

# The structural basis for agonist and partial agonist action on a $\beta_1$ -adrenergic receptor

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$\beta$ -adrenergic receptors ( $\beta$ ARs) are G-protein-coupled receptors (GPCRs) that activate intracellular G proteins upon binding catecholamine agonist ligands such as adrenaline and noradrenaline<sup>1,2</sup>. Synthetic ligands have been developed that either activate or inhibit  $\beta$ ARs for the treatment of asthma, hypertension or cardiac dysfunction. These ligands are classified as either full agonists, partial agonists or antagonists, depending on whether the cellular response is similar to that of the native ligand, reduced or inhibited, respectively. However, the structural basis for these different ligand efficacies is unknown. Here we present four crystal structures of the thermostabilized turkey (*Meleagris gallopavo*)  $\beta_1$ -adrenergic receptor ( $\beta_1$ AR-m23) bound to the full agonists carmoterol and isoprenaline and the partial agonists salbutamol and dobutamine. In each case, agonist binding induces a 1 Å contraction of the catecholamine-binding pocket relative to the antagonist bound receptor. Full agonists can form hydrogen bonds with two conserved serine residues in transmembrane helix 5 (Ser<sup>5.42</sup> and Ser<sup>5.46</sup>), but partial agonists only interact with Ser<sup>5.42</sup> (superscripts refer to Ballesteros–Weinstein numbering<sup>3</sup>). The structures provide an understanding of the pharmacological differences between different ligand classes, illuminating how GPCRs function and providing a solid foundation for the structure-based design of novel ligands with predictable efficacies.

Determining how agonists and antagonists bind to the  $\beta$  receptors has been the goal of research for more than 20 years<sup>4–11</sup>. Although the structures of the homologous  $\beta_1$  and  $\beta_2$  receptors<sup>12–15</sup> show how some antagonists bind to receptors in the inactive state<sup>16</sup>, structures with agonists bound are required to understand subsequent structural transitions involved in activation. GPCRs exist in an equilibrium between an inactive state (R) and an activated state (R\*) that can couple and activate G proteins<sup>17</sup>. The binding of a full agonist, such as adrenaline or noradrenaline, is thought to increase the probability of the receptor converting to R\*, with a conformation similar to that of opsin<sup>18,19</sup>. In the absence of any ligand, the  $\beta$ ARs exhibit a low level of constitutive activity, indicating that there is always a small proportion of the receptor in the activated state, with the  $\beta_2$ AR showing a fivefold higher level of basal activity than the  $\beta_1$ AR<sup>20</sup>. Basal activity of  $\beta_2$ AR is important physiologically, as shown by the T164I<sup>4.56</sup> human polymorphism that reduces the basal activity of  $\beta_2$ AR to levels similar to  $\beta_1$ AR<sup>21</sup> and whose expression has been associated with heart disease<sup>22</sup>.

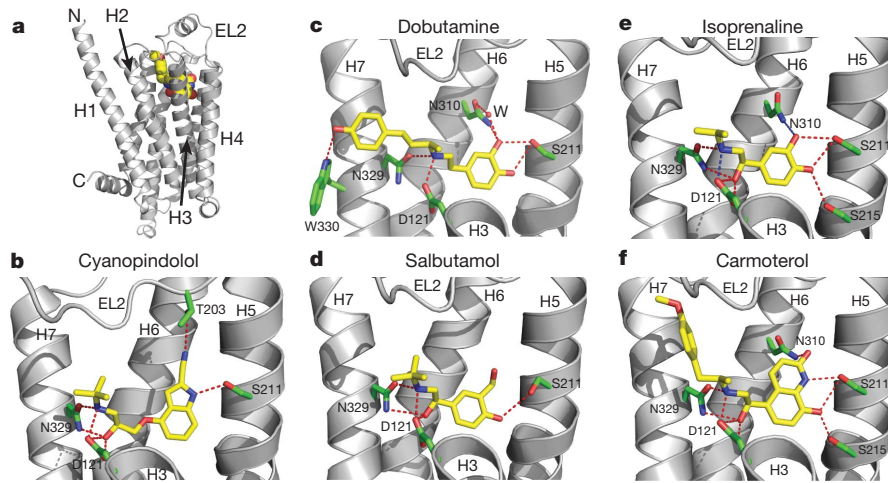
As a first step towards understanding how agonists activate receptors, we have determined the structures of  $\beta_1$ AR bound to four different agonists. Native turkey  $\beta_1$ AR is unstable in detergent<sup>23</sup>, so crystallization and structure determination relied on using a thermostabilized construct ( $\beta_1$ AR-m23) that contained six point mutations, which dramatically improved its thermostability<sup>24</sup>. In addition, the thermostabilizing mutations altered the equilibrium between R and R\*, so that the receptor was preferentially in the R state<sup>24</sup>. However, it could still couple to G proteins after activation by agonists<sup>13</sup> (Supplementary Fig. 1 and Supplementary Tables 1–3), although the activation energy barrier is predicted to be considerably higher than for the wild-type receptor<sup>25</sup>.

