Activation of CCR5 by Chemokines Involves an Aromatic Cluster between Transmembrane Helices 2 and 3*

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CCR5 is a G protein-coupled receptor responding to four natural agonists, the chemokines RANTES (regulated on activation normal T cell expressed and secreted), macrophage inflammatory protein (MIP)-1α, MIP-1β, and monocyte chemotactic protein (MCP)-2, and is the main co-receptor for the macrophage-tropic human immunodeficiency virus strains. We have previously identified a structural motif in the second transmembrane helix of CCR5, which plays a crucial role in the mechanism of receptor activation. We now report the specific role of aromatic residues in helices 2 and 3 of CCR5 in this mechanism. Using site-directed mutagenesis and molecular modeling in a combined approach, we demonstrate that a cluster of aromatic residues at the extracellular border of these two helices are involved in chemokine-induced activation. These aromatic residues are involved in interhelical interactions that are key for the conformation of the helices and govern the functional response to chemokines in a ligand-specific manner. We therefore suggest that transmembrane helices 2 and 3 contain important structural elements for the activation mechanism of chemokine receptors, and possibly other related receptors as well.

Chemokine receptors have been dragging more and more attention since the cloning of the first member of the family a decade ago. Not only are chemokines and their receptors now considered as the main organizers of leukocyte trafficking, they have also been associated to an ever increasing number of physiopathological disorders (for review, see Refs. 1 and 2). In particular, some chemokine receptors are used by the human immunodeficiency virus (HIV)1 as coreceptors (in addition to CD4) to infect target cells (reviewed in Ref. 3). Among these, CCR5 has been shown to be essential for HIV pathogenesis, as individuals homozygous for the CCR5Δ32 mutation, which results in the synthesis of a non-functional receptor, are highly (although not fully) resistant to HIV infection. Chemokine receptors belong to the rhodopsin-like family (family A) of G protein-coupled receptors (GPCR). CCR5 binds and responds to four natural chemokines, RANTES, MIP-1α, MIP-1β, and MCP-2, with nanomolar affinities (4).

Our current understanding of the activation mechanisms of GPCRs is rapidly evolving, thanks to the availability of the crystal structure of the inactive state of one of its members, rhodopsin, and to the growing amount of biochemical and physicochemical data that can help in identifying the key aspects of this process (for review, see Refs. 5 and 6). It is now well accepted that the transition from inactive to active states requires the reorganization of the transmembrane bundle made of seven imperfect α-helices. In the rhodopsin-like family, motions of transmembrane helix 3 (TM3) and TM6 during the activation process have been identified (7–9). Rigid-body movements have also been proposed for TM5 and TM7 (10, 11). Most of these conformational changes have been observed in a diverse set of receptors (mainly rhodopsin and the β2-adrenergic receptor), and it is believed that they constitute a common conformational path in the activation process, ultimately leading to the release of GDP in the bound G protein and its exchange for GTP.

A striking feature within the rhodopsin-like GPCR family is that, despite a strong sequence conservation of the transmembrane helices, there is a wide structural diversity among extracellular ligands, ranging from small neurotransmitters to large glycoproteins. The structural adaptation of a receptor to its
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1csw, 1.55 Å resolution), αα3 (1occ, 2.8 Å) and βαα3 (1ehk, 2.4 Å) cytochrome c oxidases, photosynthetic reaction center (1prr, 2.3 Å), potassium channel (1h98, 3.2 Å), mechanosensitive ion channel (1msl, 3.5 Å), rhodopsin (1f88, 2.8 Å), halorhodopsin (1el2, 1.8 Å), sensory rhodopsin (1h68, 2.1 Å), light harvesting complex (1lgh, 2.4 Å), photosystem 1 (1jbo, 2.5 Å), AQP1 (1hwo, 3.7 Å) and GlF (1f8x, 2.2 Å) channels, F-type ATPase (1eul, 2.6 Å), and fumarate reductase respiratory complex (1jbo, 2.2 Å) for α-helical segments featuring the FWXXY motif.

Molecular Dynamics Simulations of Transmembrane Helices—The model peptides Ace-Alaαα3-Thr-Ala-Pro-Alaαα3-Nme and Ace-Alaαα3-Thr-Gly-Alaαα3-Gly-Alaαα3-Ser-Gly-Alaαα3-Nme were built in the standard α-helical conformation (φ, ω = −58°, −47°). The amino acid side chains of Ser and Thr were set to the g+ conformation. Molecular dynamics trajectories of these model peptides aim to explore the role of TM2 in CCR5 triggered by the Thr-2.56-X-Pro-2.58 motif and TM3 triggered by the presence of Thr-3.29, Gly-3.30, Ser-3.35, Ser-3.38, and Gly-3.39. A similar approach was recently used to model the conformation of TM3 in the 5HT1A receptor (25). Ser and Thr residues induce a small bending angle in TM because of the additional hydrogen bond formed between the Oα atom of Ser or Thr and the i-3 or i-4 peptide carbonyl oxygen (26). Moreover, the additional flexibility provided by the adjacent Pro (because of the absence of the hydrogen bond with the carbonyl oxygen in the preceding turn of the helix) or Gly (because of the lack of the side chain) reinforces this effect. The obtained structures were placed in a rectangular box containing methane molecules to mimic the hydrophobic environment of the TM bundle. The peptide-methane systems were subjected to 500 iterations of energy minimization and then heated to 300 K in 15 ps. This was followed by an equilibration period (15–500 ps) and a production run (500–1500 ps) at constant volume using the particle mesh Ewald method to evaluate electrostatic interactions. Structures were collected for analysis every 10 ps during the production run (100 structures/simulation). To obtain a rough idea of the possible consequences that the presence of these residues in TM2 and TM3 might have on the structure of the receptor, we performed a molecular modeling exercise using the three-dimensional structure of rhodopsin as the template. The backbone of one helical turn preceding the highly conserved Asp-2.50 in TM2 and the FWXXY motif in TM3 was superimposed on the residues with the computed structures.

