mPGES-1 in prostate cancer controls stemness and amplifies epidermal growth factor receptor-driven oncogenicity

Federica Finetti1, Erika Terzuoli1, Antonio Giachetti1, Raffaella Santi2, Donata Villari3, Hiromi Hanaka4, Olof Radmark4, Marina Ziche1,5 and Sandra Donnini1,5

1Department of Life Sciences, University of Siena, Via Aldo Moro 2, 53100 Siena, Italy
2Department of Surgery and Translational Medicine, University of Florence, Largo Brambilla 3, 50134 Firenze, Italy
3Department of Clinical and Experimental Medicine, University of Florence, Viale Pieraccini 18, 50139 Firenze, Italy
4Department of Medical Biochemistry and Biophysics, Karolinska Institutet, SE-171 77 Stockholm, Sweden
5Istituto Toscano Tumori (ITT), Firenze, Italy

Abstract

There is evidence that an inflammatory microenvironment is associated with the development and progression of prostate cancer (PCA), although the determinants of intrinsic inflammation in PCA cells are not completely understood. Here we investigated whether expression of intrinsic microsomal PGE synthase-1 (mPGES-1) enhanced aggressiveness of PCA cells and might be critical for epidermal growth factor receptor (EGFR)-mediated tumour progression. In PCA, overexpression of EGFR promotes metastatic invasion and correlates with a high Gleason score, while prostaglandin E2 (PGE2) has been reported to modulate oncogenic EGFR-driven oncogenicity. Immunohistochemical studies revealed that mPGES-1 in human prostate tissues is correlated with EGFR expression in advanced tumours. In DU145 and PC-3 cell lines expressing mPGES-1 (mPGES-1SC cells), we demonstrate that silencing or ‘knock down’ of mPGES-1 (mPGES-1KD) or pharmacological inhibition by MF63 strongly attenuates overall oncogenic drive. Indeed, mPGES-1SC cells express stem-cell-like features (high CD44, β1-integrin, Nanog and Oct4 and low CD24 and α6-integrin) as well as mesenchymal transition markers (high vimentin, high fibronectin, low E-cadherin). They also show increased capacity to survive irrespective of anchorage condition, and overexpress EGFR compared to mPGES-1KD cells. mPGES-1 expression correlates with increased in vivo tumour growth and metastasis. Although EGFR inhibition reduces mPGES-1SC and mPGES-1KD cell xenograft tumour growth, we show that mPGES-1/PGE2 signalling sensitizes tumour cells to EGFR inhibitors. We propose mPGES-1 as a possible new marker of tumour aggressiveness in PCA.

Key Words

► mPGES-1
► prostate cancer
► stemness
► EGFR
► EMT

Introduction

Prostate cancer (PCA) is currently treated with androgen deprivation and chemotherapeutic agents. Resistance to chemotherapy, a mounting issue in clinical oncology due to its association with tumour recurrence, has hastened the search for new prognostic biomarkers and new therapeutic targets aimed at patient stratification in relation to treatment (Schrecengost & Knudsen 2013, Augello et al. 2014). Because ~40% of PCAs express...
epidermal growth factor receptor (EGFR), expression of which is correlated with tumour recurrence and high Gleason score, the receptor is assumed to be a potential molecular target for advanced PCa. However, clinical trials in PCa patients have shown limited efficacy of EGFR-targeted drugs (Canil et al. 2005, Hammarsten et al. 2007, Schlom et al. 2007, Gravis et al. 2008, Nabhan et al. 2009). Multiple mechanisms, such as unrestrained expression of EGFR, emergence of oncogenic mutants (KRAS, BRAF and PIK3CA) and inactivation of the PTEN tumour suppressor gene, underlie resistance to these drugs. Acquired EGFR antagonist resistance is often associated with the activation of bypass signalling pathways typically embedded in tumour cells or the surrounding tissue (Arteaga 2002, Wieduwilt & Moasser 2008, Cathomas et al. 2012, Seshacharyulu et al. 2012).

Inflammatory molecules have been shown to enhance EGFR oncogenic action in epithelial tumours, fostering their propensity to metastasize. Many studies on the link between inflammation and cancer have dealt with stimuli originating from the cancer microenvironment (extrinsic), providing strong support for this mechanism (Mantovani et al. 2008, Hanahan & Weinberg 2011, Coussens et al. 2008, 2012). Microsomal PGE synthase-1 (mPGES-1), known to be induced in inflammatory as well as tumour cells by pro-inflammatory cytokines, such as interleukin-1 beta (IL-1β) and tumour necrosis factor alpha (TNFα), contributes in a critical way to tumour progression (Jakobsson et al. 1999, van Rees et al. 2003, Takeda et al. 2004, Kamei et al. 2009, Xu et al. 2012, Sha et al. 2013, Takahashi et al. 2014). We recently found that EGFR activation up-regulates mPGES-1, which in turn promotes phosphorylation of EGFR through prostaglandin E2 (PGE2) (Donnini et al. 2012).

