Acquisition of Agonistic Properties of Nonsteroidal Antiandrogens after Treatment with Oncostatin M in Prostate Cancer Cells

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ABSTRACT

Purpose: Interleukin-6 (IL-6), a proinflammatory cytokine, activates the androgen receptor (AR) in a ligand-independent and synergistic manner. Oncostatin M (OSM) is an IL-6-type cytokine that regulates the growth of prostate cancer cells in a paracrine fashion. The present study was designed to investigate the regulation of AR expression and function by OSM, as well as the efficacy of the nonsteroidal antiandrogens hydroxyflutamide and bicalutamide in the inhibition of AR-mediated signal transduction.

Experimental Design: Expression of the OSM receptor-β in the prostate cancer cell lines LNCaP, PC-3, and DU-145 was investigated by reverse transcription-PCR. DU-145 and PC-3 cells were cotransfected with an androgen-responsive gene and AR cDNA. Reporter gene activity was measured after treatment with androgen and/or OSM in the absence or presence of antiandrogens or protein kinase inhibitors. AR expression after OSM treatment was assessed by Western blot.

Results: OSM receptor-β expression was higher in DU-145 and PC-3 than in LNCaP cells. OSM caused ligand-independent activation of the AR in DU-145 cells, and the maximal activation was 62% of that induced by the synthetic androgen methyltrienolone. In the presence of OSM, hydroxyflutamide behaved as an AR agonist. Bicalutamide down-regulated AR activation caused by OSM only at a concentration of 1 μM. The inhibitor of the protein kinase A signaling pathway PKI and Δn signal transducers and activators of transcription (STAT) 3 showed no effect on AR activation by OSM. The inhibitor of the MAPK pathway, PD 98059, caused only a minor down-regulation of OSM-induced reporter gene activity. OSM did not change AR expression in DU-145 cells transfected with AR cDNA.

Conclusions: OSM is a member of the IL-6 family of cytokines, which causes ligand-independent activation of the AR without altering receptor expression. In contrast to AR activation by IL-6, nonsteroidal AR antagonists act as agonists in the presence of OSM. This may be attributable to recruitment of different intermediary signal transduction proteins by OSM and IL-6, respectively. The acquisition of agonistic properties of AR blockers in the presence of OSM might compromise use of these drugs in prostate cancer treatment.

INTRODUCTION

Prostate cancer is the most commonly diagnosed neoplasm in men in industrialized countries. Despite the earlier diagnosis, prostate tumor incidence is increasing. Prostate cancer growth in early and intermediate stages depends on androgenic stimulation and endocrine therapy is, therefore, aimed at the inhibition of androgen signaling. Unfortunately, tumor progression, which could be detected by increased levels of serum prostate-specific antigen, occurs in most patients. The AR plays a central role in the regulation of proliferation, differentiation, and apoptosis of prostate cancer cells. The AR is up-regulated in therapy-resistant prostate cancer, as demonstrated in immunohistochemical studies and real-time PCR experiments (1, 2). The AR can be activated, even in the absence of androgens, by substances that increase intracellular cAMP levels or by growth factor receptor-mediated events (3, 4). AR transactivation potential may be increased through the activation of MAPK, as evidenced in cases of receptor stimulation by HER-2/neu (5).

There is increasing evidence for the existence of a cross-talk between the signaling pathways of IL-6 and the AR in prostate cancer cells (6–9). For several reasons, interaction between the two signaling pathways is clinically significant: (a) serum IL-6 levels are elevated in patients suffering from metastatic carcinoma of the prostate (10, 11); (b) IL-6 and its receptor were detected in prostate cancer cells on immunohistochemistry (12); (c) IL-6 concentration is increased even in tissues from patients with clinically localized prostate cancer (13); and (d) IL-6 exhibits different effects on prostate cancer cell lines, varying from the induction of G1 growth arrest to the inhibition of apoptosis (14, 15). There is limited information available on biological effects of other IL-6-type cytokines.
MATERIALS AND METHODS

Cell Culture. LNCaP, PC-3, and DU-145 cells were grown in RPMI 1640 (Hy-Clone, Logan, UT) supplemented with 10% FCS and penicillin/streptomycin (PAA Laboratories, Linz, Austria) and incubated at 37°C in a 5% CO2 atmosphere.