Molecular Dynamics Simulations of the CCR5 Receptor and Mutant Receptors—The three-dimensional model of transmembrane helices 1 and 4–7 of CCR5 was constructed by computer-aided model building techniques from the transmembrane domain of bovine rhodopsin, as determined by Palczewski et al. (27). The following conserved residues were employed in the alignment of rhodopsin and human CCR5 transmembrane sequences: Asn-55 (55 being the residue number in the 1f88 PDB file of rhodopsin) and Asn-48 (48 is the residue number in the CCR5 sequence, 1.50 in the standardized numbering); Trp-161 and Arg-165, Pro-215 and Pro-267, Pro-287 and Pro-250; Pro-303 and Pro-294. Representative structures of transmembrane helices 2 and 3, selected by automatically clustering the geometries obtained during the molecular dynamics trajectories into conformationally related subfamilies with the program NMRClust (28), were included into the model (see “Molecular Dynamics Simulations of Transmembrane Helices” above). SCWRL-2.1 was employed to add the side chains of the non-conserved residues based on a backbone-dependent rotamer library (29). All ionizable residues in the helices were considered uncharged with the exception of Asp-2.40, Asp-2.50, Asp-3.49, Arg-3.50, Lys-5.50, Arg-6.30, Arg-6.32, and Glu-7.39. To relieve residual strain resulting from suboptimal positioning of the side chains and the TM2-TM3 interface at the extracellular part, this resulting structure was placed in a rectangular box containing molecules, energy-minimized (500 steps), heated (from 0 to 300 K in 15 ps), and equilibrated (from 15 to 100 ps). During these processes, the Cα atoms were kept fixed at their positions in the rhodopsin crystal structure, with the exception of the residues forming the TM3-TM3 interface (from 2.58 to 2.59). The optimized TM2-TM3 interface accomplishes: (i) the (N′-term) of the TM2 helix is in the 10–11-Å range to allow the first extracellular loop (ECL1) of 4 residues to be shaped; (ii) Phe-82.29 interacts with Leu-104.29 and Tyr-89.62; (iii) with Thr-99.23, in a similar manner to the cytochrome c oxidase structure of helices III and VII in subunit III (see “Results”); and (iii) there are no steric clashes between the interactions of the Phe side chains of the TM2 helix and Tyr-104.29; (iv) Phe-82.29 and Thr-89.62 were further characterized by ab initio quantum mechanical calculations at the MP2/6–31G* level of theory, which is capable of describing the proposed C-H···π interactions (30).

Experimental Procedures

Numbering Scheme of GPCRs—In this work, we use a general numbering scheme to identify residues in the transmembrane segments of different receptors (24). Each residue is numbered according to the helix (1 through 7) in which it is located and to the position relative to the most conserved residue in that helix, arbitrarily assigned to 50. For instance Pro-2.58 is the proline in the transmembrane helix 2 (TM2), eight residues following the highly conserved aspartic acid Asp-2.50. For the sake of clarity, generalized numbers are italicized. When both numbers are given, the general numbering is put as superscript.

Survey of Transmembrane Helices Containing a FWXXY Motif in Known Membrane Protein Structures—We surveyed the atomic coordinates of the membrane proteins bacteriorhodopsin (PDB access number

In the case of chemokine receptors, we have recently identified a structural motif in TM2, which is central for chemokine-induced activation (13). This motif, which consists of a proline preceded by a threonine two residues ahead (T→P motif, is clearly identified a structural motif in TM2, which is central for chemokine-induced activation (13). This motif, which consists of a proline preceded by a threonine two residues ahead (T→P motif, is clearly involved in the activation process, as specific mutations of the motif led to unaffected chemokine binding but strong impairment of receptor activation (13). Moreover, modeling studies performed on this region suggested that, because of the action of the proline, the extracellular part of TM2 would strongly interact with TM3. This organization is structurally different from that of bovine rhodopsin, in which a TM2-TM1 interaction is found.

As a follow-up of these observations, we have now investigated the possible role of the TM2-TM3 interface in the activation process of chemokine receptors. A sequence alignment of chemokine receptors (Fig. 1) reveals that the extracellular parts of TM2 and TM3 contain many aromatic residues. Within the rhodopsin template, these residues are located at relatively short distances, suggesting that they might form an aromatic cluster within the three-dimensional structure of the receptors. Aromatic residues have been proposed to be involved in the activation mechanism in various GPCRs (14–22). Among other examples, a role was attributed to such residues in the ligand selectivity and ligand-induced activation of the D2 and D4 dopamine receptors (20, 23). The high density of aromatic residues at the top of TM2 and TM3 in chemokine receptors suggested that aromatic side chains could mediate interactions between these helices.

