Sub-millisecond conformational dynamics of the A_{2A} adenosine receptor revealed by single-molecule FRET

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Abstract

The complex pharmacology of G-protein-coupled receptors (GPCRs) is defined by their multi-state conformational dynamics. Single-molecule Förster Resonance Energy Transfer (smFRET) is well-suited to quantify dynamics for individual protein molecules, however, its application to GPCRs is challenging; therefore, smFRET has been limited to studies of inter-receptor interactions in cellular membranes and receptors in detergent environments. Here, we performed smFRET experiments on functionally active human A_{2A} adenosine receptor (A_{2A}AR) molecules embedded in freely diffusing lipid nanodiscs to study their intramolecular conformational dynamics. We propose a dynamic model of A_{2A}AR activation that involves a slow (>2 ms) exchange between the active-like and inactive-like conformations in both apo and antagonist-bound A_{2A}AR, explaining the receptor's constitutive activity. For the agonist-bound A_{2A}AR, we detected faster (390±80 μs) ligand efficacy-dependent dynamics. This work establishes a general smFRET platform for GPCR investigations that can potentially be used for drug screening and/or mechanism-of-action studies.

Introduction

G-protein-coupled receptors (GPCRs) constitute the largest superfamily of membrane proteins in humans containing over 800 members, which mediate critical physiological processes, such as neurotransmission, homeostasis, inflammation, reproduction, olfaction, vision, taste, and others^{1,2}. GPCRs recognize a large variety of endogenous extracellular signaling molecules transmitting their corresponding signals inside the cell, and this process can be modulated by synthetic ligands or drug molecules. In fact, over 30% of all FDA-

approved drugs target GPCRs³. Multiple lines of evidence suggest that the molecular mechanism of GPCR activation extends beyond a simple ‘on/off’ mode. First, apo receptors show basal activity that can be suppressed by inverse agonists⁴. Second, different agonists vary in efficacy and can stimulate receptor activity to a different extent⁵. Third, a single receptor can signal through several intracellular pathways, some of which could be preferentially activated by so-called “biased” ligands⁶. These three phenomena indicate that receptors are highly dynamic molecules and sample several active and inactive states stochastically (for review, see ⁷⁻⁹).

The A_{2A} adenosine receptor (A_{2A}AR) is expressed in many organs and tissues including those in the immune system, basal ganglia, heart, lungs, and blood vessels¹⁰. Throughout the body, A_{2A}AR regulates the cardiovascular tonus causing vasodilation and promotes healing of inflammation-induced injuries by suppressing immune cells^{11,12}. In the brain, A_{2A}AR modulates dopamine and glutamate neurotransmission¹². A_{2A}AR is a promising target for drugs against insomnia, chronic pain, depression, Parkinson’s disease, and cancer^{12,13}. On the molecular level, A_{2A}AR is activated by the endogenous extracellular agonist adenosine and initiates the cAMP-dependent signaling pathway via G_s and G_{olf} proteins^{12,14}. Besides G-proteins, A_{2A}AR interacts with numerous other partners including GRK-2 kinase, β-arrestin, and other GPCRs^{14,15}. One cryoEM and over 50 high-resolution X-ray crystallographic structures are available for antagonist- or agonist-bound A_{2A}AR and for its ternary complex with an agonist and an engineered G protein, making this receptor an excellent model system for investigating GPCR structural dynamics. While static structures provide critical information about the receptor’s lowest energy states, our understanding of the A_{2A}AR function remains critically incomplete without the detailed knowledge of its conformational dynamics.

The current information about A_{2A}AR conformational dynamics is based mostly on several reported NMR experiments^{16–22}. In response to ligand binding, different A_{2A}AR amino acids either alter their sole stable conformations or vary relative probabilities of coexisting stable conformations^{16,17}. On the picosecond-to-nanosecond timescale, some A_{2A}AR amino acids increase side-chain dynamics, while others become stabilized¹⁸. Sub-millisecond conformational variability was shown for both apo-form¹⁹ and agonist-bound A_{2A}AR^{16,17,20}. Large-scale conformational changes in A_{2A}AR with dwell times of seconds were also reported^{19,21}, but two independent studies described the corresponding long-lived states differently: in one report¹⁹, a 3-state model with an attributed basal activity of 70% was proposed, while in the other²¹, the authors put forward a 4-state model with a negligible basal activity. Thus, the current picture of A_{2A}AR dynamics is complex and contradictory.

Studies of A_{2A}AR dynamics face two major challenges: first, the need to cover a wide range of timescales from nanoseconds to seconds, and next, the difficulty to untangle multiple protein states within the ensemble. Single-molecule fluorescence spectroscopy provides tools to address both of these difficulties. Depending on the applied method, the fluorescence signal from individual receptors can be tracked with as low as a nanosecond temporal resolution for a total duration of either milliseconds in case of freely-diffusing molecules or even seconds to minutes using immobilized molecules^{7,23}.

Single-molecule fluorescence spectroscopy methods have been previously applied to studying GPCR conformational dynamics⁷. For example, environmentally-sensitive fluorescent dyes have been used as single-molecule reporters of conformational changes in the β_2 adrenergic receptor (β_2 AR)^{24–27} and visual rhodopsin^{28,29}. Single-label experiments are attractive because of a minimal influence of the dye on the native receptor dynamics, but the experimental readouts are often limited and lack detailed structural interpretation. Additionally, the results of single-label experiments can be obscured by multi-state dye

photophysics. Another approach, based on single-molecule Förster Resonance Energy Transfer (smFRET) between two dyes can provide more direct structural outcomes and introduce additional internal controls, however, at the expense of double-labeling. smFRET has been shown to be especially useful to investigate structural dynamics of GPCR dimers^{30–34}. To our knowledge, at the moment of this writing, ref. ³⁵ is the only published application of smFRET to quantifying intramolecular conformational dynamics in GPCRs; this study addressed structural changes on the intracellular side of immobilized β_2 AR in detergent micelles.

