Genetic engineering of the glucocorticoid receptor by fusion with the herpes viral protein VP22 causes selective loss of transactivation

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Abstract

The development of methods for engineering proteins with novel properties opens the way to manipulating intracellular processes in a therapeutically useful way. Glucocorticoids, acting via glucocorticoid receptors (GR), are potent anti-inflammatory agents, acting to oppose nuclear factor kappa B (NFκB) function. The herpes viral protein, VP22, has been reported to confer intercellular trafficking activity on ‘cargo’ proteins, potentially facilitating gene therapy with intracellular proteins.

VP22GR, resulting from the addition of VP22 to the N terminal of GR, was equipotent with the wild-type GR in opposing NFκB p65-driven expression of an NFκB reporter gene. Surprisingly, VP22GR was incapable of inducing transactivation of positive glucocorticoid reporter genes (MMTV-luc and TAT3-luc). Furthermore, the VP22GR had powerful dominant negative activity on both endogenous and exogenous GR transactivation. VP22GR was cytoplasmic in quiescent cells, and after hormone addition underwent nuclear translocation to share the same distribution as the GR. The ability of the VP22GR to selectively confer and enhance glucocorticoid-dependent transrepression of NFκB may be of use therapeutically in e.g. transplant rejection, inflammatory arthritis or asthma.

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Introduction

Glucocorticoid effects on transcription may be mediated by the direct binding of activated glucocorticoid receptors (GR) to target DNA, homodimerisation and recruitment of co-activators and also by GR interfering with other transcription factor function, including AP-1, nuclear factor kappa B (NFκB) and NUR77 (Jonat et al. 1990, Miner & Yamamoto 1991, Schule & Evans 1991, Konig et al. 1992, Heck et al. 1994, Ray & Prefontaine 1994, Caldenhoven et al. 1995, Scheiman et al. 1995a,b, Wade et al. 1995, Horwitz et al. 1996). The proinflammatory actions of NFκB are well characterised, and opposition of NFκB by GR can explain many of the anti-inflammatory actions of glucocorticoids (Scheiman et al. 1995b, Cato & Wade 1996, Nissen & Yamamoto 2000). Glucocorticoid sensitivity is related to GR expression (Vanderbilt et al. 1987) and, furthermore, opposition of NFκB action, and so anti-inflammatory activity, can be enhanced by overexpressing the GR (Mathieu et al. 1999).

The GR has a defined domain structure which comprises an N terminal transactivation domain (AF-1), a central DNA binding domain (DBD) which is the most highly conserved structure with other members of the steroid hormone receptor family, and a C terminal ligand binding domain (LBD), which also includes a second activation domain, a hinge region, and sites for interaction with heat shock proteins (Hollenberg & Evans 1988). The N terminal transactivation domain extends from amino acid 77 to 262, but a core domain from 187 to 244 contributes most of this activity (Warnmark et al. 2000). The C terminal transactivation domain depends on ligand-induced folding to recruit co-activator proteins, including members of the p160 family, GRIP1 and SRC1 (Hong et al. 1997, Collingwood et al. 1999, McKenna et al. 1999). GR inhibition of NFκB is mediated by interaction with the RelA (p65) component (Scheiman et al. 1995b). The GR DBD binds to RelA when it is bound to DNA, and the GR LBD represses the RelA transactivation domain (Nissen & Yamamoto 2000).

VP22 is a 38 kDa structural protein from the herpes simplex virus (HSV) and has the property of intercellular transport (Elliott & O’Hare 1997). VP22 trafficking appears to involve contacts with the actin architecture of the cell, and does not require the Golgi apparatus (Elliott & O’Hare 1997). It has been shown that ‘cargo’ proteins may be fused with the VP22 to allow them intercellular trafficking potential (Elliott & O’Hare 1999). Such ‘trafficked’ proteins appear to retain function, as in the case of HSV thymidine kinase, and p53 (Koelle et al. 1998, Phelan et al. 1998). An additional property of both VP22 alone and its fusion proteins appears to be its intracellular distribution.
In unstimulated cells the GR is sequestered in the cytoplasm, and removal of its C terminal, ligand binding domain generates a nuclear resident, constitutive transcriptional activator (Hollenberg & Evans 1988). As VP22 targets cargo proteins to the nucleus of transfected cells (Fang et al. 1998), or the cytoplasm of synthesising cells and the nucleus of protein recipient cells (Elliott & O’Hare 1999), by imposing additional controls on its intracellular localisation, it may alter GR function. In addition, it may be possible to influence tissue glucocorticoid sensitivity by allowing intercellular movement of GR.