In the present study, we have mutated the aromatic residues of CCR5 TM2 and TM3 into their CCR2 counterparts, either individually or in combination, and the mutants were tested for cell-surface expression, receptor conformation, ligand binding, and functional response. Molecular modeling of the transmembrane region of CCR5 has been performed, providing a structural framework to interpret these data. Integration of the experimental and molecular modeling data indicates that aromatic residues at the TM2-TM3 interface are crucial to the mechanism of receptor activation and suggests that this aromatic cluster plays a key role in the conformational changes of CCR5, leading from ligand recognition to receptor activation.
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Molecular models of the helix bundles for the mutant receptors containing the F85*-2H, L104*-2F, F109*-2F/L104*-2F, F109*-2H, F112*-2Y, and F109*-2H/F112*-2Y substitutions were constructed from the previously obtained structure of CCR5, by changing the atoms implicated in the amino acid substitutions by interactive computer graphics. Subsequently, wild-type and mutant receptors were placed in a rectangular box (~71 Å × 60 Å × 50 Å in size) containing methane molecules (~2850 molecules in addition to the transmembrane domain) to mimic the hydrophobic environment of the transmembrane helices. The density of 0.4–0.5 g cm⁻³ of the methane box is approximately half of the density observed in the hydrophobic core of the membrane. This is a result of the different equilibrium distance between carbons in the methane box and in the polycarbon chain of the lipid. However, it has been shown that this procedure reproduces several important structural characteristics of membrane embedded proteins (31). The receptor-membrane systems were subjected to 500 iterations of energy minimization and then heated to 300 K in 15 ps. This was followed by an equilibration period (15–100 ps) and a production run (100–250 ps) at constant volume using the particle mesh Ewald method to evaluate electrostatic interactions (32). Structures were collected for analysis every 10 ps during the production run (15 structures per simulation).

The molecular dynamics simulations were run with the Sander module of AMBER 5 (33), the all-atom force field (34), SHAKE bond constraints in all bonds, a 2-fs integration time step, and constant temperature of 300 K coupled to a heat bath.

Plasmids encoding the CCR5 mutants studied here were constructed by site-directed mutagenesis using the QuikChange method (Stratagene). Following sequencing of the constructs, the mutated coding sequences were subcloned into the bicistronic expression vector pEFIN3 as previously described for generation of stable cell lines (35). All constructs were verified by sequencing prior to transfection.

**Expression of Mutant Receptors in CHO-K1 Cells**—CHO-K1 cells were cultured in Ham’s F-12 medium supplemented with 10% fetal calf serum (Invitrogen), 100 units/ml penicillin, and 100 μg/ml streptomycin (Invitrogen). Constructs encoding wild-type or mutant CCR5 were transfected into CHO-K1 cells in a final volume of 0.1 ml. The mixture contained 0.05 nM 125I-RANTES (2000 Ci/mmol, Amersham Biosciences) or 0.1 nM 125I-MIP-1α (0.1 mCi, Amersham Biosciences) and microplates were shaken for 1 min and further incubated at 30 °C for 30 min. The incubation was stopped by centrifugation of the microplate for 10 min, at 800 × g and 4 °C, and aspiration of the supernatant. Microplates were counted in a TopCount (Packard, Downers, IL) for 1 min/well. Functional parameters were determined with the PRISM software (GraphPad Software) using non-linear regression applied to a sigmoidal dose-response model.

**MAP Kinase Assay**—Cells, serum-starved for 24 h, were collected and assayed as described in serum-free DMEM. After 3 min of stimulation with various concentrations of RANTES and MCP-2, cells were centrifuged and heated to 100 °C for 5 min in lysis buffer (100 mM Tris-HCl, pH 6.8, 4 mM EDTA, 4% SDS, 20% glycerol, and 0.02% β-mercaptoethanol). For Western blot analysis, solubilized proteins corresponding to ~5 × 10⁶ cells were loaded onto 10% SDS-polyacrylamide gels in a Tricine buffer system (41). After transfer to nitrocellulose membranes, proteins were probed with mouse anti-phospho-p42/p44 (1:1000) (Cell Signal) or rabbit anti-total p38 (1:2000) antibodies (Santa Cruz).

**RESULTS**

To investigate the possible role of aromatic residues at the TM2-TM3 interface in defining the structure and function of the CCR5 receptor, we have compared the amino acid sequences of CCR2 and CCR5 (Fig. 1B). These receptors are strongly related, sharing ~85% sequence identity within their TM helices. However, their extracellular domains are much more divergent, which certainly contributes to their strong selectivity toward their respective ligands (35). The TM2-TM3 aromatic cluster was found to be quite divergent between the two receptors. This could suggest that these positions are not important for the structure and/or function of the receptors and therefore highly tolerant to variability. Alternatively, these positions could be functionally important although specific to each receptor, and the various substitutions would be expected in this case to be correlated. To study the functional consequences of the differences at aromatic positions observed between CCR5 and CCR2 (Fig. 1B), we engineered CCR5 mutants in which aromatic residues were substituted by the corresponding amino acids in CCR2. The F85*-2H and Y98*-2S mutants affected TM2, L104*-2F, and F112*-3Y involved TM3. We also combined these point mutations either within TM2 (F85L/Y98S double mutant), within TM3 (F109H/L112Y/F112H) or across both helices (F85L/L104F, Y98S/L104F, F85L/Y98S/L104F).