Chemicals. Human recombinant OSM, the cAMP analogue db cAMP, and the PKA inhibitor PKI were purchased from Sigma (Deisenhofen, Germany). The nonsteroidal antiandrogen methyltrienolone (R1881) was from New England Nuclear (Dreieichenhain, Germany). The nonsteroidal antiandrogen hydroxyflutamide was from Essex Pharma (Munich, Germany), and bicalutamide was synthesized at Schering AG (Berlin, Germany). The MAPK pathway inhibitor PD 98059 was provided by Merck Eurolab (Freiburg, Germany).

RT-PCR. Total RNA from hormone-refractory (DU-145 and PC-3) and hormone-sensitive (LNCaP) prostate cancer cell lines was isolated by the SV RNA Isolation System (Promega, Madison, WI). Five hundred ng of total RNA were used for the first-strand cDNA synthesis using SuperscriptII RNase H− Reverse Transcriptase (Life Technologies, Inc. Rockville, MD). cDNA fragments of the OSMRβ and β2-microglobulin (internal control) were amplified using the following primers: OSMRβ sense (5’-CCCTGAAACACAGACGCTGCTC-3’), OSMRβ antisense (5’-GTCTTGGGCTCTAAGTACTTT-3’), β2-microglobulin sense (5’-ATGCCGTGGTGAACCAGTCT-3’), and β2-microglobulin antisense (5’-AGACTGGACACCACCAGTCT-3’; GenXpress, Vienna, Austria). For the amplification of the OSMRβ, 4 µl of cDNA solution were added to 46 µl of PCR mix containing 10× buffer (Finnzyme, Vienna, Austria; 200 nM dNTPs, 0.5 µmol of the respective oligonucleotide, 2% DMSO, and 2 units of Dynazyme II DNA polymerase). The same protocol was followed for the amplification of β2-microglobulin cDNA, with the distinction that 2 µl of cDNA solution and DMSO, at a final concentration of 4%, were added. PCR was performed in the GeneAmp PCR 9700 System thermocycler (PE Biosystems, Norwalk, CT) according to the following programs: (a) for the amplification of the OSMRβ gene: 3 min at 94°C; 33 cycles of 45 s at 94°C, 10 s at 96°C, 1 min and 5 s at 57°C, and 1 min and 20 s at 73°C; with a final extension time of 3 min at 73°C; and (b) for the amplification of the β2-microglobulin gene: 2 min at 94°C; 30 cycles of 25 s at 94°C, 10 s at 96°C, 1 min at 57°C, and 2 min and 15 s at 73°C; and a final extension for 2 min at 73°C. The fragments were visualized after electrophoresis in a 2% agarose gel in TAE buffer [0.04 mM Tris, 0.001 mM EDTA (pH 8.0)] prestained with ethidium bromide.

Transfections and CAT Assays. Transient transfection experiments by lipofection in DU-145 and PC-3 cells used the reporter plasmid ARE2-TATA-CAT and AR cDNA and were performed as described previously (6). The dn STAT3 plasmid REF STAT3CYF was kindly provided by Dr. Thomas Welte (Department of Pathology, Yale University, New Haven, CT). Reporter gene activity was measured after a 24-h incubation with OSM and/or androgen in the absence or presence of antiandrogens or PK inhibitors by CAT assay.

Western Blot. AR expression was investigated in transfected DU-145 cells and in controls. Cells were lysed as described elsewhere (17), and electrophoresis and Western blotting were performed using the NuPAGE electrophoresis system (Invitrogen, Leek, the Netherlands). For AR expression, the mouse monoclonal anti-AR antibody F39.4.1 (Biogenex, San Ramon, CA) at a dilution 1:1000 and the secondary antirabbit horseradish peroxidase-linked antibody (1:1000 dilution) were used. A mouse anti-actin monoclonal antibody from Chemicon (Harrow, United Kingdom) was used at a dilution 1:1000. Signal was detected by the enhanced chemiluminescence detection system (Amersham Pharmacia Biotech, Freiburg, Germany).

