

ARTICLE

Received 7 Mar 2012 | Accepted 30 Jul 2012 | Published 4 Sep 2012

DOI: 10.1038/ncomms2028

The allosteric vestibule of a seven transmembrane helical receptor controls G-protein coupling

Andreas Bock¹, Nicole Merten², Ramona Schrage¹, Clelia Dallanocce³, Julia Bätz⁴, Jessica Klöckner⁵, Jens Schmitz⁵, Carlo Matera³, Katharina Simon², Anna Kebig¹, Lucas Peters², Anke Müller², Jasmin Schrobang-Ley¹, Christian Tränkle¹, Carsten Hoffmann⁴, Marco De Amici³, Ulrike Holzgrabe⁵, Evi Kostenis² & Klaus Mohr¹

Seven transmembrane helical receptors (7TMRs) modulate cell function via different types of G proteins, often in a ligand-specific manner. Class A 7TMRs harbour allosteric vestibules in the entrance of their ligand-binding cavities, which are in the focus of current drug discovery. However, their biological function remains enigmatic. Here we present a new strategy for probing and manipulating conformational transitions in the allosteric vestibule of label-free 7TMRs using the M₂ acetylcholine receptor as a paradigm. We designed dualsteric agonists as 'tailor-made' chemical probes to trigger graded receptor activation from the acetylcholine-binding site while simultaneously restricting spatial flexibility of the receptor's allosteric vestibule. Our findings reveal for the first time that a 7TMR's allosteric vestibule controls the extent of receptor movement to govern a hierarchical order of G-protein coupling. This is a new concept assigning a biological role to the allosteric vestibule for controlling fidelity of 7TMR signalling.

¹ Pharmacology and Toxicology Section, Institute of Pharmacy, University of Bonn, Gerhard-Domagk-Straße 3, 53121 Bonn, Germany. ² Molecular-, Cellular- and Pharmacobiology Section, Institute of Pharmaceutical Biology, University of Bonn, Nussallee 6, 53115 Bonn, Germany. ³ Dipartimento di Scienze Farmaceutiche, Sezione di Chimica Farmaceutica 'Pietro Pratesi', Università degli Studi di Milano, Via Mangiagalli 25, 20133 Milano, Italy. ⁴ Department of Pharmacology and Toxicology, University of Würzburg, Versbacher Strasse 9, 97078 Würzburg, Germany. ⁵ Department of Pharmaceutical Chemistry, Institute of Pharmacy, University of Würzburg, Am Hubland, 97074 Würzburg, Germany. Correspondence and requests for materials should be addressed to K.M. (email: k.mohr@uni-bonn.de) or to E.K. (email: kostenis@uni-bonn.de).

Seven transmembrane helical receptors (7TMRs; G-protein-coupled receptors) are embedded into the membrane of almost all cells and are essential for orchestrating the function of cells in order to maintain homeostasis of the organism and its adaptation to environmental changes. In addition, modulation of 7TMR function by drugs is a mainstay for the prevention and treatment of disease¹. The human genome encodes about 800 7TMRs with the rhodopsin-like class A receptors representing the quantitatively dominating subfamily, which include aminergic and cholinergic receptors². 7TMR activation by an extracellular stimulus is communicated to the cell by a conformational rearrangement of the intracellular domain region that mediates recruitment and activation of adaptor proteins. These proteins control the activity of intracellular signalling pathways and include guanylnucleotide-binding proteins (G proteins) and β -arrestins³.

Heterotrimeric G proteins are subdivided into four families based on sequence homologies and functional similarities of their G α subunits, G α_i , G α_s , G $\alpha_{q/11}$ and G $\alpha_{12/13}$ (ref. 4). These can be addressed selectively, but often an endogenous activator-bound 7TMR has efficacy to activate more than one class of G protein, which is referred to as promiscuous signalling. Remarkably, especially from a therapeutic point of view, chemically modified artificial ligands may engender more precise signalling than the endogenous activator molecule, which is designated as biased signalling, functional selectivity or signal trafficking^{5–7}. It is likely that the signalling repertoire of a 7TMR corresponds to ensembles of possible conformations of the receptor protein, and that a biased activator allows only for a subset of possible receptor conformations⁸.

Crystal structures of 7TMRs give fascinating insight into the conformational rearrangements that accompany inactive-to-active transitions^{9–11}. However, the molecular principles by which a receptor protein translates the chemical properties of a ligand into a signalling fingerprint are still unresolved¹². Accumulating evidence suggests that there is no consensus sequence for adaptor protein-coupling specificity on the level of the intracellular domains¹³, and, in line with this notion, signalling is often promiscuous. On the level of the helical bundle region, 7TMR structure is highly conserved, which would correspond well to a general activation mechanism¹⁴ but not to coupling selectivity. In the extracellular loop region, however, there is rather high variability between primary structures. Remarkably, a ligand-dependent conformational coupling from the orthosteric site to the extracellular loop region has recently been described¹⁵.

Intriguingly, class A 7TMRs harbour allosteric vestibules in the extracellular entrance of their ligand-binding cavities, which receive high attention as new targets for drug discovery¹⁶. However, until now, their biological function is poorly understood. The question therefore arises whether the extracellular domains hold a function in controlling intracellular signalling of a 7TMR.

We took advantage of a new strategy for probing the allosteric vestibule's role for signal trafficking using the M₂ acetylcholine (ACh) receptor as a paradigm for small ligand 7TMRs. Muscarinic ACh receptors are important targets for novel drugs with improved selectivity¹⁷, and in particular, fundamental insights of 7TMR function have first been achieved with this receptor. These include the discovery of 7TMR allosterism¹⁸, the establishment of the ternary complex model of allosteric interactions¹⁹, its proof with a radiolabelled allosteric probe²⁰ and the identification of allosteric key epitopes as well as their spatial orientation within the allosteric vestibule^{21–23}. We designed a set of 'tailor-made' chemical probes, referred to as dualsteric agonists²⁴, to trigger graded receptor activation from the ACh-binding site (that is, the orthosteric site) while simultaneously restricting spatial flexibility of the receptor's extracellular allosteric vestibule.

Evaluation of probe actions in this study in a broad array of signalling readouts, including those in a space-providing 'rescue mutant', show that the allosteric vestibule of the M₂ receptor

controls the extent of receptor movement thereby governing a hierarchical order of intracellular adaptor protein recruitment. This is a new concept assigning an unprecedented biological role to the allosteric vestibule for controlling fidelity of receptor signalling of a small molecule 7TMR.

Results

Strategy for new chemical tool design. Muscarinic ACh receptors belong to the class A of G-protein-coupled receptors, which bind their endogenous agonist deep in the ligand-binding cleft lined by transmembrane helical domains^{25–27}. Asp103^{3,32} (Fig. 1a) serves as a docking point for the positively charged nitrogen of ACh. Iperoxo ((1) in Fig. 1b), the orthosteric building block of the dualsteric compounds, is a super-potent muscarinic agonist characterized by an affinity-enhancing Δ^2 -isoxazoline ring system and was re-synthesized as previously described²⁸. Iperoxo is 100-fold more potent than ACh as reported below and was chosen as the orthosteric building block of the dualsteric probes to ensure high-fidelity receptor activation via the orthosteric-binding site. To study the role of the allosteric substituent for signalling (shaded grey in Fig. 1b), iper-6 ((2) in Fig. 1b) was synthesized (Supplementary Methods), which carries the linker chain without the spacious allosteric building blocks. The allosteric phthalimide building block contained in iper-6-phth ((3) in Fig. 1b) has no efficacy for receptor activation²⁹ and binds to M₂-W422^{7,35} at the beginning of TM7 and to M₂-Y177^{5,32} in the extracellular loop 2 (Fig. 1a)^{21–23}. These two amino acids are located opposite to each other in the depth of the allosteric vestibule in close vicinity to the subjacent orthosteric site²³. The crystal structure of the inactive antagonist-bound M₂ receptor proves the location of M₂-Y177^{5,32} and M₂-W422^{7,35} outside of and above the aromatic cage that encloses the orthosteric ligand, respectively²⁵. Docking simulations show that the allosteric building block protrudes to the level of M₂-W422^{7,35} (ref. 24). If this binding topography causes a signalling bias, enlargement of the substituent to naphthalimide (in iper-6-naph, (5) in Fig. 1b), which by itself also has no efficacy for receptor activation²⁹, should further impair signalling. Iper-6-phth and iper-6-naph were re-synthesized precisely as described previously²⁴. In contrast, elongation of the hexamethylene middle chain to octamethylene, as realized in the newly synthesized iper-8-phth and iper-8-naph ((4) and (6) in Fig. 1b, respectively, Supplementary Methods), would be expected to restore the signalling by shifting the spacious substituent outward of this critical region.

Probe binding to cell surface hM₂-receptors. Ligand–receptor interactions in live CHO cells transfected to stably express hM₂ receptors (CHO-hM₂ cells) were measured using the non-permeant orthosteric radioligand [³H]-N-methylscopolamine ([³H]NMS) to clarify prerequisites for the analysis of probe signalling.

Dualsteric ligands may principally bind in two orientations, the dualsteric mode (pose 1 in Fig. 1c) and the purely allosteric mode (pose 2 in Fig. 1c). As receptor activation is encoded by their orthosteric agonist building block, the probes of this study will be agonistic only in the dualsteric posture. Iper-6-naph was chosen for validation of dualsteric binding, because, first, the naphthalimide residue has a higher allosteric M₂ receptor-binding affinity than phthalimide, and, second, the six-membered linker is less suited for an unconstrained dualsteric receptor interaction than the eight-membered linker (see below). Equilibrium and dissociation binding experiments revealed a pronounced preference of iper-6-naph for the dualsteric pose relative to the allosteric pose (Fig. 2).