Here we applied smFRET to investigate the conformational dynamics of A_{2A} AR in lipid nanodiscs freely diffusing in solution without immobilization. Using the MFD-PIE (multiparameter fluorescence detection with pulsed interleaved excitation) technique³⁶ (Fig. 1A), we tracked the relative movements of two dyes attached to the intracellular tip of the transmembrane helix TM6 (L225C^{6,27}, superscripts indicate Ballesteros–Weinstein numbering³⁷) and to the C-terminal intracellular helix H8 (Q310C^{8,65}) of A_{2A} AR (Fig. 1B). We observed that FRET efficiency in the double-labeled A_{2A} AR increases upon agonist-binding (Fig. 1C). Several burst-wise fluorescence analysis approaches — plot of burst-wise FRET efficiency against donor fluorescence lifetime³⁸, FRET 2-Channel kernel-based Density Estimator (FRET-2CDE)³⁹, Burst Variance Analysis (BVA)⁴⁰, and filtered Fluorescence Correlation Spectroscopy (fFCS)⁴¹ — subsequently revealed sub-millisecond conformational dynamics of A_{2A} AR. Based on quantitative analysis of the obtained data for the receptor in its apo state and upon addition of the inverse agonist ZM241385, the partial agonist LUF5834, or the full agonist NECA to the receptor, we finally propose a dynamic model of A_{2A} AR activation.

Results

Labeling and reconstitution of A_{2A}AR in nanodiscs

To track the conformational dynamics of A_{2A}AR with smFRET we chose to attach two fluorescent dyes to mutated residues L225C^{6,27} on the intracellular end of TM6 and Q310C^{8,65} on the C-terminal end of H8 (Fig. 1B, Fig. S1A). In previous A_{2A}AR FRET studies, a fluorescent protein-based FRET donor and fluorescent molecule based acceptor in similar labeling positions were shown to provide sufficient contrast between the active and inactive receptor states in live cells^{42,43}. The residue position L225^{6,27} is also homologous to the native cysteine C265^{6,27} in β₂AR that has been frequently used for fluorescent labeling^{24–26,44–48}.

We expressed the double-Cys mutant (L225C^{6,27}/Q310C^{8,65}) of A_{2A}AR in *Leishmania tarentolae* and simultaneously labeled it with two maleimide-functionalized dyes, Alexa488 and Atto643. The wild-type (WT) A_{2A}AR has six unpaired cysteines in its transmembrane helices (Fig. S1A). To achieve specific labeling of the two genetically introduced cysteines, but spare the transmembrane native cysteines, we labeled the receptors in isolated cell membranes, as described previously⁴⁹. After labeling, the receptors were purified and reconstituted in MSP1D1 nanodiscs, which can accommodate only a single monomeric receptor per nanodisc⁵⁰.

Size-exclusion chromatography confirmed a high purity and monodispersity of the nanodisc-reconstituted A_{2A}AR samples (Fig. S1B). Labeling efficiencies of 26% (Alexa488) and 8% (Atto643) were obtained for the mutant A_{2A}AR (Table S1). Labeling specificity was confirmed with the WT receptor, which showed only a marginal dye fluorescence associated with the protein after the labeling procedure (Fig. S1B). In both ensemble spectra and lifetime measurements of the fluorescently labeled A_{2A}AR FRET-sensitized acceptor

emission was readily observed, proving the existence of double-labeled FRET-active molecules in the samples (Fig. S1C-D).

To test whether the double-cysteine mutant A_{2A}AR (L225C^{6.27}/Q310C^{8.65}) is functional, we measured the ligand-induced thermostabilization of the isolated receptors as well as the agonist-induced cAMP accumulation in living cells. A fluorescent thermal stability assay⁵¹ showed that the addition of either the antagonist ZM241385 or the agonist NECA in saturating concentrations increased the melting temperature of both WT and mutant A_{2A}AR with respect to the apo-state by >7 °C, indicating ligand-binding activity of the receptor (Fig. S1E). A BRET assay of cAMP accumulation in HEK293T cells transiently expressing A_{2A}AR showed very similar pEC₅₀ values for both WT (6.41±0.15) and double-mutant (6.45±0.06) forms of the receptor upon stimulation with the agonist NECA (Fig. S1F). The mutant form of A_{2A}AR retained functional activity in nanodiscs and in cells, therefore we assume that the conformational dynamics observed for the double-labeled receptor in smFRET experiments represent the native dynamics of the WT receptor.

smFRET reveals ligand-induced conformational changes in A_{2A}AR

We diluted fluorescently labeled A_{2A}AR to single-molecule concentrations, mounted the sample on a microscope cover slip and recorded fluorescence intensity, lifetime, and anisotropy data from individual molecules diffusing freely across the femtoliter-sized observation spot (approximated by a 3D Gaussian with half-widths 0.5 μm, 0.5 μm and 2 μm) of a confocal fluorescence microscope (Fig. 1A). Inside the spot, donor and acceptor fluorophores are excited alternatingly using a two-color pulsed-interleaved excitation (PIE)⁵². The residence time of individual molecules (~1-10 ms) in the laser spot sets the upper limit of timescales approachable for the observation of A_{2A}AR conformational dynamics. Using a 4-detector MFD scheme (Fig. S2), photons detected from individual molecules were digitally

tagged with (1) the spectral band in which they were detected, (2) their global arrival time with microsecond accuracy, (3) their relative arrival time with respect to the laser pulses within a ps-ns range, and (4) their optical polarisation⁵³. PIE together with two-color detection allowed us to distinguish double-labeled receptors (simultaneously labeled with donor and acceptor) from “donor-only” and “acceptor-only” receptors (Fig. S3, see ‘Selection of double-labeled, donor-only and acceptor-only subpopulations’ in SI).

The fraction of A_{2A}ARs simultaneously labeled with donor and acceptor fluorophores showed different distributions of FRET efficiency depending on the bound ligand (Fig. 1C). The antagonist ZM241385 did not change FRET efficiency distribution within experimental error. On the contrary, both the partial agonist LUF5834 and the full agonist NECA shifted the mean FRET efficiency to larger values and increased the overall distribution width, compared to the apo-receptor. The increase in FRET efficiency was less pronounced for the partial agonist LUF5834 than for the full agonist NECA.