Here we investigated whether expression of intrinsic mPGES-1 in advanced PCa cells enhanced their aggressiveness and might be critical for EGFR-mediated tumour progression. To begin to understand the mechanism underlying enhanced prostate oncogenic drive exerted by mPGES-1, we first examined human PCa specimens obtained after prostatectomy in a group of patients showing various stages of malignancy. The close association between mPGES-1 and EGFR expression observed in these tumours led us to detailed study of this connection in experiments on hormone-independent PCa cells DU145 and PC-3 in vitro and in vivo. We demonstrate that these cells have a mesenchymal phenotype and stem-like features, which are likely to confer aggressive traits. Evidence that intrinsic mPGES-1 underpins these traits is seen in cell cultures in which ablation of the mPGES-1 gene (DU145 mPGES-1 KD) or inhibition of mPGES-1 activity prevents development of a vigorous tumorigenic phenotype. The enhanced oncogenic drive observed in vitro translated to nude mice in vivo inoculated with DU145 or PC-3 cells, as we found significantly higher tumour growth and lung metastasis formation in mice inoculated with PCa cells expressing mPGES-1. Further, blockade of EGFR in vivo with erlotinib indicated the possibility of quenching the oncogenic drive exerted by malignant cooperation of the two signals (PGE2 and EGF).

### Materials and methods

#### Tumour samples

For PCa immunohistochemical study, formalin-fixed, paraffin-embedded tissue blocks from 52 radical prostatectomy specimens were retrieved from the archives of the University of Florence (Florence, Italy). Informed consent and approval according to the Helsinki Declaration were obtained from the local ethics review board. The specimens were reviewed by two genitourinary pathologists; pathological stage and tumour grade were assigned according to the Union International Contre le Cancer (2002) classification and the Gleason score, respectively (Epstein et al. 2005, Edge et al. 2010). Twenty-five carcinomas were limited to the prostate (pT2, organ-confined PCa) and were moderately differentiated (Gleason score = 6), whereas 27 cases (advanced PCa) were non-organ-confined tumours (pT3/pT4) with a high Gleason score (≥7) (Table 1). The median age of patients was 67.7 years (range 41–78 years).

#### Immunohistochemical analysis of tumour samples

Tissue slides were deparaffinised in xylene and dehydrated in ethanol. Microwave pre-treatment in EDTA (pH 9.0) (for EGFR) or in citrate buffer (pH 6.0) (for mPGES-1 and αv-integrin) was performed for 20 min. Endogenous peroxidase activity was blocked with 3% hydrogen peroxide (v/v) for 10 min and with 3% BSA (w/v) for 30 min. The slides were incubated with primary antibodies targeting

### Table 1 Gene expression of vimentin, fibronectin and ahnak in DU145 cells expressing mPGES-1

<table>
<thead>
<tr>
<th>Gene</th>
<th>Mean ± S.D.</th>
<th>s.d.</th>
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<tbody>
<tr>
<td>Vimentin</td>
<td>12.1</td>
<td>2</td>
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<tr>
<td>Fibronectin</td>
<td>2</td>
<td>0.1</td>
</tr>
<tr>
<td>Ahnak</td>
<td>1.6</td>
<td>0.2</td>
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z6-integrin (1:100; Santa Cruz, Heidelberg, Germany), mPGES-1 (1:50; Thermo Scientific, Waltham, MA, USA) and EGFR (1:100; Cell Signalling, Leiden, The Netherlands) followed by chromogenic visualization using Immunoperoxidase Secondary Detection System kits (Chemicon, Billerica, MA, USA). In particular, sections were incubated for 15 min in the appropriate species-specific biotinylated secondary antibodies and then with streptavidin-conjugated HRP for 15 min. After incubation they were exposed to 3,3-diaminobenzidine tetrahydrochloride (Sigma) for 10 min to produce a brown reaction product. After counterstaining with hematoxylin, slides were washed thoroughly, dehydrated, cleared in xylene and mounted. Staining intensity was scored as negative (no staining) or positive (brown colour).

