Annexin A1 Interaction with the FPR2/ALX Receptor

IDENTIFICATION OF DISTINCT DOMAINS AND DOWNSTREAM ASSOCIATED SIGNALING

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Background: FPR2/ALX is activated by many ligands, including annexin A1 (AnxA1), which activates resolution circuits in inflammation.

Results: Cells transfected with FPR2/ALX and clones with domains swapped with FPR1 afforded identification of N-terminal and extracellular loop II domains as transducers of AnxA1 signaling.

Conclusion: We identified AnxA1 distinct domains of FPR2/ALX and unveiled potential specific signaling.

Significance: FPR2/ALX domain identification permits development of anti-inflammatory AnxA1 mimetics.

Understanding how proresolving agonists selectively activate FPR2/ALX is a crucial step in the clarification of proresolution molecular networks that can be harnessed for the design of novel therapeutics for inflammatory disease. FPR2/ALX, a G protein-coupled receptor belonging to the formyl peptide receptor (FPR) family, conveys the biological functions of a variety of ligands, including the proresolving mediators annexin A1 (AnxA1) and lipoxin A4, as well as the activating and proinflammatory protein serum amyloid A. FPR2/ALX is the focus of intense screening for novel anti-inflammatory therapeutics, and the small molecule compound 43 was identified as a receptor ligand. Here, we used chimeric FPR1 and FPR2/ALX clones (stably transfected in HEK293 cells) to identify the N-terminal region and extracellular loop II as the FPR2/ALX domain required for AnxA1-mediated signaling. Genomic responses were also assessed with domain-specific effects emerging, so the N-terminal region is required for AnxA1 induction of JAG1 and JAM3, whereas it is dispensable for modulation of SGPP2. By comparison, serum amyloid A non-genomic responses were reliant on extracellular loops I and II, whereas the small molecule compound 43 activated extracellular loop I with downstream signaling dependent on transmembrane region II. In desensitization experiments, the N-terminal region was dispensable for AnxA1-induced FPR2/ALX down-regulation in both the homologous and heterologous desensitization modes.

The biological properties of G protein-coupled receptors are fundamental to the control of the host inflammatory reaction mounted against an insult, in terms of both promoting the response (e.g. chemokine-mediated leukocyte recruitment) and limiting its duration and intensity. This second side of the innate immune response is currently referred to as the resolution of inflammation phase. Within this context, an important role is played by the duo annexin 1 (AnxA1) and its receptor, formyl peptide receptor (FPR) type 2, whose acronym is FPR2/ALX because it also conveys the inhibitory signals induced by lipoxin A4 and resolvin D1. FPR2/ALX is an intriguing receptor not only because it was the first one shown to mediate actions elicited by lipids, peptides, and proteins but also because it can mediate activating proinflammatory responses, such as those elicited by serum amyloid A (SAA) (2). A recent study by Ye and co-workers (3) chiefly demonstrated the ability of FPR2/ALX to modulate cell responsiveness in a ligand-biased fashion. Based on analogies with G protein-coupled receptors, several mechanistic hypotheses can be put forward to explain the versatility of this receptor, including homologous and/or heterologous dimerization, as well as the recruitment of ligand-specific signaling pathways (2, 4). Another plausible option is that ligands might activate specific receptor domains, thus promoting at least in part not overlapping downstream responses. As an example, the small lipid lipoxin A4 has been shown to activate FPR2/ALX by interacting with extracellular loop III and the associated transmembrane region (5).

This study was undertaken with a particular focus on the domains required for AnxA1-dependent receptor activation. We took advantage of recently characterized stable clones expressing chimeric receptors with different regions of FPR2/ALX and the cognate receptor FPR1 (6). Activation of these chimeric receptors by AnxA1 was analyzed and compared, in some experiments, with responses evoked by SAA and, for completion and to assess the potential impact these finding might have on drug discovery, by the small molecule agonist compound 43 (C43) (7).

The abbreviations used are: AnxA1, annexin 1; FPR, formyl peptide receptor; SAA, serum amyloid A; C43, compound 43; BAPTA-AM, 1,2-bis(o-aminophenoxy)ethane-N,N',N''-tetraacetic acid acetoxymethyl ester; mAb, monoclonal antibody.