Statistics. The Mann-Whitney t test was used for assessment of statistical significance.

RESULTS

Expression of the OSMRβ in Prostate Cancer Cell Lines. Before performing transfections of the androgen-responsive reporter gene, expression of the specific receptor OSMRβ was investigated in the prostate cancer cell lines LNCaP, PC-3, and DU-145 by RT-PCR. For OSMRβ expression studies, RT-PCR has been chosen rather than Western analysis because previous reports showed that commercially available antibodies could not be used for immunoblotting (18). The 743-bp OSMRβ fragment could be detected in all three of the cell lines. However, OSMRβ expression was much lower in the androgen-sensitive cell line LNCaP than in DU-145 or PC-3 cells (Fig. 1).

Activation of the AR by OSM in the Absence or Presence of Nonsteroidal Antiandrogens. To overcome difficulties that might occur because of the reduced OSMRβ expression in LNCaP cells, transfections were carried out in DU-145 and PC-3 cells with wild-type AR cDNA. Treatment of transfected DU-145 cells with OSM resulted in ligand-independent and
dose-dependent activation of the AR (Fig. 2). The maximal activation of the AR by OSM was 62% of the activity measured after treatment with the synthetic androgen R1881 (1 nM). Low doses of R1881 and OSM up-regulated the transcriptional activity of the AR in an additive manner. In contrast to DU-145 cells, there was no induction of reporter gene activity by OSM in the PC-3 cell line (data not shown). In previous publications, antiandrogens were shown to inhibit AR-mediated reporter gene activity induced by the related cytokine IL-6 (6, 8). To study the ability of clinically used antiandrogens to block OSM-induced activation of the AR, DU-145 cells were incubated with OSM in the absence or presence of either hydroxyflutamide or bicalutamide. Surprisingly, hydroxyflutamide enhanced the OSM-induced AR activation, whereas bicalutamide down-regulated the ligand-independent activation, only when administered at a concentration of 1 μM (Fig. 3). When bicalutamide was added at a concentration of 5 μM, OSM-induced AR activity either slightly increased (with 10 ng/ml OSM) or remained unchanged (with 50 ng/ml OSM; Fig. 3). As expected, both AR antagonists suppressed AR activity caused by R1881 (Fig. 3). Thus, action of commonly used antiandrogens substantially differs in the presence of IL-6 or OSM.

Effect of PK Inhibitors on AR Activation. Previous studies revealed that ligand-independent activation of the AR depends on the functionality of various kinase pathways (3, 6). To investigate the involvement of the PKA pathway in the up-regulation of activity of the AR by OSM, the cells were treated with the PKA inhibitor PKI. Interestingly, PKI had no influence on AR activation exerted by OSM, whereas it down-regulated the db cAMP effect on AR activation (Fig. 4). Similar to those observations, the selective inhibitor of the extracellular signal-regulated kinase 1/2 kinases, PD 98059, caused only a minor (less than 25%) inhibition of OSM-induced reporter gene activity (Fig. 5). Control experiments confirmed that AR activation by EGF could be diminished by PD 98059. In addition, administration of dn STAT3 did not counteract AR activation by OSM (data not shown).

Expression of AR in OSM-treated DU-145 Cells. Nonsteroidal activators of the AR might also up-regulate receptor expression, as demonstrated with IL-6 in LNCaP cells (8, 19). To investigate a possible effect of OSM on AR protein levels, Western blots for the detection of the AR were performed.
characteristic features of which are a decline of serum prostate-responsible for the antiandrogen withdrawal syndrome, the char-
27, 28). Each of these mechanisms may be, at least in part,
cases of receptor activation by HER-2/neu or phorbol ester (5,
24); (b) bicalutamide becomes an agonist after either prolonged
mutant AR are commonly activated by hydroxyflutamide (23,
motor of AR expression only at lower concentrations. In prostate cancer, the
presence of OSM and that bicalutamide causes inhibition of AR
activity only at lower concentrations. In prostate cancer, the
expression was analyzed in nontransfected, transfected un-
treated, and OSM-treated DU-145 cells. As expected, the AR
was not detected in nontransfected DU-145 cells. AR expression
was identical in transfected untreated and OSM-treated cells (Fig. 6). Thus, the increase in reporter gene activity caused by
OSM is solely attributable to an increase in AR activity and not
to up-regulation of AR expression.