This work describes the generation and analysis of a VP22–GR fusion protein with extreme dissociation of RelA transrepression from transactivation. Such proteins may have utility in dissecting the role of the two pathways for functional effects in vitro, and also for development as gene therapy for inflammatory disease.

Materials and Methods

Construction of plasmids

The human GR cDNA was amplified using two primers: N terminal 5′ CTG CGT CTT CAC CCT CAC TGG C and C terminal 5′ TCA CTT TGG ATG AAA CAG AAG TTT TTT TTT G and was cloned directly into the pVP22/myc-his TOPO vector (Invitrogen, Paisley, UK). This includes the full-length VP22 coding region to codon 301. This resulted in an in-frame fusion of the 3′ terminal of the VP22 coding region with codon 42 (serine) of the human GR, VP22GR. The construct was fully sequenced to confirm freedom from PCR introduced errors.

The VP22 GRmyc was generated using the same 5′ primer but with the 3′ primer lacking the endogenous translation stop codon to enable translation to continue across the myc epitope. The C terminal primer sequence was: 5′ CTT TGG ATG AAA CAG AAG TTT TTT TTT GAT A. Again, the construct was fully sequenced.

VP22 GRdim (dimerisation deficient) was generated from VP22GR by excising a cassette with BsmB1 and ClaI and replacing it with the cassette from pSGGR(A458T) (a kind gift of Dr P Herrlich, Institute for Toxicology and Genetics, Karlsruhe, Germany) (Heck et al. 1997, Reichardt et al. 1998). The 3′ terminal of VP22GR was disrupted by excising a cassette from GR nucleotides 1536 to 1921 with BstB1, and religating it to make VP22GR(-LBD). All recombinant plasmids were sequenced.

The GFPGR consists of the full-length murine GR cDNA with a 5′ fusion to the green fluorescent protein (GFP) cDNA, the kind gift of Dr Paul Housley, University of South Carolina (Galigniana et al. 1998).

The TAT3-luc reporter, consisting of three copies of the glucocorticoid response element (GRE) from the tyrosine aminotransferase gene upstream of a minimal promoter linked to luciferase, was the gift of Prof. K Yamamoto (Iniguez et al. 1997). The MMTV-luc plasmid consisted of 1-5 kb of the mouse mammary tumour virus long terminal repeat fused to the firefly luciferase gene in the pXp-2 backbone (Ray et al. 1996a). The full-length wild-type human GR expression vector (wtGR) had the human GR cDNA inserted into the pcDNA3 plasmid (Ray et al. 1999). The full-length rat GR expression vector, p6RGR, consisted of the rat GR cDNA driven by the Rous sarcoma virus (RSV) promoter. p6RGRN525 lacked the C terminal ligand binding domain by deletion from amino acid 525. p6RGR407C lacked the N terminal up to amino acid 407. All three rat constructs were the kind gifts of Dr Jorge Iniguez-Lluhi and Professor Keith Yamamoto, University of California at San Francisco, USA (UCSF) (Godowski et al. 1988). The estrogen receptor expression vector (cytomegalovirus–estrogen receptor; CMV-ER) contained the full-length human ERα in the pcDNA3 vector, and was the kind gift of Dr Maurice Needham, AstraZeneca, UK. The mineralocorticoid receptor (MR) expression vector (CMV-MR) contained the human MR cDNA inserted into pcDNA3, and was the kind gift of Dr Maurice Needham. A CMV-βGal plasmid was used to control for differences in transfection efficiency, as previously described (Ray et al. 1999).