Here we report structures of  $\beta_1$ AR-m23 (see Methods) bound to R-isoprenaline (2.85 Å resolution), R,R-carmoterol (2.6 Å resolution), R-salbutamol (3.05 Å resolution) and R-dobutamine (two independent structures at 2.6 Å and 2.5 Å resolution) (Supplementary Table 5). The overall structures of  $\beta_1$ AR-m23 bound to the agonists are very similar to the structure with the bound antagonist cyanopindolol<sup>13</sup>, as expected for a receptor mutant stabilized preferentially in the R state. None of the structures show the outward movement of the cytoplasmic end of transmembrane helix H6 by 5–6 Å that is observed during light activation of rhodopsin<sup>18,19,26</sup>. This indicates that the structures represent an inactive, non-signalling state of the receptor formed on initial agonist binding.

All four agonists bind in the catecholamine pocket in a virtually identical fashion (Fig. 1). The secondary amine and  $\beta$ -hydroxyl groups shared by all the agonists (except for dobutamine, which lacks the  $\beta$ -hydroxyl; see Supplementary Fig. 4) form potential hydrogen bonds with Asp 121<sup>3.32</sup> and Asn 329<sup>7.39</sup>, whereas the hydrogen bond donor/acceptor group equivalent to the catecholamine *meta*-hydroxyl (*m*-OH) generally forms a hydrogen bond with Asn 310<sup>6.55</sup>. In addition, all the agonists can form a hydrogen bond with Ser 211<sup>5.42</sup>, as seen for cyanopindolol<sup>13</sup>, and they also induce the rotamer conformation change of Ser 212<sup>5.43</sup> so that it makes a hydrogen bond with Asn 310<sup>6.55</sup>. The major difference between the binding of full agonists compared to the partial agonists is that only full agonists make a hydrogen bond to the side chain of Ser 215<sup>5.46</sup> as a result of a change in side chain rotamer. All of these amino acid residues involved in the binding of the catecholamine head-groups to  $\beta_1$ AR are fully conserved in both  $\beta_1$  and  $\beta_2$  receptors (Fig. 2). Furthermore, the role of many of these amino acid residues in ligand binding is supported by extensive mutagenesis studies on  $\beta_2$ AR that were performed before the first  $\beta_2$ AR structure was determined<sup>27</sup>. Thus it was predicted that Asp 113<sup>3.32</sup>, Ser 203<sup>5.42</sup>, Ser 207<sup>5.46</sup>, Asn 293<sup>6.55</sup> and Asn 312<sup>7.39</sup> in  $\beta_2$ AR were all involved in agonist binding<sup>4,5,7–9</sup> (Fig. 3). Inspection of the region outside the catecholamine binding pocket in the structures with bound dobutamine and carmoterol allows the identification of non-conserved residues that interact with these ligands (Fig. 2 and Supplementary Fig. 7), which may contribute to the subtype specificity of these ligands<sup>10,28</sup>.