**Cell Surface Expression of the Mutant Receptors**

We determined cell surface expression of the CCR5 mutants by fluorescence-activated cell sorting analysis, using five well characterized monoclonal antibodies (40, 42). The epitopes recognized by these monoclonal antibodies have been mapped to the N-terminal domain of the receptor (MC-5 and 3A9), the second extracellular loop (2D7), or a combination of extracellular domains (523 and 526).

Fig. 2 illustrates the average surface expression of the different mutants following normalization to wild-type CCR5 expression level. The TM2 mutant F85L exhibited a moderate but significantly reduced expression, reaching ~50% of the wt signal. Although the Y98S mutant was well expressed, the combination of both substitutions (F85L/Y98S) led to a reduced expression similar to that of F85L. Interestingly, although the single mutation L104F (in TM3) did not affect cell surface expression, its combination with F85L restored to normal the low expression observed for F85L alone (mutant F85L/L104F).

Strikingly, the triple mutant F85L/Y98S/L104F was not found at the cell surface, indicating that, although single and double
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Binding and Functional Properties of the Mutant Receptors

The functional consequences of the mutations and their combinations were characterized in terms of binding and intra-cellular responses, using the four natural agonists of CCR5: RANTES, MIP-1α, MIP-1β, and MCP-2. The binding affinities for the various ligands were determined by competition binding assays using $^{125}$I-RANTES or $^{125}$I-MIP-1β as labeled tracer, whereas functional responses were monitored by using the aequorin-based assay as described previously (13). Representative binding curves are shown in Fig. 3, functional concentration-action curves are shown in Fig. 4, and the data are summarized in Table I.

We first focused on residues Phe-85, Tyr-89, and Leu-104, located at the extracellular ends of TM2 and TM3. The F85L mutant was clearly impaired both at the level of binding, and in its functional response to chemokines. Although binding of RANTES was almost unaffected, the apparent affinities for MIP-1β and MCP-2 were significantly lower than those observed for wt-CCR5 (see Table I and Fig. 3), and MIP-1α binding was undetectable. The functional responses of F85L were mild (Fig. 4) and grossly consistent with the binding data. The potency of RANTES was moderately affected on this mutant, but its efficacy was reduced by 4-fold; MIP-1β and MCP-2 displayed a strong impairment of both their potencies and efficacies, whereas MIP-1α was almost inactive. The phenotype of the L104F mutant was grossly similar in terms of functional responses (with slightly better efficacies) despite binding parameters much closer to the wild-type levels. Interestingly, the addition of the L104F substitution to the F85L mutant (F85L/L104F) allowed partial restoration of the function of this mutant. Indeed, on the double mutant, MIP-1α was characterized by a binding affinity close to wt, and the functional response was significantly improved. The functional properties of MCP-2 appeared somewhat more affected by the double mutation, whereas RANTES and MIP-1β showed similar behaviors on the double mutant and on both single mutants.

Mutating Y89S affected mildly the activity of RANTES and MIP-1α (conserved potency, 2-fold reduction of $E_{\text{max}}$) decreased moderately the $E_{\text{max}}$ and potency of MIP-1β, but affected strongly the functional parameters of MCP-2, despite the normal affinity of this mutant in binding assays. Combining F85L and Y89S substitutions led to a severe impairment of functional responses, particularly for MIP-1β and MCP-2 (no response), and to a lower extent for MIP-1α, whereas RANTES was moderately affected. These effects appeared more than simply additive as compared with the single mutants. The Y89S/L104F mutant was well expressed at the cell surface, but no specific binding could be detected by competition binding assays, and this receptor was barely functional. Only RANTES could elicit a small signal at high concentrations, with a strongly reduced $E_{\text{max}}$ but a decent potency (Table I).

Modeling the TM2-TM3 Interface

The experimental data presented above demonstrate the importance of the TM2-TM3 aromatic cluster in the activation...
of CCR5. The fact that adding the L104F substitution to the F85L background partly restores the impaired function of this mutant (cell-surface expression and binding) might suggest a direct interaction between the two residues involved. Moreover, Tyr-89 might contribute to this interaction, as mutating this residue, although well tolerated by itself, is highly disruptive in the context of F85L, Y89S, or F85L/Y89S. Fig. 6A shows the location of the side chains of these residues in a CCR5 model using strictly the rhodopsin crystal as a template (27). The orientation of these side chains toward the lipidic environment is in apparent contradiction with the hypothesis that these residues are important for the structure and the activation mechanism of CCR5. We have, however, suggested previously that TM2 of CCR5 would adopt, in its outer half, a conformation that is different from that of bovine rhodopsin (13). In CCR5 and other chemokine receptors, the extracellular side of TM2 is predicted to be in close contact with TM3 (and not with TM1 as in rhodopsin) as a result of the structural action of a conserved Thr2.56-X-Pro2.58 motif (13). We now propose that this region of TM2 would be part of a structural and functional motif involving the neighboring part of TM3 (aromatic cluster and surrounding residues). To provide a structural framework allowing to understand better the experimental data presented above, we adopted a modeling procedure (detailed under “Experimental Procedures”) based on the following scheme.