Cell lines

DU145 WT (passages 5–20, ATCC HTB-81, certified by STRA) is a PCa cell line with high constitutive expression of mPGES-1 (Hanaka et al. 2009). DU145 mPGES-1 knockdown (mPGES-1KD cells, passages 8–20) and non-targeted shRNA (mPGES-1SC cells, 8–20) cells were obtained and cultured as described (Hanaka et al. 2009). PC-3 WT (passages 8–20, ATCC CRL-1435, certified by STRA) and LNCaP WT (passages 5–17, ATCC CRL-1740, certified by STRA) PCa cells were from ATCC. Cells were grown in endothelial growth medium (EGM-2) (Clonetics, Lonza) and supplemented with 10% FBS (v/v). Human umbilical vein endothelial cells (HUVEC, passages 3–10) were from Lonza (Basel, Switzerland) (C2519A, certified by expression of CD31/105, vWFVIII, and positivity for acetylated low-density lipoprotein uptake). Cells were grown in RPMI (Euroclone, Pero Milano, Italy) and supplemented with 10% FBS (v/v). Human umbilical vein endothelial cells (HUVEC, passages 3–10) were from Lonza (Basel, Switzerland) (C2519A, certified by expression of CD31/105, vWFVIII, and positivity for acetylated low-density lipoprotein uptake). Cells were grown in endothelial growth medium (EGM-2) (Clonetics, Lonza) and supplemented with 10% FBS (v/v).

Transient mPGES-1 silencing

For siRNA transfection, the siRNA sequence (human mPGES-1: 5’-CGGCTAAGAATGCAGACTTT-3’) was from Qiagen. The day before transfection, cells were trypsinized and 3 × 10^5 cells were seeded in six-well plates. Transient transfection of siRNA was carried out using Lipofectamine 2000 (Invitrogen) according to the manufacturer’s instructions. Cells were assayed 72 h after transfection.

mPGES-1 shRNA transfection

Lenti vector plasmids for mPGES-1 knock down (Sigma) and mPGES-1+/+ (p Lenti vector with C-terminal Myc-DDK tag-NM_004878) were obtained from Sigma and Origene (Origene, Rockville, MD, USA), respectively. pSAX2 packaging plasmid (12260) and pMDG.2 envelope plasmid (12259) were obtained from Addgene (Cambridge, MA, USA).

All the plasmids were sequence-verified. To generate mPGES-1 knock down (mPGES-1KD) cells or mPGES-1 overexpressing (+/+ cells, 1 × 10^6 HEK293 cells (Life Technologies) were transfected with 2.25 μg of PAX2 packaging plasmid, 0.75 μg of PMD2G envelope plasmid and 3 μg of pLKO.1 hairpin vector utilizing 12 μl of Lipofectamine 2000 on 10 cm plates. Polyclonal populations of transduced cells were generated by infection with 1 multiplicity of infectious units (MOI) of lentiviral particles. Three days after infection, cells were selected with 10 μg/ml puromycin (Gibco) or 20 μg/ml neomycin/kanamycin (Sigma) for 1 week.

Epithelial-mesenchymal transition PCR array

The expression of 88 human Epithelial-mesenchymal transition (EMT) genes was profiled in DU145 cells using the EMT-RT2 Profiler PCR Arrays (SAAB Bioscience, Qiagen). Total RNA was isolated using an RNA Mini kit (Qiagen) and reverse transcribed using an RT-PCR kit (Qiagen). Expression of 88 genes using the formula 2^(-ΔDCT). Real-time PCR

Total RNA was obtained using an RNA Mini kit (Qiagen). RNA (0.5 μg) was reverse transcribed using an RT-PCR kit (Bio-Rad). Premixed primers for vimentin, fibronectin, ahnak, ITGB1, Nanog, Oct4 and GAPDH (as internal control) were from Applied Biosystems. Real-time PCR was performed using SYBR Green Supermix (Bio-Rad). RT-PCR was performed using an iCycler iQ5 PCR Detection System. The results are expressed as 2^(-ΔDCT) or fold increase.