Fluorescence lifetime data suggest sub-millisecond conformational dynamics of A_{2A}AR

Besides fluorescence intensity, FRET is also reflected in fluorescence lifetime data. A two-dimensional plot of the per-burst FRET efficiency against the donor fluorescence lifetime provided the first insights into the receptor’s conformational dynamics (Fig. 2A,B). In theory, data for rigid molecules, in which FRET efficiency remains constant over the duration of a burst should be distributed along a curved diagonal line that intersects the lifetime axis at the lifetime of the donor-only population and the FRET efficiency axis at unity, commonly referred to as the ‘static FRET line’ (Fig. 2A). Alternatively, if receptor molecules sample different conformations during their residence time in the focal spot (1-10 ms) on a timescale that is longer than the nanosecond fluorescence lifetime, their bursts should be shifted from the ‘static FRET line’ towards the longer lifetime region. This phenomenon can be explained

by the higher weights of the lower FRET states in the fluorescence lifetime averaging due to the larger number of photons emitted by the donor. The observed rightward deviations of our burst data from the static FRET line indicate the existence of sub-millisecond conformational dynamics (beyond the fast dynamics expected for dye linkers) in the apo as well as agonist- and antagonist-bound states of A_{2A}AR (Fig. 2B).

FRET-2CDE and BVA confirm that agonists enhance conformational dynamics in A_{2A}AR compared to apo receptor

Variations of FRET efficiency within fluorescence bursts from individual receptors suggest the presence of conformational dynamics. To analyze these variations further we used two complementary approaches: FRET-2CDE³⁹ and BVA⁴⁰. Both methods assign dynamics scores to individual molecules and are sensitive to the dynamics that are slower than the time used for FRET efficiency averaging (roughly, 100 μs for both approaches).

The FRET-2CDE score provides an unbiased way for the separation of static and dynamic subpopulations of molecules and for the comparison of their fractions in different datasets³⁹. The main advantage of FRET-2CDE is that it is minimally influenced by the mean FRET efficiency in a dynamic molecule. Theoretically, static molecules should have FRET-2CDE≈10, while higher FRET-2CDE values should correspond to more pronounced conformational dynamics (Fig. 2C). In our data, neither the apo nor ligand-bound A_{2A}AR showed a clear separation of different receptor subpopulations along the FRET-2CDE axis, but the agonists did increase the mean FRET-2CDE score of A_{2A}AR compared to the apo or antagonist-bound receptors (Fig. 2D, Table S2). The fraction of receptors that exceeded FRET-2CDE>20 threshold was also higher for the agonist-bound receptors: 14±1 % for ZM241385, 15±1 % for apo, 24±2 % for LUF5834, 23±2 % for NECA. These results indicate that either the amplitude of the observed dynamics or the number of the inter-state transitions per burst increase in A_{2A}AR upon agonist binding.

BVA provides a statistically robust way to test whether the observed variations of FRET efficiency exceed fluctuations expected from the shot noise and thus to prove conformational dynamics⁴⁰. To apply BVA, we split bursts into consecutive photon-windows with $n=5$ photons in each (roughly 100 μs long), calculated standard deviations of the bin-wise FRET efficiencies within each burst, and plotted them against the mean FRET efficiency. The obtained BVA scores exceeded the 99.9% confidence interval expected from the shot noise under all four conditions (Fig. 2E,F), therefore, BVA confirmed that sub-millisecond conformational dynamics is already present in the apo and antagonist-bound $A_{2A}AR$ and further increased in the agonist-bound $A_{2A}AR$.

ffCS reveals two modes of $A_{2A}AR$ dynamics with different timescales

To estimate the timescales of $A_{2A}AR$ conformational dynamics we used the ffCS approach⁴¹. We used the photon arrival time and anisotropy information to split the photon stream from the double-labeled molecules *in silico* between the low-FRET (LF) and high-FRET (HF) channels (Fig. S4). Theoretically, if the LF and HF species are just two extremes of a heterogeneous ensemble of long-lived receptor states, then cross-correlation between the two channels will show only a diffusion-based sigmoidal component decreasing with the lag time (Fig. 2G). Contrarily, if the LF and HF species interconvert on the μs –ms timescales, the cross-correlation should decrease in the time lag region shorter than the state exchange time. (Fig. 2G) The cross-correlation function between the LF and HF channels (Fig. 2H, Table S3) showed two dynamics-conditioned anticorrelation terms on top of the positive diffusion-conditioned term:

$$G(\tau) = G_{diff}(\tau)(1 - A_1 e^{-\tau/\tau_1} - A_2 e^{-\tau/\tau_2}).$$

The ffCS analysis revealed fast microsecond-time ($\tau_1 = 3\text{--}20 \mu s$) dynamics in the apo-receptor data and in the data recorded in the presence of each of the three ligands. We believe

that the fast dynamics of the dye linkers or local fluctuations of the protein can account for this term. While a single dynamic term (A_1) described the experimental data well for the apo and the antagonist-bound $A_{2A}AR$, the agonist-bound $A_{2A}AR$ showed an additional pronounced slower dynamics term with an exchange time of $\tau_2 = 390 \pm 80 \mu s$. Since the slow term appears upon agonist addition, we attribute it to the increased dynamics of the agonist-bound protein.

PDA quantifies populations of active and inactive states in dynamic $A_{2A}AR$

Finally, to quantify the populations of $A_{2A}AR$ in different FRET-states in the ligand-free and ligand-bound forms we used the photon distribution analysis (PDA) method^{54,55}. PDA takes into account receptor dynamics and Poissonian shot noise in the measured data, and, therefore, it is preferred to a simple multi-state Gaussian fitting. For PDA, we split the fluorescence bursts into time bins of constant duration (0.5 ms, 1 ms, and 2 ms) and analyzed them globally across all apo and ligand-bound conditions. In dynamic systems, a molecule can sample several states during an individual time bin, and therefore, the FRET efficiency distribution depends on the duration of the time bin. PDA is most sensitive for picking up interconversion times on the diffusion time scale (1-10 ms); for faster or slower dynamics, PDA can be constrained *a priori* to demonstrate that the proposed model of the conformational space does not contradict the observed FRET efficiency distributions.

A model with at least three states with different FRET efficiencies was required to fit the experimental distributions. Since PDA is insensitive to fast ($< 20 \mu s$) dynamics observed in fFCS, the fitting models for the apo and antagonist-bound $A_{2A}AR$ did not include any interconversions between states. On the other hand, interconversion between two of the states with a fixed exchange time ($\tau_2 = 390 \pm 80 \mu s$) as observed in fFCS was introduced in the agonist-bound PDA models. For simplicity, one state was considered long-lived and only two

states were considered interconvertible. Finally, using PDA we determined the mean values and variances of inter-dye distances for each state and the populations of states under the apo and ligand-bound conditions (Fig. 3A, Fig. S5, and Table S4).