There are three significant differences in the  $\beta_1$ AR catecholamine binding pocket when full agonists are bound compared to when an antagonist is bound, namely the rotamer conformation changes of side chains Ser 212<sup>5.43</sup> and Ser 215<sup>5.46</sup> (Fig. 3) and the contraction of the catecholamine binding pocket by ~1 Å, as measured between the C $\alpha$  atoms of Asn 329<sup>7.39</sup> and Ser 211<sup>5.42</sup> (Fig. 4). So why should these small changes increase the probability of R\* formation? Agonist binding has not changed the conformation of transmembrane helix H5 below Ser 215<sup>5.46</sup>, although significant changes in this region are predicted once the receptor has reached the fully activated state<sup>18,19</sup>. The only effect that the agonist-induced rotamer conformation change of Ser 215<sup>5.46</sup> appears to have is to break the van der Waals interaction between Val 172<sup>4.56</sup> and Ser 215<sup>5.46</sup>, thus reducing the number of interactions

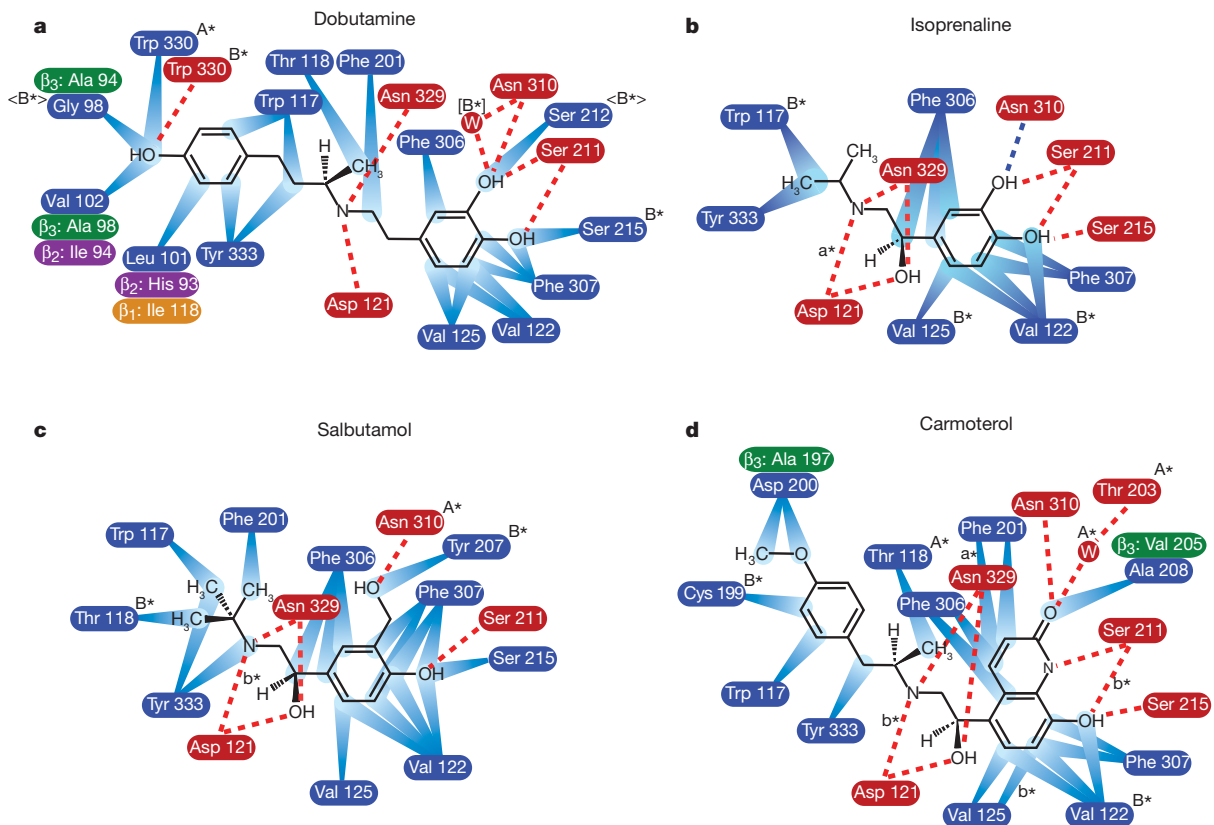
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**Figure 1 | Structure of the  $\beta_1$ -adrenergic receptor bound to agonists.**