M₂ receptors prefer coupling to G_i over G_s proteins^{30,31} (Fig. 1c). Pretreatment of CHO-hM₂ cells with pertussis toxin (PTX) to silence G_i- and disclose G_s coupling does not affect binding of iper-6-naph and other probes. Binding data are compiled in Supplementary Fig. S1 and Supplementary Table S1.

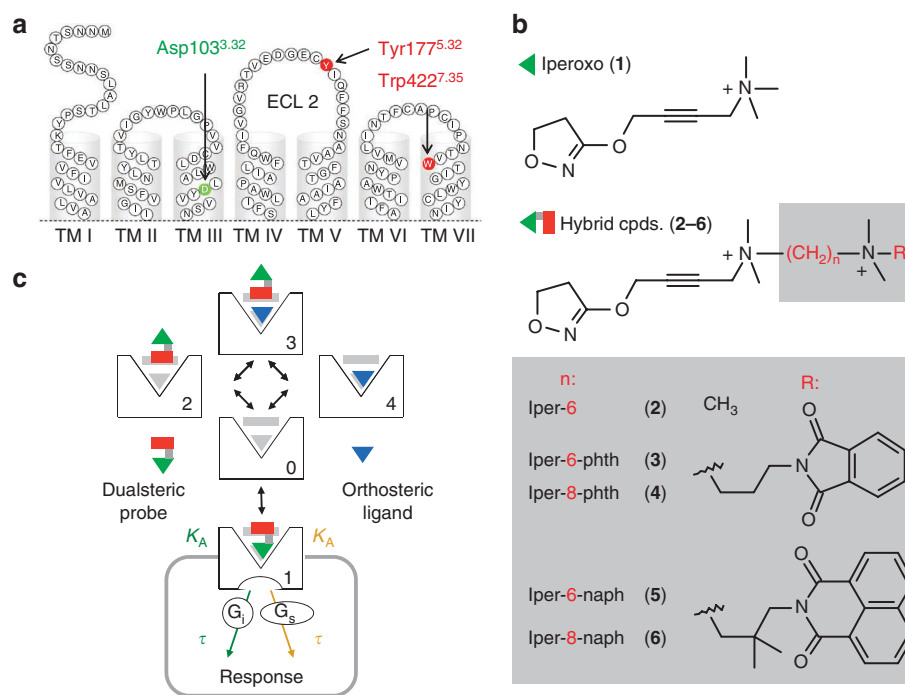


Figure 1 | Receptor binding poses of dualsteric probes. (a) Snake-like plot of the outer loop region of the muscarinic M_2 receptor along with the flanking alpha-helical domains. Arrows indicate essential epitopes for orthosteric (green) and allosteric (red) ligand binding. (b) Dualsteric hybrid design: the super high-affinity orthosteric muscarinic agonist iperoxo is linked via a hexamethylene- or octamethylene-spacer with allosteric inverse agonistic phthalimide or naphthalimide moieties. (c) Receptor binding poses of dualsteric probes in the absence (1,2) and presence (3) of an orthosteric ligand. With the hybrid design of this study, only the dualsteric binding mode (1) leads to receptor activation and downstream signalling. The coupling efficiency τ quantifies the efficacy with which agonist-occupied receptors activate G_i and G_s signalling. The coupling efficiency τ affects the binding constant K_A of the dualsteric agonist.

Dualsteric probes induce structure-dependent G_i bias. 7TMR proteins can be considered as dynamically fluctuating macromolecules that adopt ensembles of tertiary conformations⁵. Label-free techniques use native receptor proteins thus ensuring natural conformational flexibility and sensitivity to probe actions. Recently, we established dynamic mass redistribution (DMR) as a novel and powerful label-free technology to identify G-protein-dependent signalling in live cells^{32,33}. Of note, 7TMR signalling along a given pathway may yield cell type-dependent DMR responses³³. Here, we applied CHO-h M_2 cells. As CHO cells do not generate ACh esterase³⁴, they are well suited to quantify DMR responses to the endogenous transmitter ACh (Fig. 3a–d). ACh is known to activate G_i and G_s signalling of M_2 receptors^{30,31}. Native CHO-h M_2 cells respond to ACh in a concentration-dependent manner with brisk positive DMR (Fig. 3a), which suggests G_i pathway activation^{32,33}. Cells pretreated with PTX to chemically knockout G_i proteins respond to ACh with negative DMR (Fig. 3b). As this signal is completely sensitive to cholera toxin (CTX), which is known to mask G_s -mediated signalling^{32,33}, the negative DMR reflects G_s pathway activation (Supplementary Fig. S2). Therefore, positive DMR observed in unpretreated cells includes G_i - and G_s -mediated signalling events. However, the brisk peak DMR seen in unpretreated cells represents G_i activation as, first, this peak is still present in CTX-pretreated cells (Fig. 3c) and, second, potencies of ACh for inducing positive peak DMR in the absence and presence of CTX are in the same range (Fig. 3d, control: $pEC_{50} = 8.09 \pm 0.13$; CTX: $pEC_{50} = 7.49 \pm 0.16$; $P = 0.02$, unpaired t -test). The small, albeit significant, difference in potencies is likely to be caused by cellular adaptation to the CTX-induced rise of intracellular cyclic AMP (cAMP). In contrast, the potency of ACh for G_s activation ($pEC_{50} = 6.01 \pm 0.19$) is 100-fold lower compared with positive peak DMR and 30-fold lower relative to positive

peak DMR in CTX-pretreated cells ($P < 0.001$, one-way analysis of variance (ANOVA) with Dunnett's post test). Taken together, peak DMR in unpretreated cells represents G_i activation.

Dualsteric probes exhibit a G_i over G_s signalling bias compared with ACh and iperoxo (Fig. 3e–h), which induce equieffective promiscuous G_i and G_s signalling (compare DMR peaks in Fig. 3, panels e with g and f with h, respectively). Noteworthy, pretreatment with PTX did not affect receptor density ($B_{max-PTX} = 89\%$ of $B_{max-control}$, $P = 0.75$, unpaired t -test). Relative to these agonists, iper-6 has comparable maximum effects (Fig. 3g,h). In contrast, iper-6-phth and iper-8-phth, although being almost equieffective to orthosteric agonists with respect to G_i -DMR (Fig. 3e), hardly induce G_s -DMR (Fig. 3f). Of note, the middle chain-elongated iper-8-phth is more competent for G_s -DMR than iper-6-phth (Fig. 3f). In case of the spacious iper-x-naph analogues, even G_i -DMR is compromised with iper-6-naph (Fig. 3g), whereas the middle chain-elongated iper-8-naph clearly achieves maximum G_i -DMR (Fig. 3g). As found with iper-x-phth analogues, G_s -DMR is more compromised than G_i -DMR, especially in case of iper-6-naph (Fig. 3h) where G_s signalling is virtually silenced. Iper-8-naph only partially activates the G_s pathway (Fig. 3h), comparable to iper-8-phth (Fig. 3f). Peak DMR-values are enumerated in Supplementary Table S2.

Taken together, the DMR findings reveal that G_s pathway activation by the h M_2 receptor is far more sensitive to spacious allosteric substituents than G_i activation. In addition, probes indicate that the position of the spacious allosteric substituent within the extracellular binding cleft of the receptor protein is crucial for signalling.

Quantification of the probes' G_i bias. DMR measurements point to a structure-dependent signalling bias. We used standard G_i and G_s

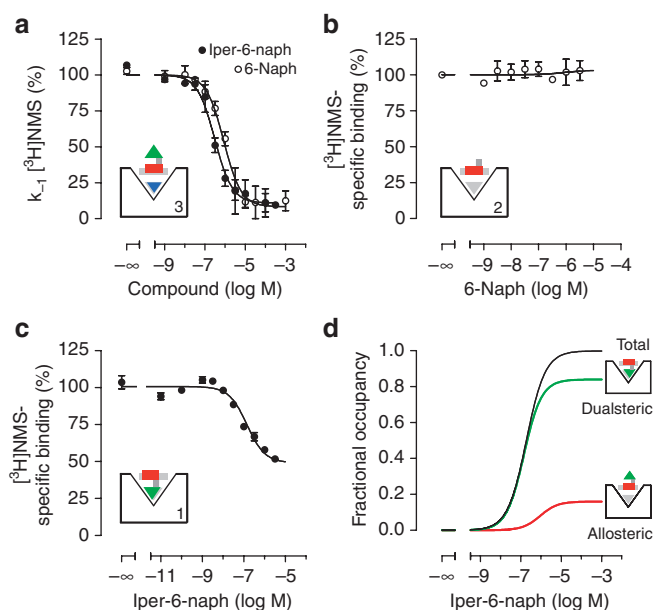


Figure 2 | Verification of dualsteric probe binding in live CHO-hM₂ cells. Iper-6-naph is chosen for verification of dualsteric binding in whole cells, because, first, the naph-building block has a higher binding affinity to the allosteric binding area than the phth-binding block and, second, signalling of iper-6-naph is most compromised among all probes, which could theoretically be due to a low probability of dualsteric binding (pose 1). ‘6-naph’ (equivalent to compound ‘A2’ in ref. 24 and re-synthesized precisely as described therein) serves as a surrogate for the allosteric building block of iper-6-naph. **(a)** After having determined radioligand ($[^3\text{H}]$ NMS)-binding characteristics (pose 4), the affinity of iper-6-naph for the radioligand-bound receptor (pose 3) is derived from probe-induced inhibition of radioligand-dissociation ($\text{EC}_{50, \text{diss}} = \alpha \times K_{\text{B, allo}}$). As to be expected iper-6-naph and 6-naph have almost the same affinity to the $[^3\text{H}]$ NMS-bound receptor. **(b)** Interaction of 6-naph with the radioligand under equilibrium binding conditions and analysis based on the ternary complex model is taken to represent the affinity of iper-6-naph for the purely allosteric binding pose 2. **(c)** Interaction of iper-6-naph with the radioligand under equilibrium conditions and data analysis including binding constants for poses 2 and 3 (modified after⁵⁵) yields the binding constant for the dualsteric mode (pose 1). **(d)** The ratio of binding constants for poses 1 and 2 displays the concentration-dependent fractional receptor occupancy of iper-6-naph (black curve) in the dualsteric (green curve, 84%) and allosteric (red curve, 16%) binding mode. Error bars indicate s.e.m. of 6–12 replicates from 2–4 independent experiments.

signalling readouts that are close downstream to receptor activation for quantifying the probes’ signalling bias.