Cell culture and transfection

COS 7, HeLa and A549 cells were obtained from ECACC (Salisbury, UK), and were cultured as previously described (Ray et al. 1999) in DMEM with glutamax (Gibco, Paisley, Strathclyde, UK) supplemented with 10% fetal calf serum. No antibiotics were used. Cells were transfected using Lipofectamine Plus (Life Technologies, Paisley, Strathclyde, UK) and were cultured as previously described [Ray et al. 1999, 1998). The MMTV-luc plasmid was used. The full-length rat GR expression vector was transfected with the TAT3-luc reporter plasmids, and combinations of 1 μg pcDNA3 GRα, 1 μg p6 RGR, 1 μg p6 RGRN525, 1 μg p6 RGR407C. The total amount of input DNA was kept constant by using either the pcDNA3 (to control experiments using the pcDNA3 vector), or the pVP22 (to control experiments using the VP22 vector) empty vectors. All experiments had 1 μg CMV-βGal added.

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The human T lymphoblast cell line, CEM C7A, was obtained from Dr G Brady, University of Manchester, UK, and cells were cultured in OptiMEM plus glutamax (Gibco) supplemented with 5% fetal calf serum. Cells were suspended at 10^7 cells per ml, and 0.8 ml was added to a 4-mm pathlength electroporation cuvette (BioRad, Hercules, CA, USA). DNA was added, and the cells were electroporated at 260 V and 1050 µF in an ‘Easyject1’ electroporator (Eurogentec, Seraing, Belgium). TAT3-luc (10 µg), CMV-βGal (1 µg) and either VP22 or VP22GR (5 µg) were used.

**Immunocytochemistry**

Cells were transfected with 2·5 µg expression vector in 10-cm tissue culture dishes and then plated onto multi-chamber slides for immunoperoxidase detection, or onto sterilised glass cover slips for immunofluorescence.

**Immunoperoxidase** Cells were fixed in 100% methanol at −20 °C for 20 min. The C terminal anti-GR antibody P20 (Santa Cruz Biotech, Santa Cruz, CA, USA) was used at 1:100 dilution in PBS+1·5% blocking serum, and the immunoperoxidase detection used the Vectastain kit (Vector Laboratories, Burlingame, CA, USA) according to the manufacturer’s suggestion. The monoclonal anti-myc antibody was purchased from Sigma (Poole, Dorset, UK), and was used at 1:500 dilution. Again detection was with the Vectastain kit.

**Immunofluorescence** Cells were plated onto glass cover slips and fixed in methanol. Cells were incubated with anti-myc 1:750 in PBS/0·2% BSA/4% goat serum, then goat anti-mouse conjugated to Texas Red 1:1000. Cover slips were mounted in 90% glycerol/10% PBS, and sealed with nail varnish.

**Immunoblotting**

Whole cell lysates were prepared using a non-ionic lysis buffer consisting of 1% Triton X-100, 150 mM NaCl, 10 mM Tris (pH 7·4), 1 mM EDTA, 0·2 mM sodium vanadate, 0·2 mM phenylmethylsulphonyl fluoride, 0·5% NP40. Protein concentration was estimated using the BioRad assay according to the manufacturer’s protocol with BSA as the standard. Twenty micrograms protein were resolved on a 7·5% SDS-PAGE gel and transferred to a PVDF (Millipore P, Millipore, Watford, UK) membrane overnight. The membrane was blocked with 5% non-fat dry milk and probed with anti-myc at 1:1000 dilution in blocking buffer. The membrane was washed in Tris-buffered saline with 0·1% Tween 20, and the secondary antibody, anti mouse-hrp (Amersham) used at 1:2000 dilution in blocking buffer. Detection was by enhanced chemiluminescence (ECL) as previously described (Ray et al. 1996b).