The PDA results (Fig. 3A) revealed that both agonists increased the population of the highest FRET efficiency state (HF state), and decreased the population of the state with intermediate FRET efficiency (MF) state – therefore, we assume that the HF and MF states correspond to the active-like and inactive-like conformations of $A_{2A}AR$, respectively. Approximately 10-20% of $A_{2A}AR$ molecules always stay in the low-FRET (LF) state independently of the added ligand – we speculate that these receptors are locked in a long-lived non-functional state or improperly folded. PDA converged to a model, where LF state is long-lived and MF state and HF states are interconvertible in agonist-bound $A_{2A}AR$.

Interestingly, the active-like HF state is also observed in the apo-receptor ensemble and even in the ZM241385-bound receptors. Additionally, the sample with the full agonist NECA has a higher population of the active-like HF state compared to the partial agonist LUF5834. The small variations in state populations between the apo-receptor and the antagonist-bound receptor are below statistical significance. We discuss below the implications of these results on the basal activity, partial agonism, and inverse agonism in $A_{2A}AR$.

Discussion

In this study, we used smFRET to investigate the conformational dynamics of A_{2A}AR. To preserve the native conformational dynamics of the receptor and minimize measurement-related artifacts we used four strategies. (1) As previous studies showed dramatic effects of commonly used detergents on GPCR conformational dynamics^{56,57} we reconstituted the receptor in nanodiscs that provide a relevant native-like lipid bilayer environment. (2) To minimize the effect of fluorophores on the receptor's dynamics we used small organic dyes attached to strategically engineered cysteines. (3) We avoided the need to remove native cysteines and associated to that potential structural perturbations by using the previously developed in-membrane labeling procedure⁴⁹. We showed that the mutant form of the receptor retains functional activity using the thermal shift assay and cAMP signaling assay in live HEK293T cells. (4) Finally, we studied receptors freely diffusing in solution and therefore excluded any artifacts related to their immobilization.

MFD-PIE fluorescence microscopy allowed us to filter out unlabeled and single-dye labeled receptors and to measure FRET efficiency with sub-millisecond temporal resolution in the double-labeled subpopulation of the receptors. Using various burst-wise fluorescence analysis techniques, we revealed sub-millisecond conformational dynamics in A_{2A}AR. Deviation of bursts from the 'static FRET line' on the FRET efficiency versus donor fluorescence lifetime plot indicated nanosecond-millisecond dynamics for the apo-A_{2A}AR and A_{2A}AR with each of the used ligands (Fig. 2E). FRET-2CDE analysis suggested more pronounced conformational dynamics in the agonist-bound A_{2A}AR than in the apo or antagonist-bound A_{2A}AR (Fig. 2F). BVA confirmed that the variations of FRET efficiency among ~100 μs time-bins exceed the level expected from shot-noise (Fig. 2G). fFCS demonstrated two components in A_{2A}AR dynamics: fast microsecond-time (3-20 μs)

dynamics present in all samples and slower ($390 \pm 80 \mu\text{s}$) dynamics evoked with agonists (Fig. 2H). This fFCS result puts all our findings in a single self-consistent picture: both fast and slow dynamics contribute to the deviation of bursts from the ‘static FRET line’, however the fast dynamics makes almost no contributions to the FRET-2CDE scores and to the BVA distribution deviations because of their 10-fold faster timescale compared to the temporal resolution of these techniques. Meanwhile, the slower dynamics evoked with the agonists explains the increased dynamics scores in FRET-2CDE and BVA for the agonist-bound $A_{2A}AR$. Finally, using dynamic PDA we proposed a three-state model of the $A_{2A}AR$ conformational dynamics that could fit the measured FRET efficiency histograms consistently with the fFCS findings (Fig. 3).

In our final three-state model of the $A_{2A}AR$ conformational space (Fig. 3B), the inactive-like MF and active-like HF states are not interchangeable in sub-millisecond time domain in the apo- and antagonist-bound $A_{2A}AR$, but can interconvert on a 300-500 μs timescale in the agonist-bound $A_{2A}AR$. The faster dynamics (3-20 μs) that contribute to conformational plasticity within these states is not sensitive to ligand binding and, therefore, likely, reflect the dynamics of the dye linkers or local fluctuations of TM6 and H8. The least populated (LF) state, presumably, corresponds to receptors locked in a long-lived non-functional state or improperly folded. Thus, our data reveal two modes of relative motions of fluorescent labels attached to residues L225C^{6,27} and Q310C^{8,65} in $A_{2A}AR$: fast microsecond-time (3-20 μs) dynamics, present in all samples regardless of the bound ligand, and slower (300-500 μs) dynamics, evoked with agonists. In a good agreement with our findings, sub-millisecond agonist-induced conformational dynamics of the intracellular part of TM7¹⁷ (Y290W^{7,55}) and the N-terminal end of H8²⁰ (I292M^{8,47}) have been shown for $A_{2A}AR$ by NMR. Although sub-millisecond dynamics between inactive-like conformations of TM6 (V229C^{6,31}) in the apo-

A_{2A}AR have been reported¹⁹, our data suggest only limited dynamics in the ligand-free and antagonist-bound A_{2A}AR.

In our experiments we could not measure slow conformational dynamics (>2 ms), because of the short residence time of individual molecules in the microscope focal spot. Our data do not indicate long-lived states in the agonist-bound A_{2A}AR (besides the ligand-insensitive LF state), but the FRET-2CDE analysis shows only moderate dynamics scores, and the observed deviations from the ‘static FRET line’ can be explained by a microsecond plasticity within a long-lived conformation. Consequently, we cannot exclude that long-lived conformations can coexist with conformations that show sub-millisecond dynamics. Keeping this in mind, we nevertheless did not introduce any additional long-lived states into our final model of the A_{2A}AR conformational space to avoid overfitting. Previous studies based on NMR provide complementary insights into the dynamics of long-lived (>2 ms) A_{2A}AR conformations^{19–21}.