Western blot

Tumour cells (5 × 10^5) were seeded in 6 cm plates in medium with 10% fetal bovine serum (FBS) (v/v) for 96 h, then lysed and analysed. Where indicated, cells were treated with PGE2 (1 μmol/l), erlotinib (10 μmol/l), NS398 (10 μmol/l) or [2-(6-chloro-1H-phenanthro-(9,10-d) imidazol-2-yl)]isophthalonitrile, MF63 (10 μmol/l). PGE2 and NS389 were from Sigma, erlotinib was from Santa Cruz and MF63 was from AbMole (Houston, TX, USA). To assess
translocation of β-catenin from cytosol to nucleus, cells were trypsinized and homogenized on ice in lysis buffer containing 0.1 mmol/l EGTA, 0.1 mmol/l EDTA, 10 mmol/l Heps, 10 mmol/l KCl, protease and phosphatase inhibitors. After incubation on ice for 15 min, Nonidet-P-40 was added to cell lysates, which were then centrifuged (3900g, 30 s). The supernatant contained the cytosolic fraction, while the pellet was solubilized in lysis buffer containing 1 mmol/l EGTA, 1 mmol/l EDTA, 20 mmol/l Heps, 10 mmol/l NaCl, 1% protease and phosphatase inhibitors (v/v), followed by incubation on ice for 10 min and centrifuging (5480g, 5 min). The supernatant contained the nuclear fraction. An equal amount of proteins was loaded on SDS-PAGE gel and transferred to a nitrocellulose membrane. Western blot was performed as described by Donnini et al. (2012). Sources of antibodies were: anti-vimentin and anti-fibronectin, Sigma; anti-α6 integrin and anti-β1 integrin, Santa Cruz; anti-P-Tyrosine, anti-EGFR, anti-caspase3 and anti-P-ERK1/2, Cell Signalling; anti-PGE2 synthases, anti-COX synthases, Nonidet-P-40 was added to cell lysates, which were then centrifuged (3900g, 30 s). The supernatant contained the cytosolic fraction, while the pellet was solubilized in lysis buffer containing 1 mmol/l EGTA, 1 mmol/l EDTA, 20 mmol/l Heps, 10 mmol/l NaCl, 1% protease and phosphatase inhibitors (v/v), followed by incubation on ice for 10 min and centrifuging (5480g, 5 min). The supernatant contained the nuclear fraction. An equal amount of proteins was loaded on SDS-PAGE gel and transferred to a nitrocellulose membrane. Western blot was performed as described by Donnini et al. (2012). Sources of antibodies were: anti-vimentin and anti-fibronectin, Sigma; anti-α6 integrin and anti-β1 integrin, Santa Cruz; anti-P-Tyrosine, anti-EGFR, anti-caspase3 and anti-P-ERK1/2, Cell Signalling; anti-PGE2 synthases, anti-COX synthases, anti-PGDH, anti-PGT and anti-EP receptors, Cayman Chemicals; anti-E-cadherin, DAKO (Milan, Italy). Images were digitalized with CHEMI DOC Quantity One software, blots were analysed in triplicate by densitometry using NIH Image 1.60B5 Software, and arbitrary densitometric units were normalized for β-actin (Sigma), tubulin (Santa Cruz) or H2A (Abcam, UK).

**Tumour growth and lung metastasis in immunodeficient mice**

Experiments were performed according to Italian and EEC guidelines for animal care and welfare (EEC Law No. 86/609). The experiments were approved by the Italian Ministry of Health (215/2011-B). To assess the contribution of mPGES-1 to the anti-tumour activity of erlotinib, immunodeficient mice (5-week-old male athymic mice, Harlan, Indianapolis, IN, USA) were inoculated s.c. in the right flank with 20×10⁶ DU145 cells (mPGES-1[^1] or mPGES-1[^2]). When tumours reached a volume of 70–100 mm³, the animals were randomly assigned to treatment with erlotinib (50 mg/kg, three times a week by gavage). The first treatment is reported as day 1. Serial calliper measurements of perpendicular diameters were used to calculate tumour volume in mm³ with the formula: shortest diameter×longest diameter×thickness of tumour in millimeter. After treatment, animals were sacrificed and tumours were collected and split in two parts. One part was immediately frozen in liquid nitrogen for western blot. The other part was embedded in Tissue-Tek O.C.T. (Sakura, Torrance, CA, USA) and frozen in liquid nitrogen for histology (Donnini et al. 2007). For histology, see Supplementary Data, see section on supplementary data given at the end of this article.

To assess the contribution of mPGES-1 to lung metastases, DU145 and PC-3 PCa cells (mPGES-1[^3] or mPGES-1[^4]) were suspended in PBS at a density of 20×10⁶ cells/ml, and 250 μl of suspension was injected into the tail vein. To investigate whether EGFR inhibition affected the invasive activity of mPGES-1, cells were pre-treated with erlotinib (10 μmol/l) for 72 h before injection. After 7 weeks, immunodeficient mice were sacrificed, lungs removed and fixed in Bouin’s solution and the number of metastatic colonies counted.

**MTT assay**

Cell proliferation was quantified by Vybrant MTT cell proliferation assay as described (Donnini et al. 2007). Briefly, tumour cells (5×10⁴) were seeded in 96-multiwell plates in medium with 10% serum for 24 h and then, where indicated, exposed to erlotinib (0.1–10 μmol/l) for 96 h in 10% FBS (v/v). Results are reported as 540 nm absorbance/well.

**Adhesion**

DU145 cells were maintained in 10% FBS (v/v) and then trypsinized; 5×10⁴ cells/ml in 1% FBS (v/v) medium were seeded in 96 multiwell plates coated with human fibronectin and incubated for 2 h at 37 °C. The wells were washed gently with PBS and adherent cells were fixed and stained with Diff-Quik. Adherent cells were counted by microscope in five random fields at 200×.