**Statistics**

Comparisons were made using analysis of variance.
Results

**VP22GR has dominant negative activity**

The VP22GR did not activate the MMTV-luc reporter in COS 7 cells (Fig. 1a), but did inhibit wild-type GR function at high concentrations of dexamethasone (Fig. 1a). Expression of VP22GR significantly inhibited MMTV-luc induction in the human pulmonary epithelial cell line A549, which contains endogenous GR, at concentrations of dexamethasone greater than 0.1 nM (Fig. 1b). In contrast, expression of the full-length, wild-type GR increased glucocorticoid sensitivity (Fig. 1b). Transfection of the empty VP22 vector did not alter the dexamethasone induction of reporter gene activity.

These experiments were repeated using a less complex glucocorticoid reporter gene, TAT3-luc. VP22GR did not allow dexamethasone induction of the reporter gene in COS 7 cells, in contrast to wild-type GR (data not shown). We found that concentrations of dexamethasone greater than 1 nM caused activation of the TAT3-luc reporter in COS 7 cells in the absence of exogenous GR, probably due to low-level expression of endogenous GR (data not shown). Therefore, dexamethasone
Figure 3 VP22GR inhibits both N and C terminal activation domains of the GR, and requires intact dimerisation and ligand binding functions. (a) COS 7 cells were transfected with 3 μg TAT3-luc, 1 μg CMV-βGal, and with either 1 μg RGR407C (407C) or RGRN525 (N525), and with either 1 μg VP22 (VP22) or VP22GR (VP22GR). Cells were treated overnight with the indicated concentrations of dexamethasone before harvest and analysis. Results are expressed as corrected relative light units (RLU), means ± standard deviation. *P<0.05 compared with the VP22 transfected control at the same concentration of ligand. A representative experiment, performed in triplicate, and repeated on three occasions.

(b) COS 7 cells were transfected with 3 μg TAT3-luc, 1 μg CMV-βGal, 1 μg wtGR, and with 1 μg VP22, 1 μg VP22GR, or 1 μg VP22GR(-LBD). (c) COS 7 cells were transfected with 3 μg TAT3-luc, 1 μg CMV-βGal, 1 μg wtGR, and with 1 μg VP22, 1 μg VP22GR, or 1 μg VP22 GRdim. Cells were treated for 16 h with dexamethasone, and harvested as described. Results are expressed as fold induction of the corrected RLU ± standard deviation. *P<0.05 compared with the VP22 transfected control treated at the same concentration of dexamethasone. A representative experiment performed in triplicate and repeated five times.
concentrations were restricted to 1 nM in COS 7 cells transfected with the TAT3-luc reporter.

VP22GR showed dominant negative activity on wild-type GR in COS 7 cells (Fig. 2a). This effect was also seen using the TAT3-luc reporter in three human cell lines which express endogenous GR: A549, CEM C7A and HeLa cells (Fig. 2b,c,d). Dexamethasone at less than 10 nM did not activate TAT3-luc in CEM C7A cells.

**VP22GR and GR interactions: structural requirements**

VP22GR inhibited GR transactivation, and so GR molecules with either the N terminal AF-1 deleted (RGR-407C), or the C terminal ligand binding domain and AF-2 deleted (RGRN525) were also examined. Both these deleted constructs are derived from the full-length, rat GR expression vector (RGR), which was inhibited by VP22GR in COS 7 cells to the same extent as the human GR (data not shown). The TAT3-luc reporter was used to analyse the function of the two deletants because MMTV-luc was not regulated by either.

RGRN525 was more active than RGR-407C at 1 nM dexamethasone, as expected, at a sub-saturating concentration of ligand (Fig. 3a). Expression of the VP22GR resulted in a dexamethasone-dependent inhibition of both RGRN525 and RGR-407C activity (Fig. 3a).

In contrast to VP22GR, VP22GR(-LBD) had no dominant negative activity (Fig. 3b), and neither did a dimerisation-deficient variant (VP22 GRdim) (Fig. 3c).

**Response element specificity of VP22GR effects**

The VP22GR opposed hydrocortisone activation of the MMTV-luc reporter in mineralocorticoid receptor transfected cells (Fig. 4a), but did not antagonise the activity of the estrogen receptor on an ERE-luc construct (Fig. 4b). Interestingly inhibition of MR function was consistently greater at higher ligand concentrations, an effect also seen with inhibition of GR transactivation (Figs 1a, 2a, 2d, 3).