The observed increase in FRET efficiency upon agonist binding was unexpected based on the available crystal structures of A_{2A}AR, which predicted a decrease in the FRET efficiency, because the distance between the C α -atoms of the labeled residues (L225 and Q310) increases from ~40 Å in the antagonist-bound structure (PDB: 3EML⁵⁸) to ~47 Å in the fully-active structure (PDB: 5G53⁵⁹). Moreover, previously reported A_{2A}AR-based FRET-sensors with fluorescent proteins at ICL3 and C-terminus showed a decrease in the FRET efficiency upon receptor activation^{42,43}. To explain this discrepancy, we proposed that the dyes are not randomly distributed (Fig. 1B), but rather occupy preferred locations within the volume accessible with their ~15 Å linkers. To test this hypothesis we performed 1- μ s long molecular dynamics (MD) simulations, which revealed that the dye attached to TM6 might preferentially locate between the intracellular tips of TM3 and TM5 in the inactive conformation and enter the G-protein-binding cavity of the receptor in the active

conformation (Fig S8). These preferred conformations of the dye would result in a significant decrease of the mean inter-dye distance upon receptor activation (from 5 nm to 3 nm), which would in turn lead to an increase in the mean FRET efficiency. Thus, our MD simulations provide a plausible explanation for the observed increase in the FRET efficiency upon A_{2A}AR activation.

The PDA analysis of our data suggests that the partial agonist LUF5834 and the full agonist NECA stabilize 53±8 % and 76±3 % of A_{2A}AR molecules, respectively, in the same active-like HF conformation. A similar mechanism for partial agonism in A_{2A}AR has recently been demonstrated via NMR with isotope-labeled methionine residues located in different structural domains (I106M^{3,54}, M140^{4,61}, M211^{5,72}, and I292M^{8,47}) of the receptor²⁰. On the other hand, two other NMR-based studies have suggested that LUF5834 either stabilizes a distinct, not a fully active conformation¹⁹, or has no effect on the A_{2A}AR conformation²¹. Our data do not support the existence of a separate partially active conformation of A_{2A}AR stabilized with LUF5834 that would be distinct from the fully active conformation stabilized with NECA. On the other hand, we cannot exclude that such partially active state could not be resolved in our data because small differences in FRET efficiency, the high photon shot-noise in single-molecule experiments, or the broadening of the FRET-distribution due to variations of photophysical parameters of the dyes.

Additionally, we observed that 20±3 % of apo A_{2A}ARs exhibit an active-like conformation, which could translate into a moderate basal activity of the receptor. Two previous NMR-based studies have addressed the molecular mechanisms of A_{2A}AR basal activity. One study reported a 70% population of pre-active and fully active states in the apo-ensemble¹⁹. Another study has reported negligible basal activity and showed that in the agonist-bound A_{2A}AR unique previously unpopulated conformations emerge²¹. A recent review suggests

that discrepancies between these two works could arise from differences in used constructs, ^{19}F reporters, their attachment sites, or in selected membrane mimicking systems (MNG/CHS versus DDM/CHS micelles)⁶⁰. The contradictory estimations of the basal activity of $\text{A}_{2\text{A}}\text{AR}$ should be put in the context of a similar heterogeneity of results provided by cell-based signaling assays. In different experiments, the basal activity of $\text{A}_{2\text{A}}\text{AR}$ was reported to reach from 0-20%^{58,61–63}, to 20-40%^{43,64,65} or even 40-70%^{66–69}. Cell assays are affected by different $\text{A}_{2\text{A}}\text{AR}$ expression levels and cell lines used^{62,68}. It has been shown that the C-terminal truncation of $\text{A}_{2\text{A}}\text{AR}$ impairs its basal activity - this can play an important role for our study as well as previous NMR-based works⁶⁴.

Finally, our measurements show that ZM241385 does not change the distribution of FRET efficiency compared to apo conditions and therefore we do not observe inverse agonism of ZM241385. Because many studies reported negligible basal activity of $\text{A}_{2\text{A}}\text{AR}$, ZM241385 is widely referred to as $\text{A}_{2\text{A}}\text{AR}$ antagonist^{11,58,70}. The recent ^{19}F NMR study, where no basal activity was detected for $\text{A}_{2\text{A}}\text{AR}$, correspondingly did not register any conformational changes induced with ZM241385²¹. On the other hand, those works that identified significant basal activity of $\text{A}_{2\text{A}}\text{AR}$ frequently reported inverse agonism of ZM241385^{43,62,63,65,68,71}. In line with these findings, the ^{19}F -NMR study that has reported 70% basal activity also showed inverse agonism of ZM241385¹⁹. Notably, it was previously shown that ZM241385 can lose inverse agonist activity if tested not in cells, but in isolated membranes⁶⁴. This latter result suggests that intracellular interaction partners can play an important role in both basal activity and inverse agonism, explaining both heterogeneity in published functional data and our results.

The multi-state conformational behavior of GPCRs delineates their complex pharmacology and, therefore, challenges modern drug design. We believe that new methods showing how

GPCR activity is modulated on a molecular level will facilitate the design and discovery of drugs with novel beneficial properties. Here we demonstrated a novel strategy to observe conformational dynamics of a GPCR in solution, yet in a close-to-physiological environment of lipid nanodiscs using intramolecular smFRET measured via the MFD-PIE approach. Our measurements combined fluorescence intensity, lifetime, and anisotropy information to characterize the sub-millisecond conformational dynamics of TM6 and H8 in A_{2A}AR and shed light on molecular mechanisms of basal activity and partial agonism in the receptor. The general strategy developed in our work can be extended to study the effects of various modulators (ligands, ions, lipids, etc.), membrane-mimicking systems (micelles, lipid nanodiscs, liposomes, etc.) and genetic modifications on the activity of A_{2A}AR and, in perspective, other GPCRs.

Methods

Protein expression, purification and labelling

The gene encoding the human A_{2A} adenosine receptor (1-316 aa) (UniProt C9JQD8) was synthesized *de novo* (Eurofins). The nucleotide sequence was optimized for *Leishmania tarentolae* expression with the GeneOptimizer software (ThermoFisher Scientific). KpnI restriction site was introduced at the C-terminus and used for polyhistidine tag (H9) fusion. The final construct was cloned into the integrative inducible expression vector pLEXSY_I-blecherry3 (Jena Bioscience, Germany) via the BglII and NotI restriction sites. L225C^{6,27} and Q310C^{8,65} mutations were introduced by PCR.

Leishmania tarentolae cells of the strain LEXSY host T7-TR (Jena Bioscience) were transformed with the A_{2A}AR expression plasmids linearized by the SmaI restriction enzyme. After the clonal selection, the transformed cells were grown at 26 °C in the dark in shaking baffled flasks in the Brain-Heart-Infusion Broth (Carl Roth, Germany) supplemented with 5 µg/mL Hemin (AppliChem), 50 U/mL penicillin and 50 µg/mL streptomycin (both antibiotics from AppliChem). When OD₆₀₀=1 was reached, 10 µg/mL tetracycline was added, and incubation continued for additional 24h.