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Experimental Procedures

Cell Culture—HEK293 cells stably transfected with FPR1 and FPR2/ALX were obtained as described (8) and maintained in culture with DMEM/F-12 supplemented with 10% FCS, gentamicin (50 μg/ml), nonessential amino acids (500 μg/ml), and Geneticin (200 μg/ml) solution. Fig. 1A shows the schematics for FPR2/ALX. HEK293 cells transfected with the FPR1-FPR2 chimera were obtained as described (6) and are listed in Fig. 1B. Cells were cultured in DMEM supplemented with 10% FCS, l-glutamine (500 μg/ml), penicillin/streptomycin (500 μg/ml), HEPES (1 mM), and Geneticin (800 μg/ml) solution.

Ligands and Inhibitors—AnxA1 was produced as described previously (9). SAA was purchased from PeproTech Laboratories (London, United Kingdom), and C43 was a generous gift from Amgen (Thousand Oaks, CA). The MEK1 inhibitor PD98059 and the calcium inhibitor BAPTA-AM were obtained from Cell Signaling Technologies (Hertfordshire, United Kingdom) and Calbiochem, respectively.

Human Polymorphonuclear Leukocyte Isolation—Peripheral blood was collected from healthy volunteers by intravenous withdrawal in 3.2% sodium citrate solution (1:10). All healthy volunteers gave oral and written consent, and cell separation was covered by Ethical Approval 05/Q0603/34 (East London and The City Research Ethics). Granulocytes (polymorphonuclear leukocytes) were isolated from blood via density centrifugation on a Histopaque 1119/1077 gradient (Sigma-Aldrich) according to the manufacturer's instructions and suspended in PBS containing 0.5% BSA. Contaminating erythrocytes were removed by lysis with cold Milli-Q water.

Calcium Mobilization Assay—HEK293 cells or freshly prepared polymorphonuclear leukocytes were incubated with 2 μM Fura 2-AM (Molecular Probes, Paisley, United Kingdom) and 1 μM Pluronic acid F-127 (Molecular Probes) in extracellular solution (13 mM glucose, 10 mM HEPES, 147 mM NaCl, 2 mM KCl, 1 mM MgCl₂, and 2 mM CaCl₂ at pH 7.3) at 37 °C for 1 h in the dark. Subsequently, cells were washed three times with the extracellular solution and added to 96-well plates prior to stimulation with agonists at the indicated concentrations. Ionomycin (1 μM) was used as a positive control. Mobilization of intracellular calcium was measured by recording the ratio of fluorescence emission at 510 nm after sequential excitation at 340 and 380 nm using the NOVOstar microplate reader (BMG LABTECH Ltd., Aylesbury, United Kingdom).

Western Blot Analysis—The protein content of the lysate was determined via Bradford protein assay (Bio-Rad). Cells lysates were boiled in 6× Laemmli buffer, analyzed by standard SDS-PAGE (12%), and electrophoretically transferred to PVDF (Millipore, Watford, United Kingdom). Membranes were incubated with rabbit anti-phospho-p44/42 MAPK (1:1000 dilution) and anti-total p44/42 (clone 137F5; 1:1000 dilution) antibodies (Cell Signaling Technologies) diluted in Tris-buffered saline containing 0.1% Tween 20 (TBST) and 1% BSA overnight at 4 °C. Membranes were washed for 30 min with TBST and incubated with HRP-conjugated goat anti-rabbit secondary antibody (1:2000 dilution; Dako, Cambridge, United Kingdom) for 2 h at room temperature. After stripping, membranes were also incubated with anti-β-actin antibody (1:10,000 dilution; Sigma-Aldrich) in TBST and 5% nonfat dry milk. Human actin was detected with HRP-conjugated goat anti-mouse antibody (1:5000 dilution). Proteins were then detected using an ECL detection kit and visualized on Hyperfilm (GE Healthcare).

Flow Cytometry Analysis—FPR1 and FPR2/ALX cell surface expression was measured by incubation with anti-FPR1 (R&D Systems, Abingdon, United Kingdom) or anti-FPR2 (Genovac, Brussels, Belgium) monoclonal antibody (mAb; 1:100 dilution in both cases); a final staining with a rabbit anti-mouse IgG (clone STAR9B; 1:200 dilution; Serotec) was then conducted. Flow cytometry analysis was performed by analyzing ≥10,000 events using a FACSCalibur flow cytometer (BD Biosciences) equipped with CellQuest software (BD Biosciences), followed by analysis using FlowJo (Version 9.2, TreeStar Inc., Stanford, CA). Results are reported as median fluorescence intensity units.