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**Figure 4** DNA template specificity for dominant negative action of VP22GR. (a) COS 7 cells were co-transfected with 3 μg MMTV-luc, 1 μg CMV-MR (MR), 1 μg CMV-βGal, and with either 1 μg VP22 (VP22) or 1 μg VP22GR (VP22GR). Cells were divided and treated with the indicated concentrations of hydrocortisone (Hc) overnight before harvest as previously described. Results are expressed as fold induction of corrected relative light units ± standard deviation. *P<0.05 compared with VP22 transfected control cells at the same concentration of ligand. (b) COS 7 cells were co-transfected with 3 μg ERE-luc, 1 μg CMV-ER (ER), 1 μg CMV-βGal, and with either 1 μg VP22 (VP22), or 1 μg VP22GR (VP22GR). Cells were divided and treated with the indicated concentrations of estradiol (E2) overnight before harvest as previously described. Results are expressed as fold induction of corrected relative light units ± standard deviation. VP22GR did not differ significantly from VP22. A representative experiment, performed in triplicate and repeated on three occasions.
Dexamethasone for the GR (Fig. 5a,b). We also examined the function of a VP22 GRmyc, which differed in having a C terminal epitope tag. This did not alter its anti-RelA activity (Fig. 5b) or its dominant negative activity on GR transactivation (data not shown).

VP22GR not only conferred glucocorticoid regulation of RelA function on GR-deficient cells (Fig. 5), but it also enhanced glucocorticoid repression of RelA function in cells with endogenous GR function, A549 (Fig. 6). Exogenous GR allowed significant repression at 1 nM dexamethasone, but the VP22GR required 10 nM dexamethasone before significant repression was seen. The inhibition with dexamethasone in vector-transfected cells did not reach significance under these conditions. There was no ligand-independent reduction in NRE-luc activity.

**VP22GR intracellular localisation**

Addition of the C terminal myc epitope facilitated detection of the VP22GR fusion protein, allowed discrimination between it and GR, and did not alter protein function for dominant negative or transrepressive activity. Western blotting revealed a major protein species at 120 kDa, the predicted mass of the VP22GR fusion (data not shown). Immunocytochemistry showed that the VP22GR was located within the cytoplasm of transfected COS 7 cells (Fig. 7a,c). Dexamethasone induced almost complete nuclear localisation of the VP22 GRmyc (Fig. 7b,d).

To compare the intracellular distribution of VP22GR with wild-type GR we used an N terminal green fluorescent protein (GFP) fused to the murine wtGR. This molecule has previously been shown to retain ligand-dependent transactivation function, and to undergo appropriate nuclear translocation following ligand exposure. Both the VP22 GRmyc (Fig. 8a) and GFPGR (Fig. 8c) show cytoplasmic distribution, and both undergo nuclear translocation with the characteristic nucleolar exclusion pattern. The pattern of staining is indistinguishable between the cells expressing the GFPGR and those expressing the VP22 GRmyc (Fig. 8b,d and e). Some GFPGR transfected cells showed perinuclear accumulation of VP22 GRmyc (Fig. 8d), suggesting that cell-to-cell transfer of VP22GR had occurred. COS 7 cells were also co-transfected with VP22 GRmyc and GFPGR. The two proteins showed complete overlap of nuclear distribution after dexamethasone (Fig. 8f).

**Discussion**

The therapeutically useful, anti-inflammatory actions of glucocorticoids result from inhibition of other transcription factor function, especially NFκB (Cato & Wade 1996, Heck et al. 1997). Problems with glucocorticoid therapy include an indiscriminate range of actions, and acquired glucocorticoid resistance. If the glucocorticoid sensitivity within a focus of inflammation could be enhanced then low, ‘safe’, doses of glucocorticoid may suffice as effective therapy.