The harvested cells were disrupted in an M-110P Lab Homogenizer (Microfluidics) at 10,000 psi in a buffer containing 50 mM $\text{NaH}_2\text{PO}_4/\text{Na}_2\text{HPO}_4$, pH 7.6, 0.2 M NaCl, 20 mM KCl, 10 mM MgCl_2 , 10% glycerol (w/v), 1 mM EDTA, 2 mM 6-aminohexanoic acid (AppliChem), 50 mg/L DNase I (Sigma-Aldrich) and cOmplete protease inhibitor cocktail (Roche). The membrane fraction of the cell lysate was isolated by ultracentrifugation at 120,000 g for 1 h at 4 °C. The pellet was resuspended in the same buffer but without DNase I and stirred for 1h at 4 °C. The ultracentrifugation step was repeated again.

Finally, the membranes were resuspended in the labelling buffer containing 50 mM HEPES, pH 7.0 10 mM MgCl_2 , 20 mM KCl, 2 mM 6-aminohexanoic acid, and cOmplete and mixed with Atto643 maleimide (ATTO-TEC) and Alexa488 maleimide (Invitrogen), dissolved in dimethyl sulfoxide (0.5 mg of each fluorescent label per 10 g of cells). Labeling reactions were carried out overnight in the dark at 4 °C on a roller mixer.

The next day, membrane fractions were pelleted by ultracentrifugation at 120,000 g for 1 h at 4 °C and washed twice with the labelling buffer for removal of unbound fluorescent labels. For solubilization, membranes were resuspended in a buffer containing 20 mM HEPES, pH 8.0, 800 mM NaCl, 5 mM MgCl_2 , 10 mM KCl, 2 mM 6-aminohexanoic acid, cOmplete with 4 mM theophylline (Sigma-Aldrich) and 1% n-Dodecyl β -maltoside (DDM) (Glycon Biochemicals) / 0.2% cholesteryl hemisuccinate (CHS) (Merck) (w/v) and left on the stirrer for 2 h at 4 °C in the dark. The insoluble fractions were removed by ultracentrifugation at 120,000 g for 1 h at 4 °C. The supernatants were loaded on an Ni-NTA resin (Cube Biotech) and incubated in the batch-mode overnight in the dark at 4 °C.

The next morning, proteins bound to Ni-NTA resin were washed with 10 column volumes of the first washing buffer: 50 mM HEPES, pH7.5, 800 mM NaCl, 25 mM imidazole, 10 mM MgCl_2 , 8 mM ATP (Sigma-Aldrich), 2 mM 6-aminohexanoic acid, 0.1 mM phenylmethylsulfonyl fluoride, 4 mM theophylline, cOmplete, 0.1% DDM / 0.02% CHS. Then, columns were washed with 10 column volumes of the second washing buffer: 50 mM HEPES, pH 7.5, 800 mM NaCl, 50 mM imidazole, 2 mM 6-aminohexanoic acid, 0.1 mM phenylmethylsulfonyl fluoride, 4 mM theophylline, cOmplete, 0.1% DDM / 0.02% CHS (w/v). Finally, proteins were eluted with 5 column volumes of the elution buffer: 25 mM HEPES, pH 7.5, 800 mM NaCl, 220 mM imidazole, 2 mM 6-aminohexanoic acid, 0.1 mM phenylmethylsulfonyl fluoride, cOmplete, 0.1% DDM / 0.02% CHS (w/v). The eluates were subjected to size-exclusion chromatography on a Superdex 200 Increase 10/300 GL column (GE Healthcare Life Sciences) in a buffer containing 20 mM HEPES, pH 7.5, 800 mM NaCl,

1 mM EDTA, 2 mM 6-aminohexanoic acid, cOmplete, 0.05% DDM / 0.01% CHS (w/v). Fractions, corresponding to A_{2A}AR monomers, were pulled and subjected to nanodisc reconstitution.

Nanodisc reconstitution

Membrane Scaffold Protein 1D1 (MSP1D1) was expressed in *E.coli* using gene with an N-terminal 6XHis-tag and up stream TEV-protease site cloned into pET28a(+) (Addgene plasmid #20061⁵⁰). MSP1D1 was purified using IMAC⁷² with further cleavage of 6xHis-tag by TEV protease (Sigma-Aldrich). The lipid mixture of 1-palmitoyl-2-oleoyl-sn-glycero-3-phosphocholine (POPC): 1-palmitoyl-2-oleoyl-sn-glycero-3-phospho-(1'-rac-glycerol) (POPG) (Avanti Polar Lipids) in chloroform was prepared at a molar ratio 7:3. The lipid film was dried under a gentle nitrogen stream, followed by removal of the solvent traces under vacuum, and then solubilized in 200 mM sodium cholate. The purified A_{2A}AR in DDM/CHS micelles was mixed with MSP1D1 and the POPC:POPG lipids at a molar ratio A_{2A}AR:MSP1D1:lipids=0.2:1:60. The final sodium cholate concentration was adjusted to 20 mM, the typical final receptor concentration was 0.1 mg/mL. After 1 h incubation at 4 °C, the mixture was incubated with wet Bio-Beads SM-2 (Bio-Rad, 0.4 g of beads for 1 mL reaction, beads were washed in methanol and equilibrated with 20 mM HEPES, pH 7.5, 800 mM NaCl, 1 mM EDTA) overnight at 4 °C in the dark. The next morning, the beads were discarded and the supernatant was supplemented with a fresh portion of Bio-Beads for an additional 4 h incubation. Finally, A_{2A}AR reconstituted into nanodiscs was subjected to size-exclusion chromatography on a Superdex 200 Increase 10/300 GL column (GE Healthcare) in a buffer containing 20 mM HEPES, pH 7.5, 150 mM NaCl, 1 mM EDTA, 2 mM 6-aminohexanoic acid, cOmplete. Fractions containing labeled receptors were combined together and used for further experiments.