**Tumour-endothelium adhesion**

Tumour-endothelium adhesion was performed using the CytoSelect tumour-endothelium adhesion assay kit according to the manufacturer’s instructions (Cell Biolabs, San Diego, CA, USA). Briefly, HUVEC cells (10×10⁵ cells/well in 48 multiwell plates) were maintained in 10% FBS (v/v) for 48 h. After monolayer formation, DU145 cells were harvested and 1×10⁶ cells/ml were suspended in serum-free medium and incubated for 1 h in the presence of CytoTracker. Cells were washed twice and added to the endothelial cell monolayer. After 1 h cells were lysed and the fluorescence read at 480 nm/520 nm. Where indicated, endothelial cells were pre-treated for 4 h with TNFα to increase cell–cell adhesion (Sheski et al. 1999).
Transendothelial migration
HUVEC cells (8 × 10^5 cells/well in the filter of a 48-well transwell plate) were maintained in 10% FBS (v/v) for 48 h. After monolayer formation, DU145 cells were harvested and 1 × 10^6 cells/ml were suspended in serum-free medium and incubated for 1 h in the presence of CytoTracker. Cells were washed twice with serum-free medium and 2 × 10^5 cells were added to the upper side of the transwell plate. After 2 h the medium in the lower side of the plate was collected and centrifuged. Migrant tumour cells were lysed and fluorescence measured at 480 nm/520 nm.

Cell viability assay
Tumour cells at a density of 5 × 10^5 cells/ml were incubated for 24 or 48 h in medium with 0.1% FBS (v/v). After incubation, the numbers of dead cells stained with trypan blue and total cells were evaluated by optical microscope. The number of dead cells was reported as a percentage of total cells.

Flow cytometry
Tumour cells were harvested and 3.5 × 10^4 cells were incubated in ice for 30 min with primary antibody, then washed in PBS and exposed to secondary antibody for an additional 30 min. Surface CD44 and CD24 (Abcam) were quantified by flow cytometry using unlabelled monoclonal or polyclonal Ab followed by fluorescein isothiocyanate (FITC) or tetramethylrhodamine (TRIC)-labelled secondary antibodies. Cells were analysed by flow cytometry using a FACScan flow cytometer (Becton-Dickinson, Franklin Lakes, NJ, USA). Data was acquired by CellQuest and plotted using FlowJo (Tree Star, Ashland, OR, USA).

Clonogenic assay
Tumour cells were plated in 60 mm culture dishes (1000 cells per dish) in medium containing 10% FBS. After 24 h cells were treated with erlotinib or MF63 (1–10 μmol/l) in 10% FBS (v/v) and kept in a humidified incubator for 10 days. Colonies (> 50 cells) were fixed and stained with 0.05% crystal violet (w/v) (Sigma) in 10% ethanol (v/v), counted and photographed.

Statistical analysis
Results were expressed as means ± S.E.M., analysed by Student’s t-test and/or one-way ANOVA with Bonferroni’s correction. A value of P<0.05 was considered to denote statistical significance.

Results
mPGES-1 promotes the mesenchymal and stem cell-like phenotype in PCa cells
We investigated the correlation of mPGES-1 expression (immunohistochemistry) with staging and grading in a series of PCa cases. Overall, mPGES-1 expression was detected in 12/25 (48%) organ-confined PCa and in 21/27 (77.7%) advanced PCa (Fig. 1A). In human advanced PCa samples, both mPGES-1 and EGFR were co-expressed in a high percentage of cases (n = 19/27, 70.3%; Fig. 1A). In the same group, α6-integrin, a stem cell marker (Marthick & Dickinson 2012, Hoogland et al. 2014), was negative or weakly stained (n = 19/27; Fig. 1B). By contrast, only 7/25 (28%) of organ-confined tumours showed co-expression of mPGES-1 and EGFR (Fig. 1A, 70.3% vs 28%, see also panel a and b from an organ confined PCa sample vs c and d from an advanced PCa sample).

In DU145 cell line isolated from a castration-resistant human brain metastasis of PCa, characterized by mPGES-1 expression (mPGES-1SC) (Hanaka et al. 2009), the knock down for mPGES-1 (stable or transient mPGES-1KD) did not affect PGE2 receptor expression (EP1-4), while it obliterated PGE2 output (> 90%, P<0.001 vs mPGES-1SC) (A and B, and Supplementary Figure 1, see section on supplementary data given at the end of this article). The large PGE2 loss occurred despite a slight increase in cyclooxygenase-2 (COX-2) expression, while other enzymes involved in PGE2 metabolism were either unchanged (e.g. COX-1), or only negligibly changed (e.g. the cytosolic isofrom, cPGES, the microsomal type 2 isofrom, mPGES-2, the prostaglandin transporter, PGT and the enzyme implicated in PGE2 degradation, 15-hydroxyprostaglandin dehydrogenase (PGDH)) (Fig. 2A). Compared to DU145 WT, transfection of cells with the empty vector (mPGES-1SC) did not affect the PGE2 signalling cascade (Fig. 2A). Similar results were obtained in experiments of transiently silenced mPGES-1 cells (Supplementary Figure 2A).