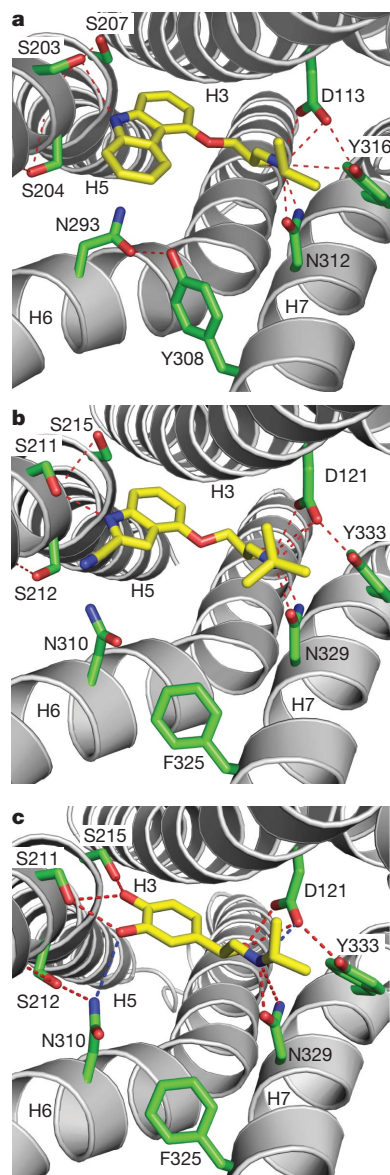
**a**, Structure of  $\beta_1$ AR shown in cartoon representation with the intracellular side at the bottom of the figure. The ligand carmoterol is shown as a space filling model (C, yellow; O, red; N, blue). The amino terminus (N), carboxy terminus (C), extracellular loop 2 (EL2), and transmembrane helices 1–4 (H1–H4) are labelled. **b–f**, The same orientation of receptor is shown in panels **b**, the antagonist cyanopindolol; **c**, **d**, the partial agonists dobutamine and salbutamol; **e**, **f**, the full agonists isoprenaline and carmoterol. The colour scheme of the

ligand and labelling of the receptor is identical in all panels, with amino acid side chains that make hydrogen bonds to the ligands depicted (C, green; O, red; N, blue). For clarity, residues 171–196 and 94–119 have been removed in **b–f**, which correspond to the C-terminal region of H4 and EL2, and EL1 with the C-terminal region of H2 and N-terminal region of H3, respectively. All structures shown are of monomer B (Supplementary Fig. 2) and were generated using Pymol (DeLano Scientific). For a comparison of the positions of the ligands when bound to the receptor, see Supplementary Fig. 5.



**Figure 2 | Polar and non-polar interactions involved in agonist binding to  $\beta_1$ -adrenergic receptor.** **a–d**, Amino acid residues within 3.9 Å of the ligands are depicted, with residues highlighted in blue making van der Waals contacts (blue rays) and residues highlighted in red making potential hydrogen bonds with favourable geometry (red dashed lines) or hydrogen bonds with unfavourable geometry (blue dashed lines). Amino acid residues labelled with an asterisk make the indicated contact either in monomer A (A\*) or in monomer B (B\*) only; for dobutamine, some contacts, labelled <B\*>, are

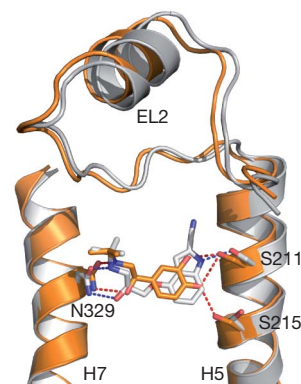
found only in monomer B of dobt92, whereas another contact, labelled [B\*], is found only in monomer B of dobt102 (Supplementary Fig. 6 and also see Supplementary Table 6 for further details and for the Ballesteros–Weinstein numbering). If specific van der Waals interactions or polar interactions are found only in monomer A or B, then the interaction is labelled a\* or b\*, respectively. Where the amino acid residue differs between the turkey  $\beta_1$ AR and the human  $\beta_1$ AR,  $\beta_2$ AR and  $\beta_3$ AR, the equivalent residue is shown highlighted in orange, purple or green, respectively (see also Supplementary Table 7).