For the glucocorticoid receptor, an intracellular protein, to be useful it would need to be expressed and active in the majority of cells in its targeted area. For this reason we evaluated VP22 as an intercellular transporter for GR protein. Previous work using other VP22 fusions had shown retained cargo protein function with either GFP...
(Elliott & O’Hare 1999) or p53 (Phelan et al. 1998), supporting the idea that GR function might be retained in a VP22GR fusion protein. Furthermore, the VP22 appeared to direct nuclear localisation (Elliott & O’Hare 1997, 1999), and thereby might alter GR function by targeting unliganded GR to the nucleus.

We initially made an almost full-length GR fusion, deleting the first 41 amino acids which do not contribute significantly to protein function (Hollenberg & Evans 1988, Scheinman et al. 1995b, Iniguez et al. 1997). Despite retaining both AF-1 and AF-2, the resulting VP22GR was unable to transactivate simple reporter genes. Not only did VP22GR lack the expected ligand-dependent transactivation of MMTV-luc and TAT3-luc, but surprisingly it exhibited dominant negative activity on endogenous GR in HeLa, A549, and CEM C7A cells. VP22GR also inhibited transfected wild-type GR in COS 7 cells (Fig. 2). The GR N terminal has been analysed by deletion and alanine scanning approaches, and more recently by mutagenesis studies in yeast (Hollenberg et al. 1987, Hollenberg & Evans 1988, Iniguez et al. 1997, Iniguez & Pearce 2000). The domain from amino acid 77 to 262 contains a major transactivation function, a c-jun interacting function, and a synergy limiting activity (Iniguez et al. 1997, Hittelman et al. 1999, Iniguez & Pearce 2000). The N terminal activation domain may interact with the C terminal activation domain to synergistically enhance transcription on some DNA templates (Hollenberg & Evans 1988, Hittelman et al. 1999). However, no function has been ascribed to the N terminal 42 amino acids, and therefore it is unlikely that deletion of these residues, in isolation, would be sufficient to abolish transactivation activity of the GR. Previously, N terminal GR fusion proteins, e.g. with GFP (27 kDa), had been shown to retain transactivation potential (Htun et al. 1996, Galigniana et al. 1999). Furthermore, fusion of GFP had been performed not only to the extreme N terminus of GR, but also up to amino acid 132 without abrogating transactivation activity (Ogawa et al. 1995). This suggests that N terminal fusions up to amino acid 132 are not sufficient to abolish the transactivation function of the GR.

The VP22GR, or a variant with a C terminal myc epitope tag, was effective at inhibiting transactivation both by a C terminal truncated, constitutively active GR (RGRN525) and also by an N terminal deleted GR (RGR407C). Dominant negative activity could be abolished by disrupting either the C terminal of the VP22GR (VP22GR(-LBD)), or dimerisation function (VP22 GRdim). As the dominant negative function of the VP22GR appeared to be dependent on DNA binding, we examined the specificity of DNA response element sequence. The MR shares both DNA element sequence

Figure 6 VP22GR enhances repression of NFκB in A549 cells. A549 cells were transfected with 3 μg NRE-luc, 1 μg CMV-βGal, and with 1 μg VP22, 1 μg wtGR, or 1 μg VP22GR. Cells were then divided and incubated with recombinant human TNFα and dexamethasone (Dex) at the indicated concentrations for 16 h before harvest. Results are expressed as corrected relative light units (RLU), means ± standard deviation. *P<0.05 compared with TNFα treatment alone, the induction with TNFα was highly significant. A representative experiment performed in triplicate, and repeated on eight occasions.
and ligand with the GR; it also has a higher affinity for hydrocortisone. VP22GR inhibited transactivation by the MR, but did not significantly inhibit the activity of the ER. This suggests that the VP22GR is inhibiting GR function specifically, and that it does so by acting through its specific DNA response element. Furthermore, such inhibition may require either homodimerisation of VP22GR or heterodimerisation between the VP22GR and GR.