**Independent Exploration of the Conformation of TM2 and TM3 by Molecular Dynamics Simulations**—Fig. 6 (B and C) shows the computed structures of TM2 (green) and TM3 (yellow). The bending of TM2 toward TM3, in its outer half, is tolerated in the context of the CCR5 helical bundle as the result of the relocation of TM3 toward TM5. It is important to note that these energetically available structures of TM2 and TM3 were obtained separately. Thus, the conformational spaces explored by these helices are the consequence of the amino acid sequence of TM2 and TM3 and not of steric hindrance between helices.

**The TM2-TM3 Interface Was Further Optimized by MD Simulations of the Seven-helix Bundle in an Apolar Environment**—Fig. 6 (D and E) shows the result of superimposing the representative structure of the MD simulation and the rhodopsin template. The specific residues in TM2 and TM3 of CCR5 generate structural differences in the extracellular part of the receptor, without modifying its more compact cytoplasmic surface.

**Membrane Protein Data Base Search**—Because stable structural motifs are likely to recur in proteins of known structure, we surveyed the data base of the structure of membrane proteins (see “Experimental Procedures”) for o-helix segments containing the aromatic cluster of TM2: the FWXXX motif. This motif is also found in subunit III of the bovine cytochrome c oxidase (PDB identification code 1occ), where transmembrane helix III of the enzyme interacts with the neighboring helix VII. Inspection of this motif in the cytochrome c oxidase structure reveals that the Phe-98 side chain in helix III interacts with Leu-252 residue in helix VII, and that the Tyr-102 side chain in helix III hydrogen bonds Ser-255 in helix VII. We propose a similar pattern to describe the interactions between TM2 and TM3 of CCR5; Phe-85 would interact with Leu-104, and Tyr-89 with Thr-99. It is important to note that Phe-85, Tyr-89, Thr-99, and Leu-104 are found specifically in CCR5, but not in CCR2, and contrib-

![Graph showing binding properties of the wt and mutant receptors.](http://www.jbc.org/)
ute therefore to a CCR5-specific motif important for the receptor structure.

Fig. 7A shows a detailed view of the TM2-TM3 interface in the model resulting from MD simulation. In this model, Leu-104 is located in an aromatic pocket formed by the side chains of Phe-85 and Trp-86, and the electron-poor C-H hydrogens of L104 interact with the electron-rich clouds of the aromatic rings. This type of C-H interaction plays a significant role in stabilizing local three-dimensional structures of proteins (43). Moreover, Phe-85 aromatic ring is located between Leu-103 and Leu-104 side chains. Thus, there is a significant interaction between the aromatic residues (Phe-85 and Trp-86) in TM2 and the hydrophobic residues (Leu-103 and Leu-104) in TM3. In addition, the TM2-TM3 interface is stabilized by a hydrogen bond between Tyr-89 and Thr-99. To evaluate the magnitude of the TM2-TM3 interaction that might be attributed to the Phe-85-Leu-104 and Trp-86-Leu-104 interactions, we performed ab initio quantum mechanical calculations on minimal recognition models consisting of the functional groups of the intervening side chains (see
**TABLE I**

Binding and functional properties of wt-CCR5 and mutant receptors

The pIC$_{50}$ values were obtained from competition binding experiments, using 125I-RANTES or 125I-MIP-1 as a tracer. Values represent the mean ± S.E. of at least two independent determinations.

| Receptor | Binding | Functional | MCP-2 |
|----------|---------|------------|-------|
| CRFC     | 9.64 ± 0.08 | 8.13 ± 0.16 | 3.32 ± 1.6 |
| L104F    | 9.74 ± 0.17 | 8.46 ± 0.24 | 4.0 ± 0.4 |
| Y89S     | 9.30 ± 0.17 | 8.40 ± 0.19 | 3.91 ± 0.17 |
| F109H    | 9.64 ± 0.08 | 8.46 ± 0.24 | 4.0 ± 0.4 |
| F112Y    | 9.92 ± 0.09 | 9.30 ± 0.09 | 6.5 ± 2.7 |

For practical reasons, some binding experiments yielding wild-type-like results both in the preliminary tests and dose-response curves were not further investigated; these values are given for illustration purposes.

**Experimental Procedures**. The energies of interaction of Phe-85 and Trp-86 with the multiple C-H hydrogens of Leu-104 are −2.4 and −2.6 kcal/mol, respectively.