DISCUSSION

OSM, which is a member of the IL-6 type of cytokines, affects the growth of prostate cells in a paracrine manner (16, 20). This is different from IL-6, for which both autocrine and paracrine loops were reported in the prostate (12, 19). It is well established that prostate cancer cells PC-3 and DU-145 express IL-6 (20–22). Those findings were supported by studies carried out on clinical specimens (12, 13). In contrast, OSM was not detected in conditioned media from the cell lines LNCaP ATCC, LNCaP GW that acquires a p53 mutation, PC-3, and DU-145 by ELISA (20, 21). On the basis of our results and those of
previous reports, we conclude that both IL-6 and OSM activate the AR (6–8). However, the unexpected findings of the present
study are that hydroxyflutamide acts as an AR agonist in the presence of OSM and that bicalutamide causes inhibition of AR
activity only at lower concentrations. In prostate cancer, the
switch of nonsteroidal antiandrogens to agonists was reported in
various clinical situations and in in vitro and in vivo models: a)
motor AR are commonly activated by hydroxyflutamide (23, 24); (b) bicalutamide becomes an agonist after either prolonged
steroid depletion or treatment with tumor necrosis factor-α (25,
26); and (c) reduced efficacy of AR antagonists was reported in
cases of receptor activation by HER-2/neu or phorbol ester (5,
27, 28). Each of these mechanisms may be, at least in part,
responsible for the antiandrogen withdrawal syndrome, the char-
acteristic features of which are a decline of serum prostate-

specific antigen levels and an improvement of clinical symp-
toms after the cessation of antiandrogen administration (29).

Although the signaling of IL-6 and OSM is similar in many
cell types, some differences in structural organization of their
receptors and in the recruitment of adapter proteins occur. On
activation, the IL-6 receptor is composed of two gp130 signal
transducing subunits and an IL-6 receptor α subunit, whereas, in
case of the OSMR, the α subunit is absent and one gp130
subunit couples to either OSMR or LIF receptor β. Recently,
differences in signal transduction between gp130 and the OSMR
were revealed: after receptor activation, the adapter protein
Shc binds only to the OSMR (and not to gp130), whereas
the tyrosine phosphatase SHP-2 binds to gp130 (and not to
the OSMR). Ref. 30). Whereas SHP-2 provides a link to the
Ras/Raf/MAPK pathway, Shc, which is recruited by OSM,
can mediate both Ras-dependent and Ras-independent re-
sponses. Therefore, the stronger effects of OSM reported in
various cell lines are not surprising.

We have confirmed the previous finding that the OSMRβ
is expressed in the androgen-insensitive prostate cancer cell
lines DU-145 and PC-3 (16). In a report by Mori et al. (16), the
OSMRβ was not detectable in the LNCaP cell line. However,
we have observed that expression of the OSMR receptor is
weaker in LNCaP than in DU-145 or PC-3 cells. These varia-
tions may reflect subtle differences in cell culture conditions and
RT-PCR procedures.

Another important difference between IL-6 and OSM is
that with the latter cytokine, inhibitors of neither PK nor dn
STAT3 substantially antagonize AR activation. These data
could be interpreted in two ways: (a) that inhibition of a single
transduction pathway is compensated by other signal transduc-
tion cascades; or (b) that the effect of OSM depends on stimu-
lation of other kinases. It is well established that the functional
activity of the AR can be stimulated in a ligand-independent
manner by activators of the PKA pathway, like forskolin (3, 31,
Weigel (3) showed that, in CV-1 and PC-3 cells, forskolin activated AR function without changing receptor levels. In summary, agonistic behavior of AR antagonists is a particularly interesting aspect of AR activation by OSM. Acquisition of agonistic properties of nonsteroidal antiandrogens in tumor cells treated with OSM may compromise their use in prostate cancer treatment. In this context, there are two possibilities for therapeutic intervention; therapy concepts aimed to interfere with signaling of the IL-6 type of cytokines or to down-regulate AR expression should be worked out (17).

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REFERENCES

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