As the VP22GR molecule had such an unexpected profile of activity on transactivation end-points, GR inhibition of NFκB activity was measured. We used the well-characterised NRE-luc reporter gene driven by overexpressed p65 (Ray et al. 1999). Surprisingly, and in contrast to the transactivation results, the VP22GR and the VP22 GRmyc were just as effective at inhibiting the p65 function as the wild-type GR. However, there was a consistent right-shift in the dose–response of this effect with both the VP22GR and the VP22 GRmyc compared with the wtGR. The VP22GR fusion proteins did not repress RelA function at less than 1 nM dexamethasone, in contrast to wild-type GR. We also examined VP22GR function in a cell line expressing endogenous GR, A549. In these cells the NRE reporter gene was activated by...
tumour necrosis factor-α (TNFα). Both wild-type GR and VP22GR were capable of significantly repressing the NRE reporter gene.

Previous work has shown that mutation to the DNA binding domain of the GR, or to the ligand binding domain can result in impaired, but not abolished transactivation (Heck et al. 1997, Ray et al. 1999). However, no changes to the N terminal of the GR have been reported to result in such dissociation of repression from transactivation, and certainly no changes have been reported to confer dominant negative activity on transactivation.

The GR is known to interact with a wide range of proteins. To transactivate target genes, co-activator molecules are recruited to the DNA bound GR, and these serve to ‘bridge’ a signal to the basal transcriptional machinery (Collingwood et al. 1999, McKenna et al. 1999). The progesterone receptor (PR) and estrogen receptor (ER) are also capable of recruiting co-repressor proteins when activated by antagonist ligands and thereby inhibiting target gene expression (Laherty et al. 1998, Lavinsky et al. 1998). The GR is proposed to recruit a co-repressor when ‘tethered’ to an NFKB response element by RelA, and this is thought to be the mechanism of GR repression of NFKB activity (Nissen & Yamamoto 2000).

It is likely that the VP22 portion of the VP22GR is responsible for modifying the activity of the GR, generating a dominant negative molecule for transactivation. However, the different mode of GR action to oppose p65 is not disturbed by the presence of the N terminal VP22. Either competition for DNA binding, and failure to recruit co-activators, or recruitment of additional factors to the VP22 portion of the VP22GR fusion and so disruption of normal complex formation would explain the dominant negative action of VP22GR on GR transactivation. The preservation of transrepressing activity seen with VP22GR suggests that this activity requires a substantially different pattern of protein–protein interaction, and that this is not disturbed by the presence of the VP22 sequence, nor any accessory proteins recruited by it. This further underlines the separation of the two pathways, and is the first example of an N terminal GR modification capable of dissociating them so completely. The minor right-shift in dose–response consistently seen with VP22GR suggests a minor reduction in ligand binding affinity, possibly by altering chaperone protein interactions in the cytosol.

As VP22 had variously been described to target cargo proteins to the cytoplasm or nucleus of transfected cells, and also appeared to interact directly with the tubulin architecture of the cells (Elliott & O’Hare 1997), we wished to determine where the VP22GR was within the cell, and how it responded to ligand treatment. In the unliganded state the VP22 GR:myc was found exclusively in the cytoplasm where its distribution was reticular. Following ligand addition, the molecule underwent almost complete nuclear translocation, to result in a homogeneous distribution with characteristic lacunae, probably nucleoli, as previously described for the GR (Htun et al. 1996). After dexamethasone treatment, the pattern of intranuclear distribution was identical for both the GFPGR and the VP22 GR:myc.

In summary, we describe the impact of N terminal fusion of the GR with the herpes viral protein, VP22. This fusion results in a protein with a novel spectrum of ligand-dependent activity. The striking finding that the fusion retained the therapeutically useful anti NFKB activity while acquiring a powerful, ligand-dependent dominant negative action on GR mediated transactivation suggests that N terminal modification of the GR may be a novel manner in which to alter the specificity of GR actions. Previous attempts to focus GR actions have resulted in selective loss of transactivation, to a greater or lesser extent (Heck et al. 1997, Reichardt et al. 1998, Ray et al. 1999). As yet, no modification has resulted in such an extreme dissociation of the two modes of GR action as this N terminal fusion. This suggests a means to alter GR function by generating N terminal fusion proteins. Moreover, such ‘designer’ proteins may find a use in gene therapy applications.

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