**Structural and Functional Role of Aromatic Residues in TM3**

Residues Phe-109 and Phe-112 are located in TM3 within the outer third of the membrane and are predicted to face toward the center of the helix bundle, as inferred from the molecular model of CCR5 (see above) or the rhodopsin template itself. The F109H mutation had little effect on the receptor function. Both the binding properties and the functional response of the mutant were closely similar to those of wt-CCR5 for all four ligands (Table I, Figs. 3 and 4). In contrast, the conservative F112Y substitution influenced strongly the activation of the receptor by its agonists. The potencies of RANTES, MIP-1α, and MIP-1β were relatively preserved, whereas that of MCP-2 was decreased by −1 log. The efficacies of all ligands were however severely affected, with $E_{\text{max}}$ values ranging from 10 to 25% of the ATP response (Table I). Remarkably, combining these two substitutions in the F109H/F112Y mutant significantly restored the functional response of the receptor, with $E_{\text{max}}$ values ranging from 54 to 90% of the maximal cell response and improved potencies for all four ligands. The L104F/F109H/F112Y change, which combine the three mutations of TM3, produced a receptor phenotypically close to that of the single L104F. Although binding affinities were wt-like, activation was affected differentially for the different agonists; RANTES was almost not affected, MCP-2 severely impaired, with MIP-1α and MIP-1β showing intermediate behaviors. This suggest an addition of effects of the L104F single mutant (significantly affected) and the F109H/F112Y double mutant (mild effect). This is not surprising considering that these motifs are located in distant part of the structure, as, in our model, Leu-104 interacts with TM2, whereas Phe-109 and Phe-112 face TM5 and TM6, respectively (see “Discussion”).

**Mutant Y108A**

Considering the selective effect on receptor function observed after mutation of transmembrane residues in TM2 and TM3, it appeared also interesting to test the putative role of residue Tyr-108. This locus, referred as position 3.22 in the generalized numbering scheme of Ballesteros (see “Experimental Procedures”), is known as an important binding site in a wide variety of GPCRs (reviewed in Refs. 6, 12, and 24). Site-directed mutagenesis has been used to demonstrate its central role in different receptors for neurotransmitters and peptides. The Y108A mutant was well recognized at the cell surface by antibodies targeting the N terminus of the receptor (mean fluorescence above 60% of signal for wt-CCR5), whereas fluorescence was significantly decreased for antibodies recognizing either the second loop or a combination of extracellular domains, possibly underlying a conformational modification of the extra cellular loops of the receptor. This mutant exhibited a rather wild-type behavior following stimulation by RANTES, MIP-1β, or MIP-1α, with slightly reduced efficacies, but was completely unreactive to MCP-2. We could, however, measure high affinity binding of this ligand by competition binding assay (Fig. 3 and Table I). As for several other mutants described above, this mutant became unresponsive to MCP-2 without affecting the affinity for this ligand, whereas other CCR5 agonists (e.g. RANTES) were still able to simulate it. It is known that GPCRs, including CCR5 (40), can adopt multiple active states that trigger different intracellular cascades and are differentially stabilized by various agonists. In this line, one could hypothesize that MCP-2 would not be able to induce Ca$^{2+}$ increase
through this mutant receptor, while being able to trigger other intracellular cascades. As shown in Fig. 5A, the four natural agonists could trigger GTPγS binding through the wt receptor. In this assay, RANTES was the most potent agonist, whereas MCP-2 appeared more potent than MIP-1α and MIP-1β. As observed with the aequorin assay, MCP-2 was unable to stimulate the Y108A mutant in the GTPγS assay. It has been shown that chemokine-induced activation of CCR5 can lead to activation of p42/p44 MAP kinases. In this assay, RANTES stimulated the Y108A mutant, although with a reduced efficiency as compared with wt-CCR5, and MCP-2 was almost inactive, except for a weak response observed at 100 nM.

The specific alteration of the mutant response to MCP-2 is reminiscent of the behavior observed in our previous study, in which mutations of the TXXP motif also affected preferentially the biological response to MCP-2, without significantly affecting the binding of this chemokine. In the wt model, the Tyr-108 side chain is positioned in the face-to-edge orientation (T-shaped) with the indole ring of Trp-86 (see Fig. 7A). This type of ω-τ aromatic- aromatic interaction has been described as stabilizing a protein structure (46). It is expected that the conformational changes induced in TM2 by mutations affecting the TXXP motif would relocate the side chain of Trp-86, located four residues apart, and as a consequence affect its interaction with Tyr-108.

**DISCUSSION**

The three-dimensional model of CCR5 based on the rhodopsin crystal structure (including the specific modeling of the TM2-TM3 interface as described above; Fig. 6) provides a coherent framework to interpret the mutagenesis data detailed under “Results.” We propose that the residues varying between CCR5 and CCR2 at the level of the TM2-TM3 aromatic cluster are forming a specific interaction motif. In the wt receptor, Phe-85 would interact with Leu-104, whereas Tyr-89 would H-bond with Thr-99, and we suggest that the substitutions observed in the CCR2 sequence are indeed correlated. It is important to note that this modeled TM2-TM3 interface provides a hydrophobic environment for Thr-82, which mutations of the TXXP motif also affected preferentially. In this context, the O atom of Thr-82 orients the Cα bond of the backbone. The additional hydrogen bond between the O atom of Thr-82 and the carbonyl group increases the magnitude of the Pro-kink and this TXXP motif is a structural determinant involved in chemokine-induced activation (13).