Thermal shift assay

To show that the A_{2A}AR mutant (L225C^{6,27}/Q310C^{8,65}) retains ligand-binding activity in lipid nanodiscs, we used the fluorescent thermal stability assay⁵¹. The studies were carried out on a Rotor-Gene Q 6 plex (QIAGEN) instrument at a heating rate of 2 °C/min and a temperature range of 25-90 °C. The excitation wavelength was set at 387 nm and the emission wavelength was 463 nm. The A_{2A}AR concentration was about 2 µM. Buffer conditions: 20 mM HEPES, 150 mM NaCl, 1 mM EDTA, 2 mM 6-aminohexanoic acid, pH 7.5. To obtain a good fluorescent intensity we used a 2.5-fold molar excess of CPM dye (7-Diethylamino-3-(4'-

Maleimidylphenyl)-4-Methylcoumarin, Invitrogen) to protein. To prepare protein for the ligand-binding measurements we added 200 μ M of ZM241385 or NECA and incubated for 1h in the dark at +4 $^{\circ}$ C. The thermal denaturation assay was performed in a total volume of 50 μ L (Fig. S1E).

Measurement of A_{2A}AR surface expression and Gs-signaling

For A_{2A}AR functional assays, the A_{2A}AR (WT or Q310C^{8.65}/L225C^{6.27} mutant) gene (GenScript) was optimized for eukaryotic expression with an N-terminal hemagglutinin signal sequence (MKTIIALSYIFCLVFA) followed by the FLAG tag epitope (DYKDDDDK) and C-terminal 10 $\times$ His tag were cloned into pcDNA3.1(-) at BamHI(5') and HindIII(3'). The surface expression of A_{2A}AR was determined by the whole-cell ELISA assay⁷³. Briefly, HEK293FT cells were seeded in a 100 mm cell culture plate and transfected separately with 10 μ g of each expression plasmid DNA (pcDNA3.1(-)_A_{2A}AR(WT), pcDNA3.1(-)_A_{2A}AR(Q310C^{8.65}/L225C^{6.27}) or pcDNA3.1(-) as a negative control) using a common Lipofectamine 3000 protocol. The plates were incubated for additional 12–18 h at 37 $^{\circ}$ C, 5% CO₂. The HRP-conjugated anti-FLAG M2 antibody (A8592, Sigma) at a dilution of 1:2000 in TBS with 1% protease-free BSA (A3059, Sigma) and TMB ready-to-use substrate (T0565, Sigma) were used for the ELISA procedure. For normalization on cells quantity Janus Green B (Sigma) staining was used, and the absorbance ratio A₄₅₀/A₅₉₅ was calculated. Measurements were performed in triplicate for WT and mutant A_{2A}AR as well as for empty-vector transfected cells. Measured values of A₄₅₀/A₅₉₅ were normalized so that the mean expression level of WT A_{2A}AR was 100% ($F_{WT} = 100 \pm 6$ %, SDs for n=3 measurements are given). The double mutant form of the receptor showed only slightly lower expression level than WT: $F_{L225C/Q310C} = 73 \pm 7$ %. Empty-vector transfected cells showed only marginal anti-FLAG antibody binding: $F_{EV} = 1 \pm 1$ %.

For evaluation of the A_{2A}AR signaling activity, we checked the effect of the agonist NECA on cAMP responses in transfected cells. For cAMP determination, we used the Bioluminescence Resonance Energy Transfer (BRET) approach with the EPAC biosensor⁷⁴. The cAMP BRET biosensor was kindly provided by professor Raul Gainetdinov⁷⁵. Transfections were carried out with Lipofectamine 3000 (Thermo) using HEK293T cells seeded in a 100 mM cell culture plate, receptor cDNA vectors pcDNA3.1(-)_A_{2A}AR(WT), pcDNA3.1(-)_A_{2A}AR(Q310C^{8.65}/L225C^{6.27}) (10 μ g each) and the EPAC biosensor cDNA vector (1 μ g) needed for evaluation of the cAMP production. Transfected cells were split into

96-well plates at 10^5 cells per well. On the following day, 70 μ L of PBS were added to each well followed by addition of 10 μ L of a 50 μ M coelenterazine-h solution (Promega). After 10-min incubation, either 10 μ L of buffer or 10 μ L of NECA at different concentrations in PBS were added, and the plate was then placed into a CLARIOstar reader (BMG LABTECH, Germany) with a special BRET filter pair (475 ± 30 nm – coelenterazine-h and 530 ± 30 nm – YFP). The BRET signal was calculated as the ratio of the light emitted at 530 nm to the light emitted at 480 nm. Three independent experiments with three technical replicas in each were conducted. For pEC₅₀ evaluation, dose-response curves from three technical replicas were averaged and analyzed. Mean and S.D. of pEC₅₀ among three biological samples were calculated (Fig. S1F).

Confocal MFD-PIE setup

For single-molecule experiments, a home-built multi-parameter fluorescence detection microscope with pulsed interleaved excitation (MFD-PIE)³⁶ was used (see scheme of the setup in Fig. S2). Two lasers were used: a pulsed 483-nm laser diode (LDH-P-C-470, Picoquant) and a pulsed 635-nm laser diode (LDH-P-C-635B, Picoquant), with alternating at 26.67 MHz pulses, delayed by 18 ns with respect to each other. Sample emission was transmitted through a pinhole and spectrally split. Both, the blue range and red range were split by polarization into two detection channels. Photons were detected by four avalanche photodiodes (PerkinElmer or EG&G SPCM-AQR12/14, or Laser Components COUNT BLUE): $B_{\parallel}$ (blue-parallel), $B_{\perp}$ (blue-perpendicular), $R_{\parallel}$ (red-parallel) and $R_{\perp}$ (red-perpendicular) (Fig. S2), which were connected to a TCSPC device (SPC-630, Becker & Hickl GmbH). Microscope alignment (excitation light guiding, objective lens correction collar, pinhole, detectors) was done using real-time fluorescence correlation spectroscopy (FCS) on freely diffusing Atto488-COOH and Atto655-COOH in water. For more details about the used equipment the reader is referred to ref.⁷⁶

smFRET data recording

Samples of double-labelled A_{2A}AR in nanodiscs were diluted in a buffer, containing 20 mM HEPES, pH 7.5, 150 mM NaCl, 1 mM EDTA, 2 mM 6-aminohexanoic acid to a protein concentration of 0.5-2 nM. To measure the effects of ligand binding, samples were supplemented with either 10 μ M ZM241385, 10 μ M LUF5834 or 10 μ M NECA and incubated for 30 min at +4 °C. After the incubation, the samples were transferred to a Nunc

Lab-Tek Chambered coverglass (Thermo). smFRET experiments were performed at 100 μ W of 483 nm and 50 μ W of 635 nm excitation. Measurements were recorded at room temperature (22 °C), samples were replenished every 30 min. With all filters applied (see *Selection of double-labeled, donor-only and acceptor-only subpopulations*), 9,000 – 12,000 bursts corresponding to double-labeled molecules were collected for each sample: 11,961 for apo, 10,167 burst for ZM241385, 9,557 for LUF5834, and 11,007 for NECA. Background scattering information was obtained via a buffer measurement under identical condition.

