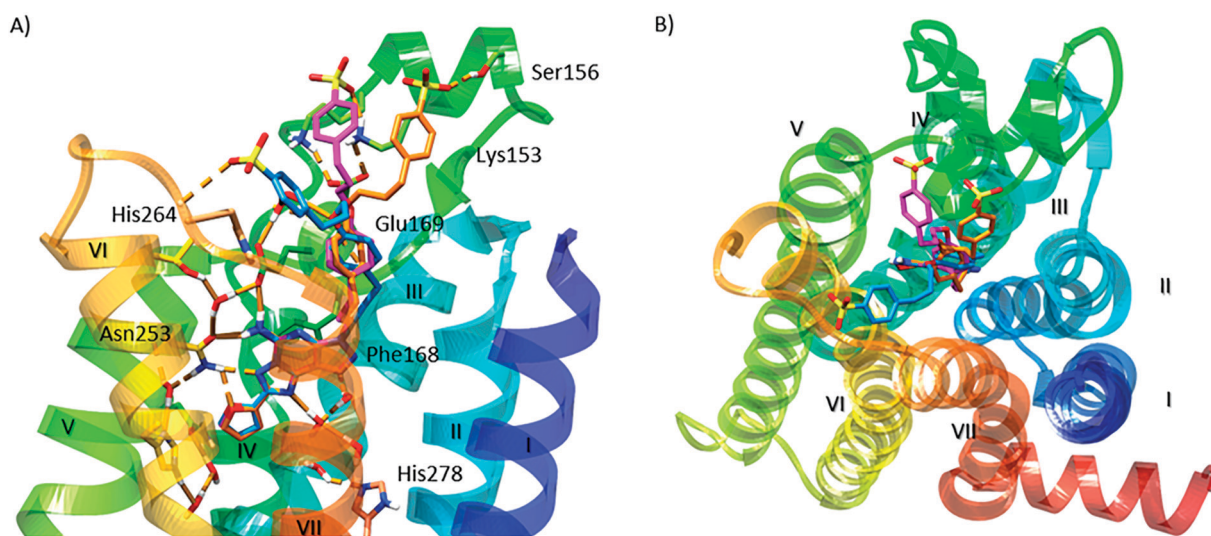
Two representative derivatives were shown to be A<sub>2A</sub>AR antagonists in a cyclic AMP assay (Fig. S6†). The activation curve for the known agonist 15 right-shifted in a parallel manner at fixed concentrations of 11 and 13. The EC<sub>50</sub> of 15 shifted from  $0.89 \pm 0.17$  nM to  $128 \pm 35$  nM in the presence of 11 (1000 nM) and to  $10.2 \pm 2.3$  nM in the presence of 13 (100 nM). Off-target binding activities at 45 diverse receptors were determined by PDSP<sup>25</sup> for selected compounds: 6c and 13 (ESI†). The only K<sub>i</sub> values below 10  $\mu$ M were: 6c, 3.36  $\mu$ M, 5HT<sub>2A</sub> serotonin receptor; 1.92  $\mu$ M, 5HT<sub>2B</sub> 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.<sup>26</sup>

Compounds 11 and 12 were tested as fluorescent tracers for flow cytometry of HEK-293 cells expressing the hA<sub>2A</sub>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<sub>d</sub> 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<sub>2A</sub>AR antagonists, nonxanthine 3 and xanthine 17, with the expected range of potency.<sup>27</sup> The K<sub>i</sub> 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<sub>2A</sub>AR.<sup>1,13</sup> Agonist inhibition of the competitive binding of 12 was complex (ESI†).

## Conclusions

We succeeded in identifying useful A<sub>2A</sub>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<sub>2A</sub>AR affinity compared to known fluorescent A<sub>2A</sub>AR ligands.<sup>15</sup> Conjugates 11 and 12 were potent and selective antagonist probes for the A<sub>2A</sub>AR, but 12 was more promising for characterization of the hA<sub>2A</sub>AR in whole cells by flow cytometry. Molecular modeling suggested that the fluorophore of 11 interacted with



**Fig. 3** Modeling of antagonist 13 binding to the hA<sub>2A</sub>AR: side (A) and top (B) views of the three possible orientations (BM1, blue; BM2, magenta; BM3, orange) of the *p*-sulfophenyl tail. Residues establishing polar (dashed orange lines) and  $\pi$ - $\pi$  interactions with the docked ligands are represented as sticks. Non-polar hydrogen atoms are omitted.