Because PGE2 regulates genes involved in EMT in tumour cell lines (Dohadwala et al. 2006), we assessed whether constitutively high mPGES-1 expression in DU145 affected their mesenchymal-cell-like phenotype with respect to mPGES-1KD cells. The results showed that ~1/4 of EMT-related genes (n = 22/88) were influenced by high expression of intrinsic mPGES-1 (Fig. 2B). Notable
changes occurred in transcription factors known to induce EMT, such as Snail, Slug (SNAI2 gene) and ZEB, and in several genes coding for cytoskeletal proteins, such as E-cadherin, vimentin, fibronectin and ahnak. Western blot and immunofluorescence analysis illustrated the marked loss of vimentin, fibronectin and ahnak and the increased expression of E-cadherin in stably and transiently silenced mPGES-1 KD cells (C and D). Reduced expression of vimentin, fibronectin and ahnak was also detected by quantitative RT-PCR (Table 1). Treatment of mPGES-1 KD cells with PGE2 (1 μmol/l, 24 h) reversed both vimentin and fibronectin expression, confirming the involvement of PGE2 in this process (Fig. 2E). In line with the role of PGE2 in cancer cell growth and E-cadherin expression (Castellone et al. 2005, Lu et al. 2012), we found massive translocation of β-catenin into the nucleus of DU145 mPGES-1 SC cells, while it remained localized in the cytoplasm of mPGES-1 KD cells (Fig. 2E and F).

We also observed a prevalence of stem-cell-like markers, regarded as indicators of tumour invasiveness, in DU145 mPGES-1 SC compared to mPGES-1 KD cells. This was indicated by the large increase in the CD44+ /CD24− ratio, the decrease in α6-integrin and the increase in β1 integrin, as well as transcription factors Nanog and Oct4 in mPGES-1 SC (Fig. 3A, B and C), all pluripotency maintaining factors overexpressed in PCa stem cells (Klarmann et al. 2009, Rentala et al. 2010, Marthick & Dickinson 2012, Nanta et al. 2013, Hoogland et al. 2014). Similar results were obtained for the PC-3 cell line. PC-3 cells expressed constitutive mPGES-1, and knocking down the enzyme significantly reduced PGE2 output (> 60%, P<0.01 vs mPGES-1 SC), promoted epithelial
Inhibition of mPGES-1 activity suppresses stem-like phenotype

Further evidence that the mPGES-1-PGE2 cascade enhances PCa cell aggressiveness was obtained by phenotype, decreased cell clonogenicity and reduced the expression of stem cell markers (Supplementary Figure 3A, B, C, D, E and F, see section on supplementary data given at the end of this article). We also recorded anchorage-independent cell viability in mPGES-1SC cells, in contrast to the drastic decline seen in mPGES-1KD cells, associated with caspase-3 activation (Fig. 3D and E, and Supplementary Figure 3G and H). Adhesion studies with fibronectin coated-wells or endothelial cells revealed that mPGES-1SC cells rapidly adhered (2 h) to the matrix or endothelium (whether or not it had been activated by TNFα to favour cell–cell interaction and transmigration), whereas mPGES-1KD cells displayed a significantly delayed adhesion (F and G). Since knock down of mPGES-1 in cells does not affect cell survival in vitro, as measured by MTT assay (Abs 0.95 ± 0.08 and 0.88 ± 0.11 for DU145 mPGES-1SC and mPGES-1KD cells; Abs 0.74 ± 0.06 and 0.79 ± 0.09 for PC-3 mPGES-1SC and mPGES-1KD cells respectively), or cell apoptosis in suspension up to 24–48 h, we conclude that the inhibition of adhesion in mPGES-1 KD cells depends on the modification of cytoskeletal organization, for example of vimentin and fibronectin.

Finally, by measuring transendothelial migration, we showed a far greater ability (nearly threefold) of mPGES-1SC than mPGES-1KD cells to cross cell layers (Fig. 3H), indicating the involvement of mPGES-1/PGE2 signalling in prostate tumour invasiveness.
mPGES-1 mediates prostate malignancy

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Research

Endocrine-Related Cancer

Figure 3

EGFR activation mediates mPGES-1/PGE2-dependent EMT of DU145 cells. (A) FACS analysis for CD24 and CD44 expression in DU145 mPGES-1SC and mPGES-1KD cells. (B) Western blot analysis and quantification of α6 and β1-integrin expression in mPGES-1SC and mPGES-1KD cells. (C) RT-PCR analysis of Nanog and Oct4 expression in mPGES-1SC and mPGES-1KD DU145 cells. (D) Cell viability of DU145 mPGES-1SC and mPGES-1KD untreated or pretreated with PGE2 (1 μmol/l, 24–48 h). (E) Western blot analysis of caspase 3 activation in DU145 cells grown in suspension for the indicated time. (F) Adhesion of DU145 mPGES-1SC and mPGES-1KD untreated or pretreated with PGE2 (1 μmol/l, 96 h) on fibronectin-coated 96 well plate.

mPGES-1 induces growth and lung metastasis formation

In light of the observed link between mPGES-1/PGE2 signalling, EMT and stemness markers in PCa cells, we investigated whether the presence of mPGES-1 in DU145 and PC-3 cells influenced their growth and metastatic invasion in vivo.