To analyze in this structural framework the consequences of the amino acid substitutions explored experimentally, several of the mutations were introduced in our model, and we specifically studied the structural and dynamical properties of the extracellular part of the TM2-TM3 interface using MD simulations (see “Experimental Procedures”). This modeling procedure allows to propose a description of the TM interface for the mutants and suggests the nature of the structural changes that might lead to alterations of the receptor function (Fig. 7).

**Structural Interpretation of Mutations Located at the Extracellular End of TM2 and TM3**—The F85L mutation weakens the interaction between the side chain of this TM2 residue and Leu-103 and Leu-104 in TM3, as a Leu-Leu interaction is not of the same magnitude as a Leu-Phe interaction. As a consequence, in the simulation, Leu-85 side chain reorients away from TM3 (Fig. 7B).

Replacing Leu-104 by Phe dramatically modifies the TM2-TM3 interface. The aromatic side chain of Phe-104 would optimally interact with the other aromatic side chains in a face-to-edge configuration (46). Thus, in the molecular dynamic simulations, the side chain of Phe-104 tends to achieve this interaction with the aromatic side chains of both Phe-85 and Trp-86 (Fig. 7C). However, the side chain of Phe-104 in this conformation is bulkier than that of Leu, resulting in a significant reorientation of Phe-85 side chain toward the periphery of the
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Fig. 6. Modeling procedure of the TM2-TM3 interface. A, side-chain positions of residues 85, 89, and 104 in a model of the transmembrane domain of CCR5 based rigorously on the rhodopsin template. The Cα trace is shown as ribbon, in red for TM2 and TM3, in blue for the other helices. In such a model, these side chains face the membrane, which is hardly compatible with the functional role suggested by the mutagenesis experiments. B, superposition of 100 structures from independent Molecular Dynamics simulations of TM2 (in green) and TM3 (in yellow) onto the rhodopsin template (see “Experimental Procedures” for details). Only the Cα trace is shown for each structure, represented as ribbon. Viewed from the extracellular side, C, same as the previous panel but viewed from the side. It appears clearly that the conformational spaces are almost not overlapping. In particular, the conformational space of TM3 moves away from that of rhodopsin, allowing room for the kinked TM2 of CCR5. D and E, extracellular and side views of the representative structures of TM2 (in green) and TM3 (in yellow) selected from the simulations and used to construct the interface.

TM bundle, and in a weakening of the TM2-TM3 interface. It is important to note that the L104F mutation increases the polarity of the Thr-82 side chain environment. Thus, the presence of the electron-rich clouds of Phe-104 aromatic ring might facilitate hydrogen bonding to the Oδ atom of Thr-82, in a manner similar to that proposed for hydrogen bonding between benzene and water (47). This would contrast with the wild-type receptor, in which a hydrogen bond is formed with the polar peptide bond of the backbone. As already discussed, the additional hydrogen bond of the Thr to the peptide bond has a significant influence on the conformation of the helix (13, 26).

In the double F85L/L104F mutant, the aromatic side chain of Phe-104 is now located between Leu-85 and Leu-103 (Fig. 7D). Thus, the electron-rich clouds of the aromatic ring interact with the electron-poor C-H hydrogens of both Leu-85 and Leu-104. Notably, this mode of interaction of Phe-104 places Leu-85 in the proximity of Thr-82, mimicking the hydrophobic environment of Thr-82 in the wild-type receptor (see above). Moreover, the aromatic side chain of Tyr-89 is interacting in a face to edge configuration with Phe-104. Thus, it appears from this simulation that the F85L/L104F mutant would partly restore the packing of the TM2-TM3 interface.

This could explain the improvement in cell-surface expression and binding affinity for MIP-1α and MIP-1β, between the F85L and F85L/L104F mutants (see Table I). As the conformation of the short ECL1 (four residues) is likely influenced by the packing of the TM2-TM3 interface, we would suggest that point mutations (like F85L) could modify the conformation of the EC domain and affect the binding of ligands.

The triple mutant F85L/L104F/Y89S is intriguing. Intuitively, one would expect that, having replaced all differing aromatic positions into the CCR2 corresponding residues, the resulting mutant would show a better functional phenotype than the single or double mutant (being closer to the functional CCR2 receptor). However, it appeared that this triple mutant is not expressed at the cell surface (see Fig. 2), suggesting severe misfolding of the protein. Our modeling suggests that the addition of the Y89S substitution to the double F85L/L104F mutant would strongly modify the TM2-TM3 interface, because the shorter and non-aromatic side chain of Ser-89 cannot fulfill the interaction with both Thr-99 and Phe-104 (Fig. 7D). We hypothesize that this major structural difference between this mutant interface and the wt structure could be a factor responsible for the weak expression, either through a perturbation of the folding process, or through destabilization of the folded protein, leading to rapid internalization and/or degradation. In the single Y89S mutant, this modification of the packing of TM2 and TM3 would not happen as the Phe-85-Leu-104 interaction maintains a proper distance between the two helices. Moreover, Thr-991,23 would tend to interact with the side chain of Gln-932,67 (Fig. 7C), changing moderately the interface. Notably, the Y89S mutant shows a normal level of expression and is moderately affected in the functional tests, suggesting that the Tyr-89-Thr-99 interaction proposed here is necessary for full functional efficiency, but not for the structural stability of the receptor. The double mutants F85L/Y89S and Y89S/L104F show a level of expression reduced by ~50%, a functional response strongly affected for F85L/Y89S and almost completely abolished for Y89S/L104F, underlining the increase in structural perturbation caused by the additional mutations.