$[^35\text{S}]$ GTP γ S binding in membrane suspensions of CHO-hM₂ cells reflects replacement of GDP by GTP at G protein α subunits as a consequence of receptor activation (Fig. 4a). This assay almost exclusively captures G_i activation³⁵. Intracellular cAMP formation in probe-activated CHO-hM₂ cells pretreated with PTX displays activation of the G_s pathway (Fig. 4b).

Regarding maximum effects, Fig. 4a,b shows—as observed in the DMR measurements—that iper-6-phth and even more so iper-6-naph are compromised with respect to the G_s assay relative to the G_i assay. Again, the middle chain-elongated derivatives show increased efficacy for G_s signalling (Fig. 4a,b and Supplementary Table S3).

The bias plot³⁶ (Fig. 4c) compares signalling via the G_i and G_s route at equal concentrations of test compound over the range of applied concentrations. The respective maximum effects of ACh define 100%. The hyperbolic curve of ACh reflects its preference for

G_i activation, which is designated as system bias. Relative to this system bias, all test compounds, except iperoxo, reveal a stronger G_i bias. On the receptor level, this may result from a higher affinity ratio of binding to G_i- versus G_s-coupled receptors. In addition or alternatively, the efficacy τ of an agonist-bound receptor may be higher for signalling via the G_i instead of the G_s pathway. Global fitting of the functional data (Fig. 4a,b) with the operational model of agonism³⁷ provides measures for both the binding constant (K_{A}) and the efficacy (τ) of the respective agonists. The τ/K_{A} -ratio³⁸ (transduction coefficient) encompasses both potential mechanisms of bias (for τ/K_{A} -ratios for both pathways normalized on ACh ($\Delta\log(\tau/K_{\text{A}})$) see Table 1). $\Delta\log(\tau/K_{\text{A}})$ -values³⁸ (Fig. 4d, Table 1) eliminate system bias and indicate true test compound bias relative to ACh. In general, dualsteric agonists are significantly G_i biased (Fig. 4d). Iper-8-naph is an exception; this compound has clearly lower efficacy on the G_s pathway (Fig. 4b) than ACh, which, however, is compensated by a higher binding affinity (low K_{A} -value; Supplementary Table S4). Figure 4e focusses on the efficacy component of ligand bias. Plotted is the efficacy $\log \tau$ of agonist-bound receptors for activating G_i and G_s signalling. The iper-6-naph-bound receptor has lowest signalling efficacy. The latter is restored by replacement of the naph-residue by the less voluminous phth-residue and even more by linker elongation from C6 to C8. Iper-8-naph and iper-8-phth attain an upper level of G_i activation, which is not surpassed by ACh and iperoxo, but are still clearly inferior with respect to G_s activation. Shortening of the dualsteric agonists to iper-6 and further to iperoxo then step-wise restores G_s signalling efficacy. These findings corroborate the notion that G_s signalling efficacy of the agonist-bound M₂ receptor is more sensitive to the allosteric residues of dualsteric probes than G_i signalling efficacy (Supplementary Table S4).

Middle chain elongation restores signalling at different levels.

As 7TMR ligands may have widely differing signalling competence depending on pathways (for review see refs 39,40), we investigated the probes’ signalling pattern in a set of additional readouts.

First, we focused on the receptor’s intracellular loops, as these mediate receptor contact with adaptor proteins. We used a FRET-sensor-hM₂-receptor, stably expressed in HEK293 cells, that has an enhanced cyan fluorescent protein (eCFP) fused to its C-terminus and a FLAsH-binding domain introduced within the third intracellular loop beneath transmembrane domain TM5, respectively^{41–44} (Supplementary Methods).

Agonist-induced conformational transitions of the inner loop region were diminished in the case of iper-6 and not significantly different from baseline in the case of iper-6-phth and iper-6-naph (Fig. 5a). Intracellular loop movements reappeared with the iper-8-X compounds. Thus, the allosteric substituents introduced into the orthosteric iperoxo and pointing to the extracellular surface of the receptor protein severely restrict inner loop region dynamics. Noteworthy, shifting the spacious residues further outward in the allosteric region by middle chain elongation unleashes orthosteric agonist-like intracellular loop rearrangement.

Next, we examined conformational transitions of the G_i protein (Fig. 5b and Supplementary Methods). Therefore, the activation-promoted rearrangement of the α -subunit relative to the $\beta\gamma$ -subunit was analysed using a protein–protein-interaction-based bioluminescence resonance energy transfer (BRET) assay⁴⁵ in COS7-cells. These were transiently co-transfected with the hM₂ receptor and the Renilla luciferase-tagged G α _{i1}-subunit along with G β and a GFP-tagged γ -subunit. All probes induce a conformational rearrangement of the G_i protein with the iper-8-X compounds displaying a higher efficacy than the respective iper-6-X probes. Compared with the inner loop-assay (Fig. 5a), efficacies are generally higher, which is likely due to a receptor reserve in the latter assay. Taken together, restricted capacity of the dualsteric probes for inducing inner loop rearrangement translates into diminished G-protein activation.

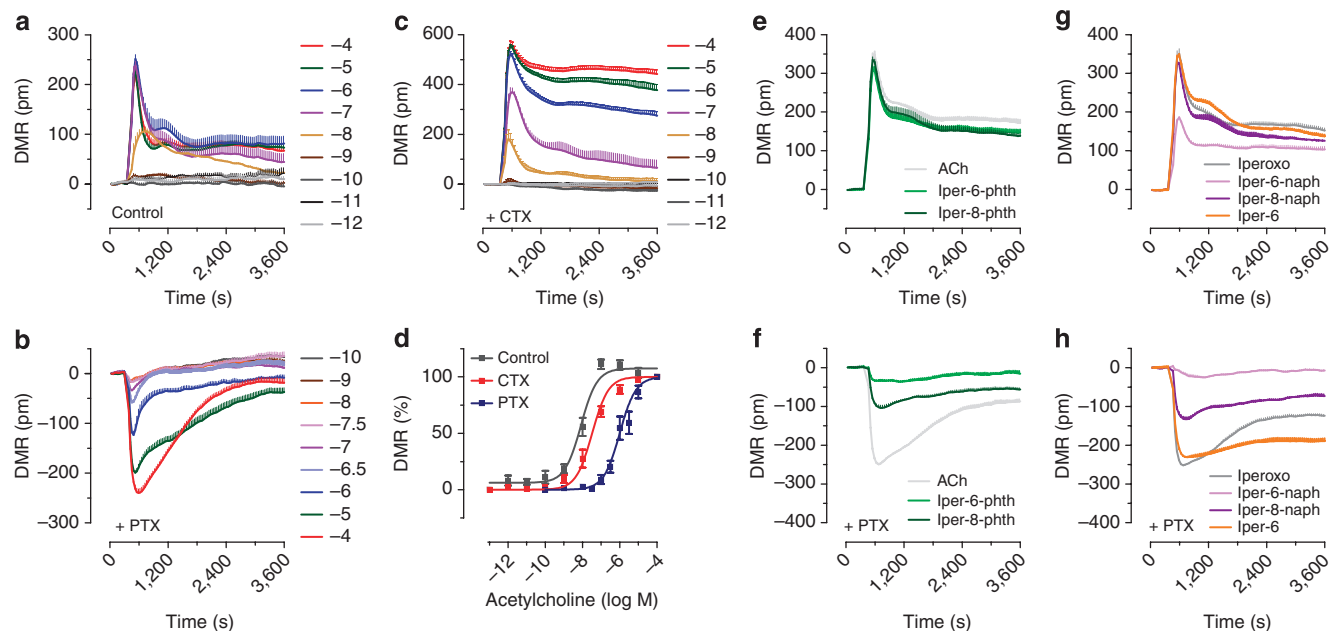


Figure 3 | Cellular DMR reveals structure-dependent G_i bias of dualsteric probes. (a–c) Shown are baseline-corrected DMR responses measured as wavelength shifts (pm) and recorded over time after addition of indicated concentrations of ACh in non-pretreated CHO-hM₂ cells (control, **a**) and either PTX (**b**)- or CTX (**c**)-pretreated CHO-hM₂ cells (means and s.e.m.) from one representative out of at least four independent experiments conducted in quadruplicates. (a) ACh exposure of control cells induces concentration-dependent positive DMR with peaks reflecting G_i signalling. (b) Pretreatment with the G_i inhibitor PTX (100 ng ml⁻¹, 16–24 h) reveals negative DMR reflecting G_s signalling. (c) Masking G_s -dependent signalling by CTX pretreatment (100 ng ml⁻¹, 8 h) induces positive DMR followed by sustained plateaus at higher concentrations. (d) Concentration-effect curves of ACh-induced peak DMR; data (means \pm s.e.m. from at least four experiments) are normalized to the response obtained with 100 μ M ACh. The pronounced sensitivity of ACh's potency to PTX indicates G_i dominance in the control DMR peaks. (e–h) DMR recordings of indicated compounds in either control cells (e,g) or PTX-pretreated cells (f,h). Applied were maximally effective concentrations: ACh 100 μ M, iper-6-phth 100 μ M, iper-8-phth 10 μ M, iperoxo 1 μ M, iper-6-naph 100 μ M, iper-8-naph 10 μ M and iper-6 100 μ M. Representative data (means and s.e.m.) of at least three independent experiments conducted in quadruplicates.