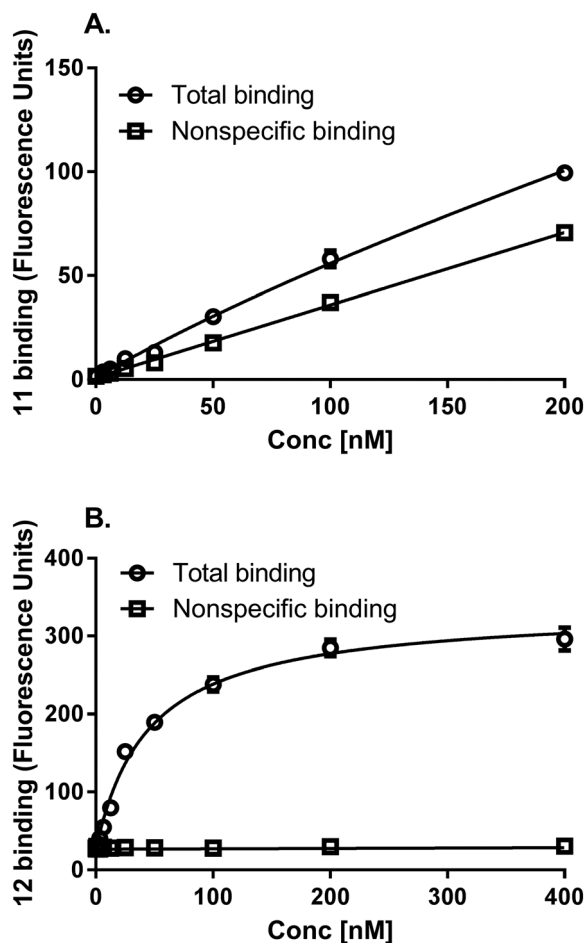


Fig. 4 Saturation of fluorescent antagonists 11 (A) and 12 (B) in hA<sub>2A</sub>AR-expressing HEK-293 cells measured using flow cytometry following a 1 h incubation at 37 °C. Non-specific binding was determined with 10  $\mu$ M 3. Results are expressed as the mean  $\pm$  SEM. The  $K_d$  value calculated from Fig. 4B is listed in the text.

hydrophobic regions between TMs. However, the hydrophilic fluorophore of 12 was coordinated to the ELs, consis-

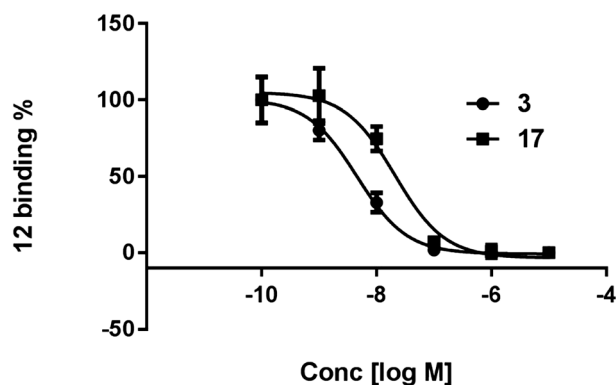


Fig. 5 Inhibition of the specific binding of fluorescent antagonist 12 (10 nM, 1 h incubation at 37 °C) in hA<sub>2A</sub>AR-expressing HEK-293 cells measured using flow cytometry. Competing antagonists were: nonxanthine 3 and xanthine 17. Non-specific binding was determined with 10  $\mu$ M 3. Results are expressed as the mean  $\pm$  SEM. The  $K_i$  values are listed in the text.

tent 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 A<sub>2A</sub>AR 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	<i>N,N</i> -Dimethylformamide
EDC	<i>N</i> -(3-Dimethylaminopropyl)- <i>N'</i> -ethylcarbodiimide
HEK	Human embryonic kidney
IE	Interaction energy
NECA	5'- <i>N</i> -Ethylcarboxamidoadenosine
PD	Parkinson's disease
TBAP	Tetrabutylammonium dihydrogen phosphate
THF	Tetrahydrofuran
TLC	Thin layer chromatography
TM	Transmembrane domain

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