Real-time PCR Using SYBR Green I Dye—The assay was performed as shown previously (10). Briefly, RNA was extracted using an RNAeasy kit (Qiagen 74104) according to the manufacturer’s instructions. Total RNA was normalized to 5 μg/reaction. cDNA was produced using SuperScript III (Invitrogen). Samples were normalized to 100 ng of cDNA/well and loaded in duplicate for each gene using Power SYBR Green PCR Master Mix (AB 4367659). QuantiTect primer assays (Qiagen) were used: human GAPDH (QT01192646), human SGPP2 (QT00041832), human JAG1 (QT00031948), and mouse JAM3 (QT00024997). GAPDH was used as an endogenous control. PCR ramping protocols were standardized for QuantiTect primer assay sets at 95 °C for 15 min, followed by 40 cycles at 94 °C for 15 s, 55 °C for 30 s, and 72 °C for 30 s.

Statistical Analysis—Data are expressed as means ± S.E. of experiments conducted at least three times. Data were analyzed using Student’s t test (two groups) or one-way analysis of variance, followed by Dunnett’s post hoc tests (more than two groups). In all cases, a p value of <0.05 was taken as significant.

Results

FPR1, FPR2/ALX, and Chimeric Receptors—Human FPR1 and FPR2/ALX display ~69% homology (2) as illustrated in Fig. 1A. To evaluate the contribution of single domains in FPR2/ALX activation, we used eight different chimeric receptors with domains swapped between FPR1 and FPR2/ALX, generated as described previously (6) and detailed in Fig. 1B. The degree of receptor expression, as demonstrated in Ref. 6, was assessed using specific mAbs against FPR1 or FPR2. Supplemental Fig. 1 reports these results and shows that chimeras C, D, and G were recognized by the anti-FPR2/ALX mAb, whereas all others were detected with the anti-FPR1 mAb (supplemental Fig. 1).

Calcium Mobilization Induced by FPR1 and FPR2/ALX Agonists—AnxA1 and its bioactive peptides are able to mobilize intracellular Ca²⁺ in primary polymorphonuclear leukocytes (11, 12) and cells transfected with FPR2/ALX and/or FPR1, respectively (12, 13). Using native FPR2/ALX, AnxA1 afforded concentration-dependent responses, with an EC₅₀ value of ~6 nm (Fig. 1C). SAA and C43 also produced canonical concentration-dependent responses, with EC₅₀ values of ~30 and ~100 nm, respectively (Fig. 1C).
To compare the ligand-binding domains activated by the different agonists, Ca$^{2+}$ mobilization in chimeric HEK293 cells was studied. AnxA1 (selected concentration, 10 nM) promoted Ca$^{2+}$ flux in native FPR2/ALX, with similar responses in clones C, G, and H (all ranging from 40 to 60% of the maximal response), with little or no effect on FPR1-expressing cells and the other clones (Fig. 2A). The addition of SAA (100 nM) afforded ~50% of the ionomycin response with both native FPR2/ALX cells and clones E, F, G, and H, whereas no responses were detected with the other clones, including clone B, which bears extracellular domain III of FPR2/ALX (Fig. 2B). The small molecule C43 (1 μM) produced a similar degree of response in FPR2/ALX cells and was active only in clones E and F (Fig. 2C). For completion, we also tested formyl-methionyl-leucyl-phenylalanine (300 nM; as selected from preliminary analyses (not shown)), which, as expected, was active only in native FPR1 (but not FPR2/ALX) cells; with respect to the clones analyzed, formyl-methionyl-leucyl-phenylalanine was particularly active in clones A and B (supplemental Fig. 2).

**ERK Activation Induced by AnxA1 and the Two FPR2/ALX Agonists**—Next, we determined whether the clone specificity that emerged so far could be confirmed with another readout for FPR2/ALX activation, such as phosphorylation of ERK (8, 12). As sometimes evident with transfected cells, the overexpressed receptor was, in some instances, already (partially) activated, yet the addition of AnxA1 (10 nM) provoked clear ERK phosphorylation in both FPR2/ALX and clones C, G, and H as monitored at 5 and 10 min post-addition (Fig. 3A). Clone E was also tested because it was unresponsive to AnxA1 for transient Ca$^{2+}$ flux, and the same inactivity held true for phospho-ERK. For completion, we also tested SAA and C43. The positive asso-
FPR2/ALX Domains for AnxA1-mediated Activation

**FIGURE 3. Clone-specific ERK1/2 phosphorylation induced by AnxA1, SAA, and C43.** HEK293 cells expressing native and chimeric clones were treated with AnxA1 (10 nM; A), SAA (0.1 μM; B), or C43 (1 μM; C) (concentrations selected from Fig. 1) for 5 or 10 min as indicated, and the phosphorylation of ERK1/2 was monitored by Western blot analysis. Representative blots from three or more experiments with distinct cell preparations are shown. tERK, total ERK.