ERK1/2 phosphorylation is triggered by the $\beta\gamma$ -subunit of active G_i proteins⁴⁶. The active $\beta\gamma$ -subunit directly interacts with phosphoinositide 3-kinase γ inducing a series of downstream events resulting in increased ERK1/2-phosphorylation and eventually in modified gene expression⁴⁷. M₂ receptor-mediated ERK1/2-phosphorylation in CHO cells is exclusively G_i -dependent, as it is completely abolished after PTX pretreatment (Supplementary Fig. S3). As iper-6-X compounds are less efficacious than their middle chain-elongated iper-8-X fellows (Fig. 5c), the structure-dependent efficacy pattern is similar to that encountered in G_i readouts (Figs 3e,g and 4a).

β -Arrestin recruitment was studied in HEK293 cells stably expressing GFP- β -arrestin2 and transiently coexpressing the hM₂-Rluc fusion protein and GRK2 (Supplementary Methods). The construct has full signalling competence with respect to G_i and G_s signalling (Supplementary Fig. S4). β -Arrestin recruitment is partially restored in the middle chain-elongated derivatives iper-8-X relative to the hexamethylene compounds iper-6-X (Fig. 5d).

Taken together, dualsteric probe-induced reduction of the receptor's allosteric domain flexibility translates into restricted intracellular loop rearrangement and, subsequently, to impaired downstream signalling events.

An allosteric space 'rescue mutation' boosts signalling. In order to further test the hypothesis that M₂ receptor signalling critically depends on a conformational narrowing in the core region of the receptor's allosteric vestibule near M₂-W422^{7,35}, we replaced the spacious tryptophan by the small alanine. This should unleash

signalling of the dualsteric agonists. We focused on iper-6-phth and iper-6-naph, as these probes displayed the weakest signalling efficacy in the preceding experiments.

M₂-W422^{7,35}A reduces binding affinities of ACh (Fig. 6a) and iperoxo (Fig. 6b). The respective potencies for G-protein activation are diminished to the same extent (Fig. 6c,d). Both, iper-6-phth and iper-6-naph lose binding affinity and, to a lesser extent, potency at mutant receptors (Fig. 6a–d and Supplementary Table S5). Most importantly, the allosteric mutation leads to a gain in dualsteric probes' efficacy (Fig. 6c,d). As the surface expression of mutant receptors was about one-third lower compared with wild type (Supplementary Fig. S5), evaluation based on maximum effects would even underestimate the ligand's true capacity for receptor activation. Comparison of log τ -values reveals that efficacy for G_i activation of iper-6-phth and iper-6-naph in the mutant are indistinguishable from that of iperoxo (Fig. 6e).

Probe-induced maximum DMR was compared between CHO-hM₂ cells (Fig. 7a,b) and CHO-hM₂-W422^{7,35}A cells (Fig. 7b,c). Both, PTX pretreatment and the allosteric mutation lead to a pronounced loss of orthosteric agonist potency (Figs 3d and 6c,d). Under these conditions, iperoxo—due to its high potency—still achieves a clear maximum effect on the G_s pathway at the mutant receptor and was therefore used instead of ACh to define full efficacy and balanced agonism (for original DMR traces see Supplementary Fig. S6). As reported above (Fig. 6 and Supplementary Table S5), dualsteric probe potency is little affected by the mutation. Probes were applied at maximally effective concentrations with respect to both, G_s and G_i pathway activation. Under these conditions, maximum

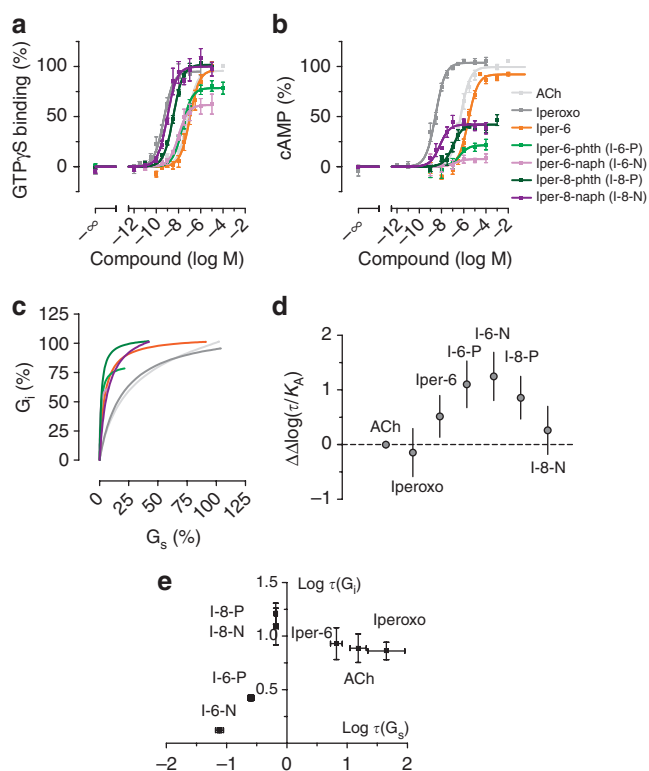


Figure 4 | Quantifying the G_i bias of dualsteric probes. (a) M₂ receptor-mediated G_i protein activation is reflected by [³⁵S]GTP γ S binding to membranes of CHO-hM₂ cells. Maximum G-protein activation induced by ACh was set 100%. Data are means \pm s.e.m. from at least four independent experiments conducted in quadruplicates. (b) M₂ receptor-mediated G_s activation is reflected by an increase of intracellular cAMP in CHO-hM₂ cells pretreated with PTX (50 ng ml⁻¹, 16–24 h). Maximum cAMP formation induced by ACh was set 100%. Data are means \pm s.e.m. from at least three independent experiments conducted in quadruplicates. Concentration-effect curves were obtained by global fits applying the operational model of agonism³⁷ to the data points in **a** and **b**. (c) Visualization of ligand bias: fractional [³⁵S]GTP γ S binding and cAMP accumulation (**a,b**) are plotted for equal concentrations of the respective agonists on the ordinate and abscissa, respectively. The hyperbolic shape of the ACh graph indicates system bias. The G_i protein is preferred by the M₂ receptor. Ligands whose bias exceeds system bias display graphs that differ substantially in their curvature radii and the height of their plateaus. (d) Quantification of ligand bias: $\Delta\Delta\log(\tau/K_A)$ values compensate for system bias and display true ligand bias relative to ACh, which is taken as balanced agonist. Ligands with positive values are G_i biased when the limits of their 95% confidence intervals (CIs) do not contain zero. Data are means with 95% CIs. (e) Efficacies of the agonist-occupied receptor to activate the G_i pathway ($\log \tau(G_i)$) and the G_s pathway ($\log \tau(G_s)$) are plotted on the ordinate and abscissa, respectively. Log τ values are derived from global fits applying the operational model to the data points in **a** and **b**. Data are means \pm s.e.m.

effects can be taken to reflect coupling efficacy τ . In the mutant receptor, maximum G_s-DMR induced by the probes is increased in case of the iper-6-X compounds and iper-8-phth (Fig. 7b). Only in the case of iper-8-naph, which has the highest G_s-DMR among the dualsteric probes at wild-type hM₂ receptor, a further increase in maximum G_s-DMR was not observed. The most impressive gain in G_s pathway activation is found with iper-8-phth, yet full G_s signalling is not achieved in the mutant (Fig. 7b,c). As mentioned previously (Fig. 6), the mutant receptor also allows for a gain in efficacy

Table 1 | Parameters characterizing ligand bias of muscarinic agonists.

Ligand	[³⁵ S]GTP γ S binding	cAMP accumulation	Log bias
	$\Delta\log(\tau/K_A)$, 95% CI	$\Delta\log(\tau/K_A)$, 95% CI	$\Delta\Delta\log(\tau/K_A)$, 95% CI
ACh	0	0	0
Iperoxo	2.11, 1.83 to 2.38	2.25, 1.84 to 2.66	-0.15, -0.59 to 0.29
Iper-6	-0.26, -0.51 to -0.01	-0.78, -1.09 to -0.46	0.52, 0.13 to 0.90
Iper-6-phth	0.37, 0.11 to 0.62	-0.74, -1.15 to -0.33	1.10, 0.68 to 1.53
Iper-6-naph	0.63, 0.35 to 0.91	-0.62, -1.03 to -0.21	1.25, 0.81 to 1.69
Iper-8-phth	1.21, 1.02 to 1.41	0.36, -0.05 to 0.77	0.86, 0.47 to 1.25
Iper-8-naph	1.78, 1.50 to 2.06	1.52, 1.11 to 1.93	0.26, -0.18 to 0.70

CI, confidence interval.

Transduction coefficients $\Delta\log(\tau/K_A)$ relative to the reference agonist ACh with 95% confidence intervals (95% CI) for the G_i and the G_s pathway as represented by [³⁵S]GTP γ S binding and cAMP accumulation assays, respectively. $\Delta\Delta\log(\tau/K_A)$ denotes true ligand bias.

for G_i signalling, which is most pronounced for iper-6-naph (see ordinates in Fig. 7a,c).