In tumours induced by inoculating nude mice with mPGES-1SC or mPGES-1KD cells, the volume measurements (here reported at days 12 and 21) showed significant differences, growth being 2.3-fold higher in DU145 mPGES-1SC than mPGES-1KD tumours, and 4.9-fold higher in PC-3 mPGES-1SC than mPGES-1KD tumours (Fig. 5A). Consistently, Ki-67 showed denser immunostaining in mPGES-1SC than mPGES-1KD tumours (Fig. 5B). Accordingly, the Ki67 score was 45 ± 2.9% and 57 ± 3.8% for DU145 and PC-3 mPGES-1SC respectively and 21 ± 1.9% and 29 ± 1.8% for DU145 and PC-3 mPGES-1KD respectively, indicating a higher proliferation rate in cells overexpressing mPGES-1.

Western blot (Fig. 5C) and immunohistochemistry of tumours (Fig. 5D) revealed abundant mPGES-1 and vimentin expression in mPGES-1SC, contrasting with the absence of these two proteins in mPGES-1KD tumours.

We also investigated the contribution of mPGES-1 expression to PCa metastasis by injecting DU145 and PC-3 cells (mPGES-1SC and mPGES-1KD cells for both cell lines) via free access
whether mPGES-1/PGE$_2$ signalling modulates EGFR expression levels in the mPGES-1SC group than in the mPGES-1KD group (Fig. 6D), suggesting that EGFR signalling plays a role in mPGES-1 enhancement of tumour growth.

mPGES-1 induces EGFR expression in prostate tumours and mediates EGFR-dependent tumour growth

It is well documented that PGE$_2$ favours EGF/EGFR-induced oncogenicity by directly phosphorylating EGFR (Buchanan et al. 2003, Donnini et al. 2007). However, it is unknown whether mPGES-1/PGE$_2$ signalling modulates EGFR expression levels. We measured EGFR expression in DU145 mPGES-1SC and mPGES-1KD cells, in tumours from mice inoculated with the respective cell lines and in DU145 and PC-3 treated with MF63. We found distinctly higher EGFR expression levels in DU145 and PC-3 mPGES-1 SC, the number of colonies was 18 ± 2 and 20 ± 3.8, respectively. In contrast, knock down of mPGES-1 significantly reduced the numbers of colonies (10 ± 1.9 and 3.3 ± 2.8 respectively, Fig. 5E).

These results demonstrate that mPGES-1/PGE$_2$ signalling plays a significant role in prostate tumour growth and metastasis development.

In vivo, erlotinib administration to DU145 cells (10 μmol/l) abrogated EGFR phosphorylation and downregulated vimentin expression (Fig. 6G). Phosphorylation of EGFR was independent of EGF expression, which was affected in mPGES-1KD cells but not in mPGES-1SC (Supplementary Figure 5, see section on supplementary data given at the end of this article). Erlotinib also reduced cell viability in DU145 mPGES-1KD cells and functionally halved the number of cell colonies (number of colonies: mPGES-1SC = 58 ± 4 vs mPGES-1KD = 27 ± 5) (Fig. 6H, I, J, K and L). In vivo, erlotinib decreased tumour growth in DU145 mPGES-1SC and mPGES-1KD-bearing mice with respect to the vehicle-treated group (area under curve: mPGES-1SC + erlotinib = 12360 vs mPGES-1KD + erlotinib = 5664, Fig. 6M). Erlotinib treatment was more effective in reducing tumour volume when mPGES-1 was knocked down (Fig. 6I), but it did not affect the number of metastases for DU145 and PC-3 cell lines (number of metastases: mPGES-1SC = 20 ± 4.8 and mPGES-1KD = 7.5 ± 3.5 for DU145; mPGES-1SC = 17.7 ± 4.2 and mPGES-1KD = 5.3 ± 3.4 for PC-3).

All together, this data provides clear evidence of the role played by the mPGES-1/PGE$_2$ pathway in inducing a mesenchymal phenotype and stemness in PCa cells, thus reinforcing EGFR tumorigenic drive.
Discussion

The present study shows that by eliciting mesenchymal and stem-cell-like traits and EGFR expression, the tumour intrinsic inflammatory mPGES-1/PGE2 pathway cooperates with the EGFR oncogene to promote an aggressive PCa phenotype.