**DISCUSSION**

FPR2/ALX is a G protein-coupled receptor that is able to recognize many ligands, often of a very different chemical nature (2, 5). Following binding with their agonists, FPR2/ALX undergoes a conformational change that allows functional interaction with Gαi1, Gαi2, and Gαi3 and association with Gq, Gq, and Gα16 (16). This event triggers activation of a variety of signaling pathways, including fluxes in intracellular calcium and activation of phospholipases A2, C, and D, PI3K, and MAPK (17). Receptor activation would also cause rapid phosphorylation, leading to phospholipase C-mediated receptor desensitization and internalization (18). FPR2/ALX downstream activation of the MAPK pathway, in particular ERK1/2, regulates cell chemokinesis (19). However, a major conundrum in the biology of this receptor is, on one hand, the large numbers of putative ligands and, on the other hand, the apparent ability to elicit contrasting cellular responses in a ligand-biased fashion. The latter point has been chiefly demonstrated in a recent study in macrophages, where both putative “proinflammatory” (or rather activating) FPR2/ALX agonists (such as SAA) and inhibitory agonists (such as AnxA1 and lipoxin A4) were able to convey their message through this receptor and afford functional antagonism to each other (3). This study aimed to shed light on this heavily ligand-biased FPR2/ALX pharmacology by assessing the possibility that distinct receptor domains might interact with distinct ligands and potentially elicit different...
receptor downstream responses. We took advantage of well characterized stably transfected cells with varied FPR1-FPR2/ALX chimeric receptors (6) and began by assessing robust readouts, such as Ca²⁺ fluxes and ERK phosphorylation. An important control useful also for identifying the epitope recognized by commercially available mAbs was the overall amount of receptor expression. It appears that the anti-FPR2/ALX mAb likely binds to a tridimensional epitope formed between the receptor N-terminal domain and extracellular loops II and III. When the signal was positive, this mAb revealed similar receptor density on the cell surface as in native FPR2/ALX-transfected cells. All other chimeras were detected solely by anti-FPR1 mAb. It is evident that the mAbs are able somehow to penetrate the transmembrane domains because the anti-FPR2/ALX antibody was active in clone D (containing sequence 40–86 of the native receptor), whereas the anti-FPR1 mAb was not active in clone F (containing sequence 106–145 of the FPR2/ALX receptor).

The combined data produced with the clones for AnxA1 indicate that the N-terminal domain of FPR2/ALX is sufficient for this ligand to elicit rapid downstream responses, whereas the second extracellular loop is required to provoke more sustained changes, such as those leading to modulation of gene expression. Here, we selected three genes as readouts from our previous study (10) and confirmed modulation upon FPR2/ALX engagement; intriguingly, clone C, which could transduce Ca²⁺ and phospho-ERK, modulated JAG1 and JAM3, but clone H, with both extracellular loops II and III, was nearly as active as the native receptor in SGPP2 induction. This dichotomy of responses led us to propose the model discussed below and shown in Fig. 7. It is worth noting that, in a previous study using

![Figure 4](image-url)  
**FIGURE 4.** Clone-specific genomic responses evoked by AnxA1. Real-time PCR data for SGPP2 (*A*), JAG1 (*B*), and JAM3 (*C*) gene product expression in HEK293 cells expressing native and the indicated chimeric clones following 4 h of incubation with AnxA1 (0.5 μM). The response produced with control CMV empty plasmid-transfected HEK293 cells was taken as 1. Data (mean ± S.E.) are from three distinct experiments performed in duplicate. *, *p* < 0.05 versus control CMV-transfected HEK293 cells.