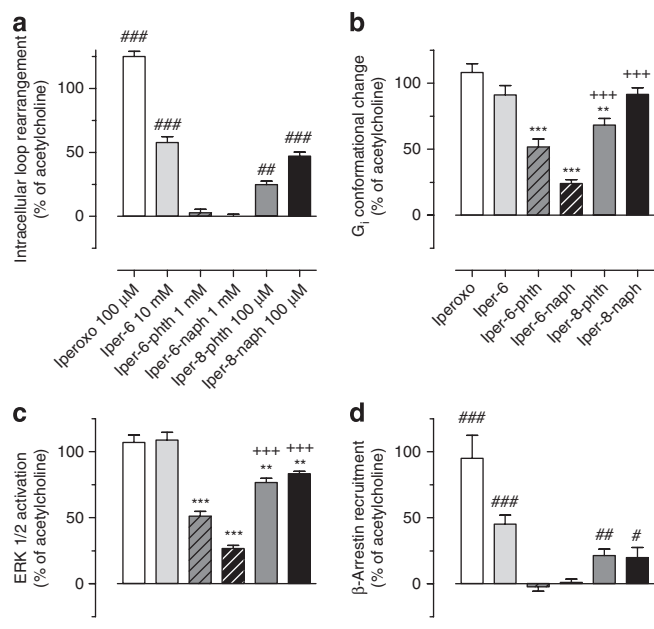
The partial agonists pilocarpine and McN-A-343 are included in Fig. 7 for comparison. Pilocarpine has an orthosteric binding topography²⁹, McN-A-343 is a bitopic allosteric/orthosteric partial agonist at hM₂ receptors⁴⁸. Pilocarpine and McN-A-343 respond differently to the mutation compared with the dualsteric probes (Fig. 7a,c), as the efficacy of pilocarpine is almost completely lost, whereas the efficacy of McN-A-343 does not differ from hM₂ wild type (all E_{max} values are enumerated in Supplementary Table S6). These findings underpin that the dualsteric probes exhibit a unique molecular mechanism of receptor interaction.

Importantly, the increase in G_i/G_s signalling of the dualsteric compounds in CHO-hM₂-W422^{7,35}A cells (Fig. 7c) strictly follows the hyperbolic relationship defined for CHO-hM₂ cells (Fig. 7a). Obviously, nearly full efficacy for G_i signalling is required before additional G_s signalling can emerge. We conclude that the receptor conformation mediating G_s signalling is reached via an intermediate conformation, the signalling efficacy of which is confined to G_i activation.

Discussion

In recent years, accumulating evidence suggests that signalling promiscuity is a general principle of 7TMR biology. For many 7TMRs, ligands are available that selectively activate a subset of signalling pathways^{3,40}. The mechanisms underlying multiple signalling pathway activation and functional selectivity are far from being understood¹². Applying a new methodological approach, this study provides hitherto unprecedented insights into the role of the allosteric vestibule for the discrimination between signalling pathways of a 7TMR. In order to allow for the subtle conformational transitions that underlie signalling promiscuity and selectivity⁸, we laid emphasis on the study of a native, unlabelled receptor protein. To probe the role of the receptor's extracellular domain for discriminating signalling pathway activation, we applied small molecules designed to interfere in a graded manner with activation-related extracellular domain rearrangements. Our study shows that this strategy allows capturing conformational dynamics underlying differential signalling pathway activation.

Several studies show that extracellular loops, especially the second extracellular loop, take part in the binding and activation of



rhodopsin-like class A 7TMRs⁴⁹. Until now, however, the molecular events have not been elucidated by which the extracellular region is functionally linked with the intracellular control of adaptor protein recruitment and the consecutive differential activation of signalling pathways. Here we show for the first time that relay of signalling triggered by orthosteric agonist binding critically depends on spatial freedom for concurrent extracellular conformational transitions. Allosteric substituents—attached to the orthosteric agonist iperoxo—that protrude into the free volume of the extracellular loop region severely hamper signalling of the M₂ receptor.

Structure–activity relationships of dualsteric probes and employment of the hM₂-W422^{7.35A} mutant receptor allowed to narrow down a critical region of the M₂ receptor in the vicinity of W422^{7.35},

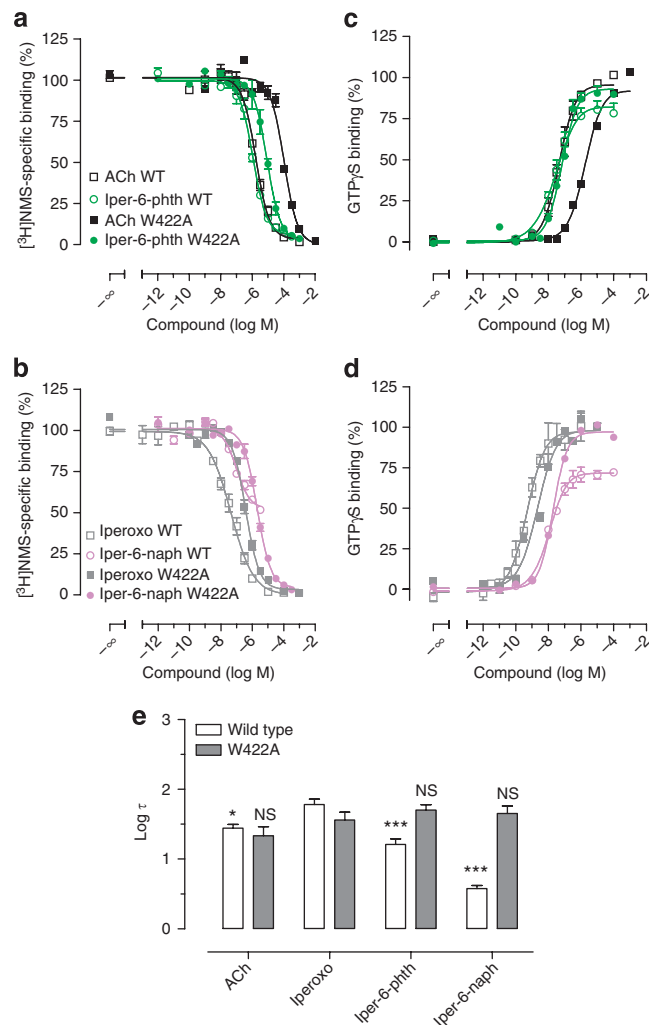


Figure 6 | **Recovery of G_i signalling of dualsteric probes in the allosteric M₂ W422^{7.35A} receptor mutant.** (a, b) Equilibrium cell surface receptor binding of ACh, iperoxo, iper-6-phth and iper-6-naph competing against [³H]NMS in live CHO-hM₂ cells (open symbols) and CHO-hM₂ W422^{7.35A} cells (filled symbols). Data are means ± s.e.m. from at least three independent experiments conducted in triplicates. Binding curve fitting was based either on the four-parameter logistic function or on the allosteric ternary complex model in case of iper-6-phth wild type (WT) and iper-6-naph WT. Binding constants for the M₂ WT receptor and M₂ W422A mutant receptor were 5.86 ± 0.04 and 4.36 ± 0.02 for ACh, 7.49 ± 0.16 and 6.81 ± 0.01 for iperoxo, 6.11 ± 0.06 and 5.45 ± 0.09 for iper-6-phth and 7.07 ± 0.08 and 6.05 ± 0.04 for iper-6-naph, respectively.

(c, d) M₂ receptor-mediated G_i protein activation was measured as reflected by [³⁵S]GTPγS binding to membranes of CHO-hM₂ cells (open symbols) and CHO-hM₂ W422^{7.35A} cells (filled symbols). Maximum G-protein activation induced by ACh was set 100%. Data are means ± s.e.m. from at least three independent experiments conducted in quadruplicates. Curve fitting in c and d was based on the operational model of agonism using the ligand-binding affinities derived from a and b.

(e) Coupling efficiencies of indicated ligands at the M₂ WT receptor and the M₂ W422^{7.35A} receptor mutant. (*****P* < 0.001, **P* < 0.05), significantly different from iperoxo at the M₂ WT receptor according to two-way ANOVA with Bonferroni's multiple comparison test, NS, not significantly different from iperoxo at the mutant receptors.

which is located above the orthosteric-binding pocket at the contact between the third extracellular loop and the beginning of TM7 (ref. 25). This epitope is conserved among the five subtypes of the

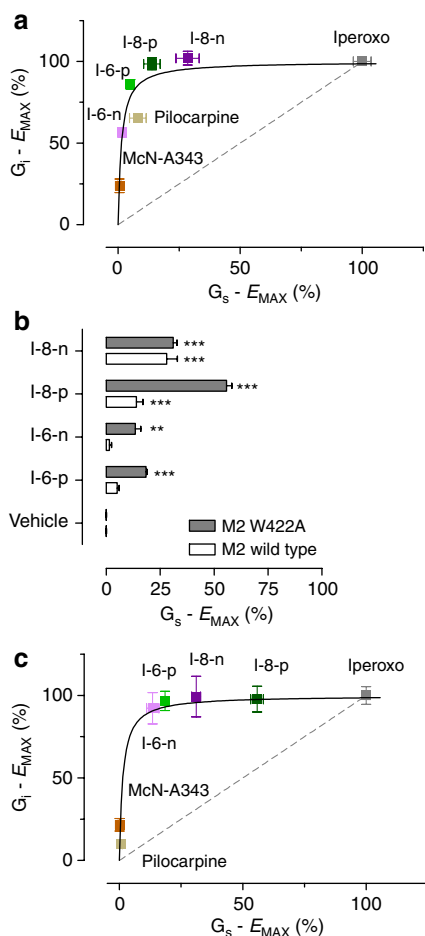


Figure 7 | G_s signalling of dualsteric probes is augmented in the W422^{7.35}A mutant. (a,c) Maximum G_i - and G_s -DMR effects of the indicated compounds in wild-type CHO-hM₂ cells (a) or CHO-hM₂ W422^{7.35}A cells (c). Data represent means \pm s.e.m. from three independent experiments in a and c. Cells were pretreated with PTX 100 ng ml⁻¹ for 16–24 h to disclose G_s signalling. All concentrations used provide maximum effects for both the G_i and the G_s pathway according to GTP γ S binding and cAMP accumulation assays, respectively. Data were normalized on the respective effect of iperoxo 10 μ M, which was set 100% for each condition. Dashed lines represent balanced agonism. Continuous lines define a binodal curve resulting from a best fit to the indicated data points in a. Concentrations of ligands: iperoxo 10 μ M, iper-6-phth (i-6-p) 100 μ M, iper-8-phth (i-8-p) 10 μ M, iper-6-naph (i-6-n) 100 μ M, iper-8-naph (i-8-n) 10 μ M, pilocarpine 100 μ M and McN-A-343 100 μ M. (b) Iper-6-phth and iper-6-naph induce significant G_s signalling at the hM₂ W422^{7.35}A mutant in contrast to wild-type receptors. Shown are maximal DMR responses normalized to the DMR of iperoxo 10 μ M (set 100%) in CHO-hM₂ cells (wild type, open bars) and CHO-hM₂ W422^{7.35}A cells (filled bars). Data are means \pm s.e.m. of at least three independent experiments. * $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$, significant to vehicle according to a two-way ANOVA with Bonferroni's multiple comparison test.