**Figure 3 | Comparison of the ligand-binding pockets of the  $\beta_1$  and  $\beta_2$  adrenergic receptors.** The ligand-binding pockets are shown as viewed from the extracellular surface with EL2 removed for clarity (same colour scheme as in Fig. 1). **a**,  $\beta_2$ AR with the antagonist carazolol bound (PDB code 2RH1); **b**,  $\beta_1$ AR with the antagonist cyanopindolol bound (PDB code 2VT4); **c**,  $\beta_1$ AR with the agonist isoprenaline bound.

between H4 and H5. As there is only a minimal interface between transmembrane helices H4 and H5 in this region (Supplementary Table 8 and Supplementary Fig. 8), this loss of interaction may be significant in the activation process. In this regard, it is noteworthy that the naturally occurring polymorphism in  $\beta_2$ AR at the H4–H5 interface, T164I<sup>4,56</sup>, converts a polar residue to a hydrophobic residue as seen in  $\beta_1$ AR (Val 172<sup>4,56</sup>), which results in both reduced basal activity and reduced agonist stimulation<sup>21</sup>. This supports the hypothesis that the extent of interaction between H4 and H5 could affect the probability of a receptor transition into the activated state.

In contrast to the apparent weakening of helix–helix interactions by the agonist-induced rotamer conformation change of Ser 215<sup>5,46</sup>, the agonist-induced rotamer conformation change of Ser 212<sup>5,43</sup> probably results in the strengthening of interactions between H5 and H6. Upon agonist binding, Ser 212<sup>5,43</sup> forms a hydrogen bond with Asn 310<sup>6,55</sup> (Fig. 3) and, in addition, hydrogen bond interactions to Ser 211<sup>5,43</sup> and Asn 310<sup>6,55</sup> mediated by the ligand serve to bridge H5 and H6. The



**Figure 4 | Differences in the ligand-binding pocket between antagonist- and agonist-bound  $\beta_1$ -adrenergic receptor.** An alignment was performed (see Methods) between the structures of  $\beta_1$ AR-m23 bound to either cyanopindolol (grey) or isoprenaline (orange) and the relative positions of the ligands and the transmembrane helices H5 and H7 are depicted. The 1 Å contraction of the ligand-binding pocket between H5 and H7 is clear.

combined effects of strengthening the H5–H6 interface and weakening the H4–H5 interface could facilitate the subsequent movements of H5 and H6, as observed in the activation of rhodopsin.

Stabilization of the contracted catecholamine binding pocket is probably the most important role of bound agonists in the activation process (Fig. 4). This probably requires strong hydrogen bonding interactions between the catechol (or equivalent) moiety and both H5 and H6, and strong interactions between the secondary amine and  $\beta$ -hydroxyl groups in the agonist and the amino acid side chains in helices H3 and H7. Reduction in the strength of these interactions is likely to reduce the efficacy of a ligand<sup>29</sup>. Both salbutamol and dobutamine are partial agonists of  $\beta_1$ AR-m23 (Supplementary Table 3) and human  $\beta_1$ AR. In the case of salbutamol, there are only two predicted hydrogen bonds between the headgroup and H5/H6, compared to three–four potential hydrogen bonds for isoprenaline and carmoterol. Dobutamine lacks the  $\beta$ -hydroxyl group, which similarly reduces the number of potential hydrogen bonds to H3/H7 from three–four seen in the other agonists to only two. We propose that this weakening of agonist interactions with H5/H6 for salbutamol and H3/H7 for dobutamine is a major contributing factor in making these ligands partial agonists rather than full agonists.