Software

All simulations and analyses of experimental data were performed in the software package PAM (PIE Analysis with MATLAB)⁷⁷. The software is available as a source code, requiring MATLAB to run, or as pre-compiled standalone distributions for Windows or MacOS at <http://www.cup.uni-muenchen.de/pc/lamb/software/pam.html> and hosted in Git repositories under <http://www.gitlab.com/PAM-PIE/PAM> and <http://www.gitlab.com/PAM-PIE/PAMcompiled>. A detailed manual is located under <http://pam.readthedocs.io>. Details of smFRET data treatment are given in Supplementary methods.

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Author contributions

OV performed receptor expression. IM and OV performed receptor labeling, purification, and nanodisc reconstitution. TG and IM performed ensemble-TCSPC measurements. IM and QC measured fluorescence spectra. QC and JHe prepared the instrumentation for smFRET experiments. IM performed smFRET experiments, analyzed smFRET data and drafted the manuscript. JHe, ABa, SW, ABo and VB contributed to the analysis of smFRET data. AGe performed cell functional assay. PKh performed TSA functional assay with the contribution of AM, AL, AGu and PKu. PO performed molecular dynamics simulations. IM, VB, JHe,

VC, TG, ABo, AGu, AM, PKh, VG discussed the data, analysis, and contributed to writing the manuscript. IM, VB, JHe, VC, TG, JHo conceived the study. All authors commented on and edited the manuscript. VB, and JHe supervised the project.

Competing interests

Authors declare no competing interests.

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FIGURES AND TABLES

Fig. 1.

Agonist-induced conformational changes in $A_{2A}AR$ are revealed by smFRET. (A) Schematic illustration of the MFD-PIE smFRET experiment on $A_{2A}AR$ embedded in lipid nanodiscs and stochastically labeled with the donor (Alexa488) and the acceptor (Atto643) fluorescent dyes at TM6 and H8. Eight coexisting labeling variants of $A_{2A}AR$ are shown as shadowed receptors in both sides of the image, 'D' and 'A' correspond to donor and acceptor dyes, respectively. $A_{2A}AR$ s diffuse in solution and stochastically cross the focal spot of an inverted

fluorescence microscope. Bursts of fluorescence from donor and acceptor fluorophores are recorded within the 1-10 ms residence time of individual A_{2A}ARs crossing the focal spot. Only those receptors labeled with both donor and acceptor produce FRET signal. In the PIE approach, two spatially overlapped and alternately pulsing lasers are focused by the microscope objective to excite donor and acceptor fluorescence consequently. Using the MFD approach, fluorescence signals of donor and acceptor are recorded separately, and the fluorescence lifetime and anisotropy of each dye are determined. **(B)** The labeled sites (L225^{6,27}, Q310^{8,65}) and the volume accessible for the dyes (simulated using FPS software⁷⁸) are shown on the A_{2A}AR structure (PDB: 3EML⁵⁸), the membrane boundaries (dashed lines) are obtained from the PPM web server⁷⁹ and shown as dashed lines **(C)** Burst-wise distributions show an agonist-induced increase in FRET efficiency in the double-labeled A_{2A}AR. The number of bursts used for the analysis (N) is given for each condition.

Fig. 2.

Four complementary burst-wise analysis approaches suggest an agonist-induced increase in the sub-millisecond conformational dynamics of A_{2A}AR. Contour plots are two-dimensional histograms of different fluorescence burst parameter distributions. Simulated plots for 'static' and 'dynamic' molecules (**A,C,E,G**), and experimental data for double-labeled A_{2A}AR (**B,D,F,H**) are shown. (**A, B**) The FRET efficiency is plotted against donor fluorescence lifetime. The 'static FRET' line is shown in red. A shift of burst distribution to the right from the red line indicates dynamic FRET. (**C, D**) The FRET-2CDE dynamics score is plotted against FRET efficiency. The FRET-2CDE=20 threshold is indicated as red dashed lines, and the percentage of bursts with FRET-2CDE>20 is shown in red text. (**E, F**) BVA dynamics-scores are plotted against FRET efficiency. Red diamonds show the centers of burst

subgroups equally spaced along the FRET efficiency axis. The solid black lines show mean BVA-scores, and the transparent grey areas demonstrate 99.9% confidence intervals expected for static molecules, given the shot-noise present in the data. **(G, H)** The cross-correlation fFCS function is plotted against time lag. Experimental points are shown in black; the error bars were estimated by statistical bootstrapping. The fitted curve is shown in orange; the exchange times derived from the fit are highlighted with vertical red lines. The number of fluorescence bursts used for the analysis are the same as for fig. 1C: 10,167 for ZM241385, 11,961 for apo-state, 9,557 for LUF5834, 11,007 for NECA.

Fig. 3.

PDA quantifies parameters of the $A_{2A}AR$ three-state action model by fitting FRET efficiency distributions. **(A)** Experimental distributions of 1-ms-long time-bins derived from fluorescence bursts of double-labeled $A_{2A}AR$ (grey area) were fitted with a three state-model. The resulting fit (black line) is a sum of distributions simulated for molecules that stay in the LF ($R = 57.9 \pm 6.0$ Å, light green line), MF ($R = 50.0 \pm 2.1$ Å, dark cyan line), or HF ($R = 45.1 \pm 4.9$ Å, red line) state during the entire simulated time-bin, and the distribution for molecules that sample both MF and HF states within a time-bin (orange line). The fitting residuals are shown on the top of each panel. The bar charts on the right show relative populations of the three states, with error bars representing SD of $n = 3$ independent data subsets. **(B)** The three-state action model of $A_{2A}AR$ and corresponding energy landscapes for the apo and agonist-bound receptor demonstrate relative populations of the states and inter-

state exchange times. $\tau_{ex}=(k_{12}+k_{21})^{-1}$ is the relaxation time of the exchange between the MF and HF states. TM6 is colored on the schematic (cylinder) representation of active (PDB: 5G53⁵⁹, red) and inactive (PDB: 3EML⁵⁸, dark cyan) structures of A_{2A}AR.