muscarinic receptor. M₂-W422^{7.35} has been suggested to form a link with M₂-Y177^{5.32} of the second extracellular loop and to undergo a conformational rearrangement on receptor activation²⁹. Interestingly, comparison of the crystal structures of the inactive and the active β_2 receptor revealed that receptor activation involves a movement of the corresponding β_2 -Y308^{7.35} towards β_2 -F193^{5.32}

of the second extracellular loop, thereby narrowing the free volume above the orthosteric site¹⁰. Our strategy of probing conformational freedom including the 'rescue mutant' M₂ W422^{7.35}A demonstrates that the corresponding region of the M₂ receptor is critical for G-protein coupling.

The physiological function of allosteric binding areas of class A 7TMR is only beginning to be understood. Evidence for a mechanism of general importance has recently been provided for β_1 and β_2 adrenergic receptors. Molecular dynamics simulations imply that ligand entry into and exit from the orthosteric site is a two-step event that includes intermediate ligand binding to a vestibule formed in the receptor's extracellular domains^{50,51}. The first step, entry into the vestibule, appears to be important for the removal of water molecules from the ligand⁵⁰. The subsequent passage leading from the allosteric vestibule into the orthosteric-binding cavity of the β_2 receptor is lined by β_2 -Y308^{7.35} and β_2 -F193^{5.32} (ref. 50). As these epitopes are rearranged in the course of the inactive-to-active receptor transition¹⁰, it is likely that this rearrangement occurs parallel to or is even caused by the ligand moving along the passage into the orthosteric site. Our findings disclose that spatial rearrangement of this passage in the M₂ receptor is critical for receptor movements required for appropriate unfolding of the intracellular domain region for G-protein coupling. In particular, we show that the allosteric vestibule of the muscarinic M₂ receptor has to provide room for a 'standard' conformational rearrangement allowing G_i coupling and an 'extended' movement for G_s coupling. Beyond that we show that the G_s activation-competent receptor conformation is reached via an intermediate conformation with full efficacy for G_i activation. Hence, for the first time we assign a biological role to the allosteric vestibule for controlling G-protein signalling of a 7TMR.

Crystal structures of active class A 7TMRs point to a common activation mechanism including cytosolic outwards movement of transmembrane helices and a corresponding inwards movement of extracellular domains^{9–11}. In the end, the novel chemical-biological strategy introduced here offers a widely applicable label-free approach for clarifying the dependence of promiscuous signalling pathway activation on extracellular domain rearrangements. This does not only apply to 7TMRs that are already known to bind allosteric or even dualsteric ligands⁵². The new strategy will be of even broader relevance if the extracellular vestibule-mediated ligand-binding process represents a more general mechanism^{50,51}.

With respect to drug discovery, our findings likely disclose a general route for the design of a new type of biased agonist. It is unique as it addresses two distinct receptor sites simultaneously with opposing functional messages, that is activation of the orthosteric receptor region by the agonist moiety and inactivation of an allosteric domain by an antagonist or even inverse agonist moiety^{29,53}. When applied un-tethered, the components reveal strong negative binding cooperativity as previously shown with orthosteric ACh and allosteric naph- or phth-residues²⁹ reflecting a severe conformational mismatch. On covalent linkage of these building blocks, dualsteric binding is enforced leading to a receptor conformation, which is unlikely to occur with un-tethered ligands and has specific signalling properties. The choice of building blocks and linker-length allows designing custom-made receptor activators endowed with desired signalling phenotypes.

We conclude that the allosteric vestibule serves to fine-tune the nature of signals generated on orthosteric ligand activation of a 7TMR. The activation of differential signalling pathways piloted by extracellular domains is due to sequential conformational states of the receptor protein and eventually results in a hierarchical order of intracellular adaptor protein recruitment. As several family A 7TMRs harbour an allosteric vestibule within their outer domains, the findings also suggest that controlling G-protein coupling is a general biological role of 7TMR allosteric vestibules.

Methods

Cell culture. Flp-In-Chinese hamster ovary cells (Flp-In-CHO) stably expressing the hM₂ receptor (CHO-hM₂ cells) or the hM₂ W422A mutant receptor (CHO-hM₂ W422A cells) and CHO-K1 cells were cultured in Ham's nutrient mixture F-12 supplemented with 10% (v/v) fetal calf serum (FCS), 100 U ml⁻¹ penicillin, 100 µg ml⁻¹ streptomycin and 2 mM L-glutamine. COS-7 cells were grown in DMEM supplemented with 10% (v/v) FCS, 100 U ml⁻¹ penicillin and 100 µg ml⁻¹ streptomycin. HEK293 cell clones stably expressing either GFP2-β-arr2 or the hM₂ receptor FRET-sensor, respectively, were maintained in DMEM supplemented with 10% (v/v) FCS, 100 U ml⁻¹ penicillin, 100 µg ml⁻¹ streptomycin and 200–500 µg ml⁻¹ G418. All cell lines were kept at 37 °C in a humidified 5% CO₂ atmosphere (7% CO₂ for HEK293 cells stably expressing the hM₂ FRET-sensor).

Radioligand-binding assay. CHO-hM₂ and CHO-hM₂ W422A cells were seeded on cell culture dishes and grown for 2 days. For PTX experiments, cells were pre-treated with PTX (100 ng ml⁻¹) for 16–24 h before the assay. Cells were collected and adjusted to give a final concentration of 750,000 cells per ml in assay buffer (Hank's balanced salt solution (HBSS) supplemented with 20 mM HEPES; pH 7.0). All experiments were carried out with 75,000 cells per well in a 96-well microtitre-plate (Abgene, Germany) at 28 °C in a final volume of 500 µl. Experiments were terminated by rapid vacuum filtration and bound radioactivity was determined by solid scintillation counting.

For two-point kinetic dissociation experiments, cells were pre-incubated with 2 nM [³H]NMS for 45 min before addition of dualsteric compounds. Reassociation of [³H]NMS was prevented by 10 µM atropine and reactions were terminated after 10 min.

For equilibrium binding experiments, cells were incubated for 4 h with 0.2 nM [³H]NMS and agonists at different concentrations to reach equilibrium. To make sure that binding equilibrium was attained in presence of dualsteric compounds, incubation time was calculated as described previously²³.

DMR assay. DMR assays were performed on a beta version of the Corning Epic biosensor. A detailed protocol is published elsewhere³³. Briefly, CHO-hM₂ wild-type, CHO-hM₂W422A cells and CHO-K1 cells transfected to transiently express the hM₂ wild-type receptor or a FLAG-tagged hM₂ receptor-Rluc fusion protein were grown to confluence for 20–24 h on Epic biosensor 384-well microplates. Cells were washed twice with HBSS containing 20 mM HEPES (pH 7.0) and kept for 2 h in the Epic reader at 28 °C. DMR was monitored before (baseline read) and after the addition of compound solutions for 3,600 s. Pre-incubation time for CTX (100 ng ml⁻¹) pretreatment and PTX (100 ng ml⁻¹) pretreatment was 8 and 16–24 h, respectively.

[³⁵S]GTPγS-binding assay. [³⁵S]GTPγS-binding experiments were conducted as described previously^{24,29,32}. In brief, homogenates of membranes of CHO-hM₂ wild-type or CHO-hM₂W422A cells (40 µg ml⁻¹) were incubated with 0.07 nM [³⁵S]GTPγS and maximum agonist-induced [³⁵S]GTPγS incorporation was measured after 1 h.

cAMP accumulation assay. Agonist-induced rise of intracellular cAMP was captured in CHO-hM₂ cells pretreated with 50 ng ml⁻¹ PTX for 16–24 h as described previously⁵⁴ using the HTRF-cAMP dynamic kit (Cisbio, Bagnols-sur-Cèze, France) following the manufacturer's instructions. Fluorescence was quantified on a Mithras LB 940 reader (Berthold Technologies, Bad Wildbad, Germany).

ERK1/2 phosphorylation assay. Quantification of phosphorylated ERK1/2 levels was performed using the HTRF-Cellul'erk kit (Cisbio). CHO-hM₂ cells were seeded into 96-well plates at a density of 40,000 cells per well and cultured overnight. For PTX pretreatment, cells were dispensed in medium containing 100 ng ml⁻¹ PTX. The next day, cells were washed twice in HBSS buffer supplemented with 20 mM HEPES and starved for 4 h in serum-free medium. After starvation, cells were incubated with agonists diluted in medium for 10 min at 37 °C. Following treatment, agonists were removed and cells were lysed in 50 µl of the supplemented lysis buffer. The plates were incubated for 10 min at room temperature with shaking to lyse the cells and then frozen overnight at -20 °C. Lysates (16 µl) were transferred to a white 384-well plate. Anti-phospho-ERK1/2-d2 (2 µl) and anti-ERK1/2-Eu³⁺-Cryptate (2 µl) were added to each well and plates were incubated in the dark for 2 h at room temperature. Time-resolved FRET signals were measured after excitation at 320 nm using the Mithras LB 940 multi-mode reader (Berthold Technologies). Data analysis was based on the fluorescence ratio emitted by the d2-labelled anti-phospho-ERK1/2 (665 nm) over the light emitted by the Eu³⁺-Cryptate-labelled anti-ERK1/2 (620 nm). Levels of phosphorylated ERK1/2 were normalized to those obtained by 100 µM ACh.