The agonist-bound structures of  $\beta_1$ AR indicate there are three major determinants that dictate the efficacy of any ligand: ligand-induced rotamer conformational changes of (1) Ser 212<sup>5,43</sup> and (2) Ser 215<sup>5,46</sup> and (3) stabilization of the contracted ligand-binding pocket. The full agonists studied here achieve all three. The partial agonists studied here do not alter the conformation of Ser 215<sup>5,46</sup> and may be less successful than isoprenaline or carmoterol at stabilizing the contracted catecholamine binding pocket due to reduced numbers of hydrogen bonds between the ligand and the receptor. The antagonist cyanopindolol acts as a very weak partial agonist and none of the three agonist-induced changes are observed. In contrast to partial agonists, neutral antagonists or very weak partial agonists such as cyanopindolol may also have a reduced ability to contract the binding pocket owing to the greater distance between the secondary amine and the catechol moiety (or equivalent). For example, the number of atoms in the linker between the secondary amine and the headgroup of cyanopindolol is four, whereas the agonists in this study only contain two (Fig. 1 and Supplementary Fig. 4). A ligand with a sufficiently bulky headgroup that binds with high affinity and which actively prevents any spontaneous contraction of the binding pocket and/or Ser<sup>5,46</sup> rotamer change, would be predicted to act as a full inverse agonist. This is indeed what is observed in the recently determined structure<sup>15</sup> of  $\beta_2$ AR bound to the inverse agonist ICI 118,551.

The significant structural similarities amongst GPCRs suggests that similar agonist-induced conformational changes to those we have



observed here may also be applicable to many other members of the GPCR superfamily, though undoubtedly there will be many subtle variations on this theme.

## METHODS SUMMARY

**Expression, purification and crystallization.** The  $\beta_{44}$ -m23 construct was expressed in insect cells, purified in the detergent Hega-10 and crystallized in the presence of cholesterol hemisuccinate (CHS), following previously established protocols<sup>30</sup>. Crystals were grown by vapour diffusion, with the conditions shown in Supplementary Table 4.

**Data collection, structure solution and refinement.** Diffraction data were collected from a single cryo-cooled crystal (100 K) of each complex in multiple wedges at beamline ID23-2 at ESRF, Grenoble, France. The structures were solved by molecular replacement using the  $\beta_1$ AR structure<sup>13</sup> (PDB code 2VT4) as a model (see Methods). Data collection and refinement statistics are presented in Supplementary Table 5.

**Full Methods** and any associated references are available in the online version of the paper at [www.nature.com/nature](http://www.nature.com/nature).

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**Supplementary Information** is linked to the online version of the paper at [www.nature.com/nature](http://www.nature.com/nature).

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**Author Contributions** T.W. devised and performed receptor expression, purification, crystallization, cryo-cooling of the crystals, data collection and initial data processing. P.C.E. helped with crystal cryo-cooling and data collection. J.G.B. performed the pharmacological analyses on receptor mutants in whole cells and R.N. performed the ligand binding studies on baculovirus-expressed receptors. R.M. and A.G.W.L. were involved in data processing and structure refinement. Manuscript preparation was performed by T.W., C.G.T., A.G.W.L. and G.F.X.S. The overall project management was by G.F.X.S. and C.G.T.

**Author Information** Coordinates and structure factors have been submitted to the PDB database under accession codes 2y00, 2y01, 2y02, 2y03 and 2y04 for  $\beta_{44}$ -m23 bound either to dobutamine (dob92 and dob102), carmoterol, isoprenaline or salbutamol, respectively. Reprints and permissions information is available at [www.nature.com/reprints](http://www.nature.com/reprints). The authors declare no competing financial interests. Readers are welcome to comment on the online version of this article at [www.nature.com/nature](http://www.nature.com/nature). Correspondence and requests for materials should be addressed to C.G.T. ([cgt@mrc-lmb.cam.ac.uk](mailto:cgt@mrc-lmb.cam.ac.uk)) or G.F.X.S. ([gebhard.schertler@psi.ch](mailto:gebhard.schertler@psi.ch)).