![Figure 5](image-url)  
**FIGURE 5.** N-terminal domain of FPR2/ALX is dispensable for homologous and heterologous desensitization. *A*, HEK293 cells expressing native and chimeric clones C, G, and H were treated with vehicle, AnxA1 (10 nM), and SAA (0.1 μM) for 5 min, followed by a second treatment with AnxA1 (10 nM). Data (means ± S.E. of more than three distinct experiments done in triplicate) are reported as percent ionomycin (1 μM) response. *, *p* < 0.05 versus vehicle pretreatment. *B*, selected clones (see Fig. 1 for details).
a similar approach but different chimeric strategy, the FPR2/ALX agonist lipoxin A4 was shown to interact with extracellular loop III (5).

SAA is an acute-phase protein that is able to activate a variety of receptors, including FPR2/ALX (20, 21). Our data confirm the genuine agonistic activity of SAA on FPR2/ALX without any activation of FPR1. In terms of receptor domains, analysis of Ca²⁺ fluxes and ERK phosphorylation indicates an interaction with extracellular loops I and II, whereas the N-terminal and extracellular loop III domains are dispensable for the Ca²⁺ response. Concerning C43, in line with the ability of small molecules to “penetrate” inside G protein-coupled receptor loops (22), this FPR2/ALX agonist (7, 23) did not elicit Ca²⁺ flux responses in FPR1-transfected cells and was active in native FPR2/ALX, as well as in clones E and F; therefore, extracellular loop I, transmembrane region III, and intracellular loop II contain the binding site for C43. It should be noted here that C43 produces anti-inflammatory effects when given in vivo (7), which are lost in Fpr2/Fpr3 null mice (15); however, other studies have claimed an involvement of Fpr1 (the mouse counterpart of human FPR1) in its anti-inflammatory profile (24, 25), yet activation of human FPR1 could not be detected in these experimental settings with the stably transfected cells. Next, studies should establish whether C43 permits FPR1 and FPR2/ALX dimerization.

We completed this investigation by determining FPR2/ALX homologous and heterologous desensitization because they had already been reported for AnxA1 and SAA (3) and also because of the functional consequences these processes may have in modulating cell responsiveness in inflammatory settings (12, 15). It emerged that AnxA1 engagement of FPR2/ALX extracellular loop II is required for desensitization to occur, as determined by Ca²⁺ readout. As shown recently (26), the C terminus (present in clone H) may also have an important role in FPR2/ALX desensitization. Analyses conducted with...
FPR2/ALX Domains for AnxA1-mediated Activation

clone C, which contained solely the FPR2/ALX N-terminal region linked to the nearly full FPR1 sequence, corroborated the specificity of full-length AnxA1 for FPR2/ALX and its inability to engage FPR1. This is in contrast to the ability of short AnxA1-derived peptides to activate all members of the human FPR family (12, 27). We have recently discussed how this dichotomy of protein/peptide behavior may be of biological significance in the context of resolution of inflammation (28).

In summary, we studied the domains of FPR2/ALX engaged by AnxA1 in comparison with SAA, which elicits opposite cellular responses (3), and the small molecule C43 in view of the potential exploitation of the pharmacology of this receptor for the development of novel anti-inflammatory therapeutics. The picture that emerges is illustrated in Fig. 7, which shows that AnxA1 can interact with the N-terminal domain and extracellular loop II. Because of the large structure of this protein (29) and the fact that, in the presence of calcium, it undergoes a conformational change with exposure of its N-terminal domain (30, 31), we hypothesize that at least two FPR2/ALX sites are required to accommodate this agonist (Fig. 7), a hypothesis proposed quite a few years ago when structural data were scarce (32). A recent study proposed a three-point binding model for small molecule ligand interaction with FPRs, for FPR2/ALX (33), confirming original studies with FPR (14). These binding sites are within the non-conserved amino acid residues 84, 85 (point 1), 163 (point 2), and 284 (point 3) (14, 33). Also in our model, AnxA1 binds the FPR2/ALX receptor in the binding pocket that comprises residue 163 (present in clone G). Residues 84, 85, and 163 are in the FPR2/ALX-binding region required by C43 and SAA to induce receptor activation.

In conclusion, information has been obtained regarding the ligand-specific domains required for classical FPR2/ALX readouts, such as Ca²⁺ fluxes and ERK phosphorylation. Because of the emerging importance of FPR2/ALX in the regulation of the inflammatory process, we reasoned that a better appreciation of the binding domains and the ensuing signaling characteristic of the interaction with the endogenous agonist AnxA1 is not only of biological importance but will also be significant in the development of AnxA1 mimetics able to engage this effector of resolution.

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