BRET assay. β-Arrestin2 recruitment assays were performed on HEK293 cells stably expressing GFP2-β-arr2 and transiently coexpressing the human M₂ receptor-Rluc fusion protein and GRK2. For measurement of G-protein subunit rearrangements, COS-7 cells transiently coexpressing the hM₂ receptor, Gα_{i1}-91Rluc, GFP10-Gγ₂ along with the complementary subunit Gβ₁ were used. Forty-eight hours after transfection, cells were detached and resuspended in

HBSS with 20 mM HEPES at a density of 1.06×10⁶ cells per ml. Cell suspension (170 µl) was distributed in white 96-well microplates and incubated in the presence of 10 µl buffer or agonist for 2 min (β-arrestin2 recruitment) or 1 min (G-protein activation) before substrate addition. DeepBlueC coelenterazine (Gold Biotechnology, MO, USA; 20 µl per well) was injected by injector 3 to yield a final concentration of 5 µM. To detect BRET, light emission at 400 and 515 nm was measured sequentially using a Mithras LB 940 instrument. The BRET signal (milliBRET ratio) was determined by calculating the ratio of the light emitted by the fluorescence acceptor (515 nm) and the light emitted by Rluc (400 nm).

FlAsH labelling and single cell-FRET measurements. FlAsH labelling was performed as published previously⁴³. In brief, cells were grown to near confluence on glass coverslips. Cells were washed twice with labelling buffer containing 150 mM NaCl, 10 mM HEPES, 2.5 mM KCl, 4 mM CaCl₂, 2 mM MgCl₂ supplemented with 10 mM glucose (pH 7.3). After incubation with labelling buffer supplemented with 500 nM FlAsH and 12.5 µM 1,2-ethanedithiol (EDT) at 37 °C for 1 h, cells were flushed twice with labelling buffer. To reduce non-specific binding, the cells were again incubated for 10 min with labelling buffer supplemented with 250 µM EDT. Finally, the cells were rinsed twice with labelling buffer.

FRET measurements were performed using a Zeiss Axiovert 200 inverted microscope endowed with a PLAN-Neofluar oil immersion 100-objective, a dual emission photometric system and a Polychrome IV light source (Tiil Photonica, Gräfelfing, Germany) as described previously^{42,44}. Experiments were conducted at room temperature using live HEK293 cells stably expressing the hM₂ receptor FRET-sensor that were maintained in assay buffer. Single cells were excited at 436 nm (dicroic 460 nm) with a frequency of 10 Hz. Emitted light was recorded using 535/30 nm and 480/40 nm emission filters and a DCLP 505 nm beam splitter for FlAsH and CFP, respectively. FRET was observed as the ratio of FlAsH/CFP, which was corrected offline for bleedthrough, direct FlAsH excitation and photo-bleaching using the 8.0 version of the Origin software as described recently⁴⁴. To investigate changes in FRET on ligand addition, cells were continuously superfused with FRET buffer complemented with various ligands in saturating concentrations as indicated. Superfusion was done using the ALA-VM8 (ALA Scientific Instruments).

Calculations and data analysis. Quantification of probe-induced DMR signals was based on the signal's peak value between 500 and 1,200 s. All DMR recordings were solvent corrected.

All nonlinear regression analyses were performed using Prism 5.01 (GraphPad Software, San Diego, CA). Equilibrium binding data from ACh, iperoxo, iper-6 and iper-8-phth were analysed by a four-parameter logistic function yielding IC₅₀ values, which were subsequently converted into apparent equilibrium dissociation constants K_i using the Cheng-Prusoff correction as described previously²⁴. In case of allosteric incomplete displacement of [³H]NMS by iper-6-phth, iper-6-naph and iper-8-naph, curve analysis was based on the allosteric ternary complex model as described elsewhere^{24,29}. Data obtained from [³⁵S]GTPγS-binding and cAMP accumulation assays were subjected to a two-step analysis. First, data were fitted by a four-parameter logistic function yielding parameter values for a ligand's potency (pEC₅₀) and maximum effect (E_{max}). Second, data points were analysed by the operational model of agonism³⁷,

$$E = \frac{E_{\max} \cdot \tau \cdot [A]}{\tau \cdot [A] + ([A] + K_A)} \quad (1)$$

where E_{max} is the maximal response of the system induced by the maximal effect of a full agonist, K_A is the equilibrium dissociation constant of the agonist-receptor complex and reflects affinity, and τ is the 'coupling efficiency', which indicates how efficient agonist binding is transduced into a signalling response and therefore reflects agonist efficacy. For quantification of ligand bias, we used a scale log(τ/K_A) that comprises information about both affinity and efficacy of the respective agonist for activating a signalling pathway³⁸. These transduction coefficients were then expressed relative to ACh for each pathway according to

$$\Delta \log(\tau / K_A)_{\text{agonist, pathway}} = \log(\tau / K_A)_{\text{agonist, pathway}} - \log(\tau / K_A)_{\text{ACh, pathway}} \quad (2)$$

To offset the effects of system bias and to display true ligand bias, a 'Log bias' scale is given by

$$\Delta \Delta \log(\tau / K_A)_{\text{agonist}} = \Delta \log(\tau / K_A)_{\text{agonist, pathway 1}} - \Delta \log(\tau / K_A)_{\text{agonist, pathway 2}} \quad (3)$$

Computation of transduction coefficients, ratios thereof and the bias factors included error propagation.

Statistical analysis. Data are shown as means \pm s.e.m for n observations. Comparisons of groups were performed using one-way ANOVA or two-way ANOVA with Bonferroni's multiple comparison test.