## METHODS

**Expression, purification and crystallization.** The turkey (*M. gallopavo*)  $\beta_1$ AR construct,  $\beta_36$ -m23, contains six thermostabilizing point mutations and truncations at the N terminus, inner loop 3 and C terminus<sup>30</sup>. Here we used the  $\beta_44$ -m23 construct, which differs from the previously published  $\beta_36$ -m23 construct only by the presence of two previously deleted amino acid residues at the cytoplasmic end of helix 6 (H6), Thr 277 and Ser 278. Baculovirus expression and purification were all performed as described previously<sup>30</sup>, but with the detergent exchanged to Hega-10 (0.35%) on the alprenolol affinity column. Purified receptor was competitively eluted from the alprenolol Sepharose column with 0.2 mM agonist ((R)-isoprenaline, (R,S)-salbutamol, (R,S)-dobutamine or (R,R)-carmoterol). The buffer was exchanged to 10 mM Tris-HCl, pH 7.7, 100 mM NaCl, 0.1 mM EDTA, 0.35% Hega-10 and 1.0 mM agonist during concentration to 15–20 mg ml<sup>-1</sup>. Before crystallization, CHS and Hega-10 were added to 0.45–1.8 mg ml<sup>-1</sup> and 0.5–0.65%, respectively. Crystals were grown at 4 °C in 200 nl sitting drops and cryo-protected by soaking in either PEG 400 or PEG 600 for ~5 min (Supplementary Table 4) before mounting on Hampton CrystalCap HT loops and cryo-cooling in liquid nitrogen.

**Data collection, structure solution and refinement.** Diffraction data were collected at the European Synchrotron Radiation Facility, Grenoble, France, with a Mar 225 CCD detector on the microfocuss beamline ID23-2 (wavelength, 0.8726 Å) using a 10- $\mu$ m focused beam. The microfocuss beam was essential for the location of the best diffracting parts of single crystals, as well as allowing several wedges to be collected from different positions. Images were processed with MOSFLM<sup>31</sup> and SCALA<sup>32</sup>. The isoprenaline complex was solved by molecular replacement with PHASER<sup>33</sup> using the  $\beta_1$ AR structure (PDB code 2VT4) as a model<sup>13</sup>. This structure was then used as a starting model for the structure solution of the carmoterol complex. Finally, the carmoterol complex was used as a starting model for both the dobutamine complexes and for the salbutamol complex. Refinement and rebuilding were carried out with REFMAC5<sup>34</sup> and COOT<sup>35</sup> respectively. The dob92 dobutamine crystal diffracted to a higher resolution (2.5 Å) than the dob102 crystal (2.6 Å), but the dob102 data set was more complete and less anisotropic than dob92 and gave a lower Wilson B factor (Supplementary Table 5). Dictionary entries for the agonists were created using Jligand and PRODRG<sup>36</sup>. During refinement with REFMAC5 tight non-crystallographic restraints ( $\sigma = 0.05$  Å) were applied to the majority (172) of the residues in the two molecules in the asymmetric unit, with their selection based on improvements in  $R_{\text{free}}$  values. For the salbutamol complex, where the resolution was lower (3.05 Å), all three standard rotamers were modelled for Ser 211 and Ser 215 side chains, and the final choice was made on the basis of the local stereochemistry and features in the difference maps. Hydrogen bond assignments for the ligands were determined using hbplus<sup>37</sup> but allowing a maximum hydrogen-acceptor distance of 2.7 Å and a minimum angle of 89 degrees. Superposition of the different complexes was achieved by determining an initial transformation based on the 12 C-terminal residues of helix 2 (90–101) and then using the lsq\_imp option of the program O<sup>38</sup> to find the largest number of residues that could be superposed without a significant increase in the root mean squared deviation (r.m.s.d.). Cutoff values of between 0.2–0.5 Å for residues to be included in the superposition were found to produce the largest number of residues while maintaining a small r.m.s.d. (<0.15–0.3 Å), depending on the structures being compared. This was repeated using the uppermost residues of helices 3, 6 and 7 to determine the initial transformation, and all cases converged to give the same solution, with 147 residues superposed and a final r.m.s.d. of 0.28 Å for the superposition of the carmoterol and cyanopindolol structures, and lower r.m.s.d. values for superposing different agonist structures on one another. The convergence to a common solution validates this procedure for determining the optimal transformation. Validation of the final refined models was carried out using Molprobit<sup>39</sup>. Omit densities for the ligands and the surrounding side chains are shown in Supplementary Fig. 3.