Combining the experimental results with our modeling approach leads to the following picture; the function of the CCR5 receptor is strongly dependent on the interface between the extracellular ends of TM2 and TM3, which is determined by


series of interhelical interactions. The structural and dynamical properties of this interface are maintained by, at least, two polar interactions: Tyr-89-Thr-99 and Phe-85-Leu-104. These interactions are in equilibrium with each other, providing balanced structural constraints. Perturbing this interface by mutating one of these residues would modify this equilibrium and affect the function of the receptor.

This model implies that, in the case of the CCR2 receptor, the TM2-TM3 interface is organized differently, and that the substitutions tested here are counterbalanced by other changes in the sequence. In particular, one can spot the T993.23 change between the two receptors (see Fig. 1), but other changes (like A922.66 or Q932.67) could also be important. Interestingly, a chimeric construct involving ICL1, TM2, ECL1, and TM3 of CCR2 in a CCR5 background appeared to be well expressed and functional (35), suggesting that interactions between TM2 and TM3 interface and the TM3-TM6 interaction for wt and mutant receptors. A, model of the TM3 interface obtained for the wt receptor. Only interacting side chains are shown, highlighting the proposed Phe-85-Leu-104 and Tyr-89-Thr-99 interactions (see text for details). Panels B and C show the models for F85L and L104F mutants, respectively, illustrating the loss of interaction at the 85–104 locus for these receptors. D, model of the double F85L/L104F mutant showing the proposed recovered interaction. E, wt model showing the location of Phe-109 and Phe-112 and featuring the conformation of Asn-252 interacting with the backbone in TM6. F, model of the F112Y mutant, showing how this mutant could modify the conformation of Asn-252 in TM6. G, model of the double F109H/F112Y, illustrating how the F109H additional mutation would recover the structural effect of the F112Y mutant.

Fig. 7. Molecular models of TM2-TM3 interface and the TM3-TM6 interaction for wt and mutant receptors.
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and TM3 are sufficient to obtain structural stability and functionality (as the other helices were strictly from CCR5).

The Extracellular Ends of TM2 and TM3 Are Involved in the Activation Mechanism—By mutating the Thr-2.56-X-Pro-2.58 motif, we previously have shown that the structural integrity of TM2 is crucial for CCR5 function (13). According to the nature of the substitution (various residues in place of Thr-82\(^{2.56}\), or Ala instead of Pro-84\(^{2.58}\), we could modulate the extent of the structural perturbation, which was translated into a functional defect. However, the different agonists of CCR5 were affected differently by these mutations, suggesting that this part of the receptor is involved in the activation process in a ligand-specific manner.

We proposed that this TXP motif governs the structural and dynamical properties of the extracellular end of TM2. By looking at the specific role of Phe-85\(^{2.56}\) and Tyr-89\(^{2.52}\) of a Pro could significantly modulate the structure of the pro-\(H_9251\). Line-induced deformation, and therefore be functionally rele-

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We proposed that this TXP motif governs the structural and dynamical properties of the extracellular end of TM2. By looking at the specific role of Phe-85\(^{2.56}\) and Tyr-89\(^{2.52}\), we now probe the elements involved in the ligand-induced activation in this particular part of the TM bundle. Interestingly, we find the same trend as that observed after mutating the TXP motif; the ligand sensitivity to the mutations is variable, RANTES being the least affected agonist, whereas MCP-2 is the most sensitive.

Structural Modeling of Mutants Involving the Middle of TM3—Phe-109 and Phe-112, variable between CCR5 and CCR2, are not facing TM2 in the CCR5 model, but are oriented toward TM5 and TM6. We therefore undertook full bundle MD simulation to model (see “Experimental Procedures”) the possible effects of the F109\(^{2.59}\)H, F112\(^{2.59}\)Y, and F109H/F112Y mutations, and the putative interaction between these residues and TM5 and/or TM6.

Although the F109H change does not modify the pattern of helix-helix interactions as compared with the wt receptor (data not shown), changing Phe-112 for Tyr modifies the interactions between TM3 and TM6 (Fig. 7, panels E and F). In our simulation, the hydroxyl group added by the mutation forms a H-bond with Asn-252\(^{3.36}\) in TM6. In the wt receptor, this Asn H-bonds back to the backbone carbonyl of Trp-248\(^{3.33}\) and Tyr-112 interacts through its H-atom with the backbone carbonyl of Trp-248\(^{3.33}\). We therefore undertook full bundle MD simulation to probe the elements involved in the ligand-induced activation in this particular part of the TM bundle. Interestingly, we find the same trend as that observed after mutating the TXP motif; the ligand sensitivity to the mutations is variable, RANTES being the least affected agonist, whereas MCP-2 is the most sensitive.

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Activation of CCR5 by Chemokines Involves an Aromatic Cluster between Transmembrane Helices 2 and 3

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