References

- Overington, J. P., Al-Lazikani, B. & Hopkins, A. L. How many drug targets are there? *Nat. Rev. Drug Discov.* **5**, 993–996 (2006).
- Lagerstrom, M. C. & Schiöth, H. B. Structural diversity of G protein-coupled receptors and significance for drug discovery. *Nat. Rev. Drug Discov.* **7**, 339–357 (2008).
- DeWire, S. M., Ahn, S., Lefkowitz, R. J. & Shenoy, S. K. β -Arrestins and cell signaling. *Annu. Rev. Physiol.* **69**, 483–510 (2007).
- Simon, M. I., Strathmann, M. P. & Gautam, N. Diversity of G proteins in signal transduction. *Science* **252**, 802–808 (1991).
- Kenakin, T. & Miller, L. J. Seven transmembrane receptors as shapeshifting proteins: the impact of allosteric modulation and functional selectivity on new drug discovery. *Pharmacol. Rev.* **62**, 265–304 (2010).
- Rajagopal, S., Rajagopal, K. & Lefkowitz, R. J. Teaching old receptors new tricks: biasing seven-transmembrane receptors. *Nat. Rev. Drug Discov.* **9**, 373–386 (2010).
- Smith, N. J., Bennett, K. A. & Milligan, G. When simple agonism is not enough: emerging modalities of GPCR ligands. Signalling and regulation of G protein coupled receptors. *Mol. Cell. Endocrinol.* **331**, 241–247 (2011).
- Deupi, X. & Kobilka, B. K. Energy landscapes as a tool to integrate GPCR structure, dynamics, and function. *Physiology (Bethesda)* **25**, 293–303 (2010).
- Scheerer, P. *et al.* Crystal structure of opsin in its G-protein-interacting conformation. *Nature* **455**, 497–502 (2008).
- Rasmussen, S. G. F. *et al.* Structure of a nanobody-stabilized active state of the β_2 adrenoceptor. *Nature* **469**, 175–180 (2011).
- Lebon, G. *et al.* Agonist-bound adenosine A2A receptor structures reveal common features of GPCR activation. *Nature* **474**, 521–525 (2011).
- Reiter, E., Ahn, S., Shukla, A. K. & Lefkowitz, R. J. Molecular mechanism of β -arrestin-biased agonism at seven-transmembrane receptors. *Annu. Rev. Pharmacol. Toxicol.* **52**, 179–197 (2012).
- Rasmussen, S. G. F. *et al.* Crystal structure of the β_2 adrenergic receptor-Gs protein complex. *Nature* **477**, 549–555 (2011).
- Oldham, W. M. & Hamm, H. E. Heterotrimeric G protein activation by G-protein-coupled receptors. *Nat. Rev. Mol. Cell. Biol.* **9**, 60–71 (2008).
- Bokoch, M. P. *et al.* Ligand-specific regulation of the extracellular surface of a G-protein-coupled receptor. *Nature* **463**, 108–112 (2010).
- Christopoulos, A. & Kenakin, T. G protein-coupled receptor allostery and complexing. *Pharmacol. Rev.* **54**, 323–374 (2002).
- Wess, J., Eglén, R. M. & Gautam, D. Muscarinic acetylcholine receptors: mutant mice provide new insights for drug development. *Nat. Rev. Drug Discov.* **6**, 721–733 (2007).
- Lüllmann, H., Ohnesorge, F. K., Schauwecker, G. C. & Wassermann, O. R. Inhibition of the actions of carbachol and DFP on guinea pig isolated atria by alkane-bis-ammonium compounds. *Eur. J. Pharmacol.* **6**, 241–247 (1969).
- Stockton, J. M., Birdsall, N. J., Burgen, A. S. & Hulme, E. C. Modification of the binding properties of muscarinic receptors by gallamine. *Mol. Pharmacol.* **23**, 551–557 (1982).
- Tränkle, C. *et al.* Interactions of orthosteric and allosteric ligands with [3 H]dimethyl-W84 at the common allosteric site of muscarinic M2 receptors. *Mol. Pharmacol.* **64**, 180–190 (2003).
- Voigtländer, U. *et al.* Allosteric site on muscarinic acetylcholine receptors: identification of two amino acids in the muscarinic M2 receptor that account entirely for the M2/M5 subtype selectivities of some structurally diverse allosteric ligands in N-methylscopolamine-occupied receptors. *Mol. Pharmacol.* **64**, 21–31 (2003).
- Huang, X. -P., Prilla, S., Mohr, K. & Ellis, J. Critical amino acid residues of the common allosteric site on the M2 muscarinic acetylcholine receptor: more similarities than differences between the structurally divergent agents gallamine and bis(ammonio)alkane-type hexamethylene-bis-[dimethyl-(3-hydroxypropyl)ammonium]dibromide. *Mol. Pharmacol.* **68**, 769–778 (2005).
- Prilla, S., Schrobang, J., Ellis, J., Hölte, H. -D. & Mohr, K. Allosteric interactions with muscarinic acetylcholine receptors: complex role of the conserved tryptophan M₂ 422Trp in a critical cluster of amino acids for baseline affinity, subtype selectivity, and cooperativity. *Mol. Pharmacol.* **70**, 181–193 (2006).
- Antony, J. *et al.* Dualsteric GPCR targeting: a novel route to binding and signaling pathway selectivity. *FASEB J.* **23**, 442–450 (2009).
- Haga, K. *et al.* Structure of the human M2 muscarinic acetylcholine receptor bound to an antagonist. *Nature* **482**, 547–551 (2012).
- Hulme, E. C., Lu, Z. L. & Bee, M. S. Scanning mutagenesis studies of the M1 muscarinic acetylcholine receptor. *Receptors Channels* **9**, 215–228 (2003).
- Shi, L. & Javitch, J. A. The binding site of aminergic G protein-coupled receptors: the transmembrane segments and second extracellular loop. *Annu. Rev. Pharmacol. Toxicol.* **42**, 437–467 (2002).
- Kloeckner, J., Schmitz, J. & Holzgrabe, U. Convergent, short synthesis of the muscarinic superagonist iperoxo. *Tetrahedron Lett.* **51**, 3470–3472 (2010).
- Jäger, D. *et al.* Allosteric small molecules unveil a role of an extracellular E2/transmembrane helix 7 junction for G protein-coupled receptor activation. *J. Biol. Chem.* **282**, 34968–34976 (2007).
- Haga, K. *et al.* Functional reconstitution of purified muscarinic receptors and inhibitory guanine nucleotide regulatory protein. *Nature* **316**, 731–733 (1985).
- Michal, P., Lysíková, M. & Tuček, S. Dual effects of muscarinic M(2) acetylcholine receptors on the synthesis of cyclic AMP in CHO cells: dependence on time, receptor density and receptor agonists. *Br. J. Pharmacol.* **132**, 1217–1228 (2001).
- Schröder, R. *et al.* Deconvolution of complex G protein-coupled receptor signaling in live cells using dynamic mass redistribution measurements. *Nat. Biotech.* **28**, 943–949 (2010).
- Schröder, R. *et al.* Applying label-free dynamic mass redistribution technology to frame signaling of G protein-coupled receptors noninvasively in living cells. *Nat. Protoc.* **6**, 1748–1760 (2011).
- Lazareno, S. & Birdsall, N. J. Detection, quantitation, and verification of allosteric interactions of agents with labeled and unlabeled ligands at G protein-coupled receptors: interactions of strychnine and acetylcholine at muscarinic receptors. *Mol. Pharmacol.* **48**, 362–378 (1995).
- Milligan, G. Principles: extending the utility of [35 S]GTP γ S binding assays. *Trends Pharmacol. Sci.* **24**, 87–90 (2003).
- Gregory, K. J., Hall, N. E., Tobin, A. B., Sexton, P. M. & Christopoulos, A. Identification of orthosteric and allosteric site mutations in M₂ muscarinic acetylcholine receptors that contribute to ligand-selective signaling bias. *J. Biol. Chem.* **285**, 7459–7474 (2010).
- Black, J. W. & Leff, P. Operational models of pharmacological agonism. *Proc. R. Soc. Lond. B. Biol. Sci.* **220**, 141–162 (1983).
- Kenakin, T., Watson, C., Muniz-Medina, V., Christopoulos, A. & Novick, S. A simple method for quantifying functional selectivity and agonist bias. *ACS Chem. Neurosci.* **3**, 193–203 (2012).
- Galandrin, S., Oligny-Longpré, G. & Bouvier, M. The evasive nature of drug efficacy: implications for drug discovery. *Trends Pharmacol. Sci.* **28**, 423–430 (2007).
- Kenakin, T. Collateral efficacy in drug discovery: taking advantage of the good (allosteric) nature of 7TM receptors. *Trends Pharmacol. Sci.* **28**, 407–415 (2007).
- Maier-Puschel, M. *et al.* A fluorescence resonance energy transfer-based M2 muscarinic receptor sensor reveals rapid kinetics of allosteric modulation. *J. Biol. Chem.* **285**, 8793–8800 (2010).
- Hoffmann, C. *et al.* A FRET-based approach to determine G protein-coupled receptor activation in living cells. *Nat. Meth.* **2**, 171–176 (2005).
- Hoffmann, C. *et al.* Fluorescent labeling of tetracycline-tagged proteins in intact cells. *Nat. Protoc.* **5**, 1666–1677 (2010).
- Villardaga, J. -P., Bünnemann, M., Krasel, C., Castro, M. & Lohse, M. J. Measurement of the millisecond activation switch of G protein-coupled receptors in living cells. *Nat. Biotech.* **21**, 807–812 (2003).
- Gales, C. *et al.* Probing the activation-promoted structural rearrangements in preassembled receptor-G protein complexes. *Nat. Struct. Mol. Biol.* **13**, 778–786 (2006).
- Crespo, P., Xu, N., Simonds, W. F. & Gutkind, J. S. Ras-dependent activation of MAP kinase pathway mediated by G-protein beta gamma subunits. *Nature* **369**, 418–420 (1994).
- Lopez-Illasaca, M., Crespo, P., Pellici, P. G., Gutkind, J. S. & Wetzker, R. Linkage of G protein-coupled receptors to the MAPK signaling pathway through PI 3-kinase γ . *Science* **275**, 394–397 (1997).
- Valant, C. *et al.* A novel mechanism of G protein-coupled receptor functional selectivity. *J. Biol. Chem.* **283**, 29312–29321 (2008).
- Peeters, M. C., van Westen, G. J. P., Li, Q. & IJzerman, A. P. Importance of the extracellular loops in G protein-coupled receptors for ligand recognition and receptor activation. *Trends Pharmacol. Sci.* **32**, 35–42 (2011).
- Dror, R. O. *et al.* Pathway and mechanism of drug binding to G-protein-coupled receptors. *Proc. Natl Acad. Sci. USA* **108**, 13118–13123 (2011).
- González, A., Perez-Acle, T., Pardo, L. & Deupi, X. Molecular basis of ligand dissociation in β -adrenergic receptors. *PLoS ONE* **6**, e23815 (2011).
- Valant, C., Robert Lane, J., Sexton, P. M. & Christopoulos, A. The best of both worlds? Bitopic orthosteric/allosteric ligands of G protein-coupled receptors. *Annu. Rev. Pharmacol. Toxicol.* **52**, 153–178 (2012).
- Hilf, G. & Jakobs, K. H. Agonist-independent inhibition of G protein activation by muscarinic acetylcholine receptor antagonists in cardiac membranes. *Eur. J. Pharmacol.* **225**, 245–252 (1992).
- Schröder, R. *et al.* The C-terminal tail of CRTH2 is a key molecular determinant that constrains G α i and downstream signaling cascade activation. *J. Biol. Chem.* **284**, 1324–1336 (2009).
- May, L. T., Leach, K., Sexton, P. M. & Christopoulos, A. Allosteric modulation of G protein-coupled receptors. *Annu. Rev. Pharmacol. Toxicol.* **47**, 1–51 (2007).

Acknowledgements

We thank Corning Inc. for their support on the Epic system. This work was funded by the Deutsche Forschungsgemeinschaft (DFG) by grants to K.M. (MO 821/2-1), U.H. (HO 1368/12-1), E.K. (KO 1583/3-1) and C.H. (SFB487 TPA1). A.K. was a member of the graduate school GRK677. The University of Milan and 'Dote ricerca': FSE, Regione Lombardia financed the postdoctoral position of C.M. A.B. is member of the graduate school THEME (Theoretical and Experimental Medicine), University of Bonn. R.S. and L.P. are members of the graduate school BIGS DrugS, University of Bonn. R.S. is funded by the German National Merit Foundation.

Author contributions

A.B. designed and performed experiments and contributed to the design of research and to writing the manuscript. N.M., R.S., C.D., J.B., J.K., J.S., C.M., K.S., A.K., L.P. and A.M. designed and performed experiments. J.S.-L. edited the manuscript and contributed to discussion. C.T. contributed to data analysis, edited the manuscript and contributed to discussion. C.H., M.DeA. and U.H. designed research, contributed to

discussion and edited the manuscript. E.K. and K.M. designed research and wrote the manuscript.

Additional information

Supplementary Information accompanies this paper at <http://www.nature.com/naturecommunications>

Competing financial interests: The authors declare no competing financial interests.

Reprints and permission information is available online at <http://npg.nature.com/reprintsandpermissions/>

How to cite this article: Bock, A. *et al.* The allosteric vestibule of a seven transmembrane helical receptor controls G-protein coupling. *Nat. Commun.* 3:1044 doi: 10.1038/ncomms2028 (2012).

License: This work is licensed under a Creative Commons Attribution-NonCommercial-NoDerivative Works 3.0 Unported License. To view a copy of this license, visit <http://creativecommons.org/licenses/by-nc-nd/3.0/>