The two dobutamine crystals (dob92 and dob102) differed in the crystallization buffer and pH (Supplementary Table 4) and this resulted in slightly different unit cell parameters (Supplementary Table 5) and packing arrangements. The differences between these two structures (overall r.m.s.d. 0.21 Å for monomer A, 0.21 Å for monomer B) provides a measure of the influence of crystal packing forces on the detailed conformation of the receptors. The observed differences in the ligand-binding pocket for monomer B, where there are no direct lattice contacts, emphasizes the conformational flexibility of this region (Supplementary Fig. 6).

**Pharmacological analysis of agonist binding to the thermostabilized  $\beta_1$ AR mutants in whole cells.** Stable CHO-K1 cell lines expressing either the wild-type turkey truncated receptor ( $\beta$ trunc), or the  $\beta_36$ , or the  $\beta_6$ -m23 or the  $\beta_36$ -m23 receptors and a CRE-SPAP reporter were used<sup>40</sup>. See Supplementary Table 1 for a description of the constructs. Cells were grown in Dulbecco's modified Eagle's medium nutrient mix F12 (DMEM/F12) containing 10% fetal calf serum and 2 mM D-glutamine in a 37 °C humidified 5% CO<sub>2</sub>; 95% air atmosphere.

To analyse the affinities of agonist binding to  $\beta_1$ AR mutants <sup>3</sup>H-CGP 12177 saturation binding and competition binding experiments were performed on whole cells (Supplementary Table 1). Cell lines were grown to confluence in white-sided tissue culture 96-well view plates. <sup>3</sup>H-CGP 12177 whole-cell competition binding was performed as previously described<sup>40</sup> using <sup>3</sup>H-CGP 12177 in the range of 0.82–1.80 nM. The  $K_D$  values for <sup>3</sup>H-CGP 12177 were 0.32 nM ( $\beta$ trunc), 0.85 nM ( $\beta_6$ -m23), 0.34 nM ( $\beta_36$ ) and 0.88 nM ( $\beta_36$ -m23). For the competition assays, all data points on each binding curve were performed in triplicate and each 96-well plate also contained six determinations of total and non-specific binding. In all cases, the competing ligand completely inhibited the specific binding of <sup>3</sup>H-CGP 12177. A one-site sigmoidal response curve was then fitted to the data using GraphPad Prism 2.01 and the IC<sub>50</sub> was then determined as the concentration required to inhibit 50% of the specific binding as described previously<sup>40</sup>.

The ability of the receptors to couple to G proteins and induce an increase in cAMP concentrations was determined by measuring the increase in secreted alkaline phosphatase (SPAP) under the transcriptional control of a cAMP response element (CRE). Cells were grown to confluence in clear plastic tissue culture treated 96-well plates and CRE-SPAP secretion into the media measured between 5 and 6 h after the addition of agonist as described previously (Supplementary Fig. 1 and Supplementary Table 3)<sup>40</sup>.

**Binding of agonists to  $\beta_1$ AR mutants expressed in insect cells for structural studies.** Receptors  $\beta_36$  and  $\beta_36$ -m23 were expressed using the baculovirus expression system in insect cells (High Five) as described previously<sup>30</sup>. Cells were disrupted by freeze-thaw and membranes prepared by centrifugation. Saturation binding and competition binding experiments were performed using <sup>3</sup>H-dihydroalprenolol as described previously<sup>41</sup>. Non-specific binding of radioligand to the receptor was determined by including 100  $\mu$ M unlabelled alprenolol. The assay mixtures were incubated for 2 h at 30 °C and then filtered on a 96-well glass-fibre filter plates (Millipore) pre-treated with polyethyleneimine. The filters were washed three times with ice-cold buffer (Tris 20 mM pH 8, NaCl 150 mM), dried, and counted in a Beckmann LS 6000 scintillation counter. The apparent IC<sub>50</sub> values were determined by nonlinear regression analysis using a one-site competition model in Prism software and  $K_i$  values were determined using the Cheng–Prusoff equation<sup>42</sup>.

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