As an experimental paradigm we used DU145 and PC-3 cells in which mPGES-1 was stably or transiently knocked down by mRNA silencing (mPGES-1KD), comparing them with prostate cells containing a negative control non-targeting shRNA plasmid (mPGES-1SC). Further, evidence of the specificity of mPGES-1/PGE2 signalling in PCa aggressiveness was obtained by pharmacological inhibition of the enzyme with the selective MF63 inhibitor (Xu et al. 2008).

Inflammation plays a role in the development and progression of many cancers, including PCa, and multiple pro-inflammatory molecules are associated with PCa recurrence (Ørsted & Bojesen 2013). Here we investigated the contribution of the mPGES-1/PGE2 pathway to EMT, a process that promotes acquisition of mesenchymal traits, such as enhanced growth and migratory capacity, invasiveness and resistance to apoptosis, by epithelial cells. We observed that mPGES-1 knockdown influenced a set of genes promoting EMT in tumour cells, such as genes coding for transcriptional activity (Snail, Slug and ZEB), which were significantly downregulated. The effect on the cytoskeletal protein vimentin, which became undetectable in mPGES-1 KD and in cells treated with MF63, was particularly striking. A number of other changes occurred
in mPGES-1 KD cells. They included downregulation of insulin growth factor binding protein-4 (IGFBP-4) and the integrin β1 (ITGB1) gene, re-localization of β-catenin and overexpression of E-cadherin. All these markers are known to be associated with PCa aggressiveness (Damon et al. 1998, Miyake et al. 2000, Mohan & Baylink 2002, Bijnsdorp et al. 2013, Carbonell et al. 2013). Regulation of these markers may therefore be causally related to the loss of function parameters observed in mPGES-1 KD cells.

mPGES-1 expression in PCa cells was clearly associated with stem-like features as demonstrated by the greater survival ability of mPGES-1 SC cells in suspension compared to mPGES-1 KD cells. Additional evidence of mPGES-1-linked stemness was a significant shift in the CD44/CD24 ratio, decreased β1-integrin, increased...
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z6-integrin expression and increased transcription factors Nanog and Oct4 (Nanta et al. 2013). Collectively, these results provide evidence of involvement of mPGES-1/PGE2 in promoting EMT and stemness in mPGES-1SC tumour cells, since its ablation and inhibition impairs their inherent potential to survive in suspension and to transmigrate to endothelial cells.

EGFR studies provided another means to examine the interplay between PGE2 input and oncogenic drive mediated by the EGFR system. The picture emerging is one of reciprocal activation producing vigorous PCA progression when both components (PGE2 and EGF) of this circuit are maximally expressed, as in mPGES-1SC1KD tumours. Further, ablation of PGE2 production interferes with tumour progression and in contributing, in association with the EGFR pathway, to aggressiveness in PCA. Indeed, erlotinib treatment reduced tumour development in the mPGES-1SC group to approximately the same level as that observed in mPGES-1KD tumours. Further, ablation of PGE2 production significantly reduced lung metastasis in mice, indicating that mPGES-1 signalling plays a key role in EMT and stemness of PCa cells (Oskarsson et al. 2014). The combination of mPGES-1 knockdown or pharmacological inhibition of mPGES-1 by MF63 with erlotinib results in a significant reduction of lung metastasis in mice, indicating a low Gleason score. Thus, mPGES-1/PGE2 signalling empowers tumour cells to disseminate and seed metastases by activating EMT and stemness in PCA.

The present findings illustrate the role of mPGES-1 signalling in influencing EGFR-mediated oncogenicity and in contributing, in association with the EGFR pathway, to aggressiveness in PCA. Indeed, erlotinib treatment reduced tumour development in the mPGES-1SC group to approximately the same level as that observed in mPGES-1KD tumours. Further, ablation of PGE2 production significantly reduced lung metastasis in mice, indicating that mPGES-1 signalling plays a key role in EMT and stemness of PCa cells (Oskarsson et al. 2014). The combination of mPGES-1 knockdown or pharmacological inhibition of mPGES-1 by MF63 with erlotinib results in a more effective strategy to inhibit PCA cell growth. Ability to undergo EMT and acquire stem-cell-like features enables tumours to expand, spread and resist chemother-apy/target therapy (Clevers 2011, Hoggatt et al. 2013). PGE2 release in the tumour microenvironment was recently shown to be responsible for tumour initiation and repopulation, hence inhibitors of PGE2 production interfere with tumour progression and chemoresistance (Kurtova et al. 2015). Our study demonstrates that mPGES-1 ablation or inhibition in human PCA cells suppresses their overall oncogenic drive and reduces their stemness and invasiveness. We conclude that the mPGES-1 gene may be considered a signature gene for identifying a subtype of rapidly progressing prostate tumours, which, in the case of overexpression of EGFR, may benefit from combined treatment with inhibitors of EGFR, tyrosine kinase and PGE2.

Supplementary data

This is linked to the online version of the paper at http://dx.doi.org/10.1530/ERC-15-0277.

Declaration of interest

The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

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