

# ErbB3 Binding Protein 1 Represses Metastasis-Promoting Gene *Anterior Gradient Protein 2* in Prostate Cancer

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## Abstract

Dysregulation of the developmental gene *anterior gradient protein 2* (*AGR2*) has been associated with a metastatic phenotype, but its mechanism of action and control in prostate cancers is unknown. In this study, we show that overexpression of *AGR2* promotes the motility and invasiveness of nonmetastatic LNCaP tumor cells, whereas silencing of *AGR2* in the metastatic derivative C4-2B blocks invasive behavior. ErbB3 binding protein 1 (EBP1), a putative repressor of *AGR2*, is attenuated in prostate cancer. We show that the anti-invasive effect of EBP1 occurs, at least in part, through its ability to inhibit *AGR2* expression. Mechanistic investigations indicate that EBP1 downregulates Foxa1- and Foxa2-stimulated *AGR2* transcription and decreases metastatic behavior. In contrast, EBP1 ablation upregulates *AGR2* via Foxa1- and Foxa2-stimulated *AGR2* promoter activity and increases metastatic behavior. In both prostate cell lines and primary tumors, we documented an inverse correlation between EBP1 and *AGR2* levels. Collectively, our results reveal an EBP1-Foxa-*AGR2* signaling circuit with functional significance in metastatic prostate cancer. *Cancer Res*; 70(1); 240–8. ©2010 AACR.

## Introduction

Metastasis is the major cause of death from prostate cancer. Interruption of early metastases is crucial for a majority of individuals with prostate cancer. Given the complex cascade of events in metastasis, including invasion, intravasation, extravasation, and colonization, defining the molecules and underlying mechanisms required for these steps may help identify therapeutic targets (1–3).

Aberrant or untimely activation of developmentally regulated genes contributes disproportionately to metastasis. *Anterior gradient protein 2* (*AGR2*), isolated in a screen for differentially expressed genes in neural development of *Xenopus laevis*, plays an essential role in the formation of the forebrain and the mucus-secreting cement gland (4). In humans, *AGR2* was first identified in breast cancers and forced expression of *AGR2* cDNA conferred a metastatic phenotype on benign nonmetastatic rat breast carcinoma cells (5). Aberrant *AGR2* expression has been found in adenocarcinomas of breast (6–8), esophagus (9, 10), pancreas (11–13), and prostate (14, 15), but how *AGR2* promotes the invasive phenotype in prostate cancer remains unknown.

Foxa (Foxa1, Foxa2, and Foxa3) proteins belong to a superfamily of forkhead transcription factors, which act as genetic potentiators that facilitate the binding of other transcriptional factors during development and differentiation (16, 17). A recent report suggests that Foxa1 and Foxa2 positively regulate *AGR2* promoter activity (18). In addition, Foxa proteins are important in prostate carcinogenesis. In particular, Foxa2 may be involved in progression of prostate cancer to androgen independence (19).

ErbB3 binding protein 1 (EBP1), cloned by our group, is the human homologue of a previously identified cell cycle-regulated mouse protein p38-2G4 (20). EBP1 is a conserved molecule with multiple roles in cell growth, differentiation, and apoptosis (21). In particular, EBP1 has been characterized as a negative regulator of cross-talk between HRG-triggered signaling and the androgen receptor (AR) axis in prostate cancer (22–24). The expression of the *EBP1* gene is reduced in clinical and preclinical models of prostate cancer (24, 25). Restoration of EBP1 expression ameliorates the hormone-refractory phenotype both *in vitro* and *in vivo* (25). In conducting a microarray analysis of *EBP1*-transfected prostate cancer cells to determine the spectrum of differentially expressed genes contributing to the hormone-refractory phenotype, we found that *AGR2* is an EBP1-downregulated gene (25). Our current work identifies a previously uncharacterized signaling circuit that involves EBP1-Foxa-*AGR2*, which has potential mechanistic and functional significance in therapeutic management of metastatic prostate cancer.

## Materials and Methods

**Cell culture, gene transfection, and luciferase reporter assay.** The LNCaP cell line was obtained from the American

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Type Culture Collection. C81, C4-2, and C4-2B cells were gifts of Dr. Yun Qiu (University of Maryland) per permission of Drs. Lin (26) and Chung (27). C81 sublines stably transfected with *EBP1* cDNA or vector control, EBP1-null C13, and control A16 cells were established previously in our lab (25). Cells were maintained at 37°C in a humidified atmosphere of 5% CO<sub>2</sub> in air. Cell lines were routinely cultured in RPMI 1640 (LNCaP and C81) or T medium supplemented (C4-2 and C4-2B) with 10% fetal bovine serum (Sigma).

pGL3B-AGR2, a reporter construct containing the *AGR2* promoter, was a gift from Dr. J. Hampe (18). Foxa1 or Foxa2 expression plasmids were gifts of Dr. R. Matusik (19). Cells were seeded in six-well plates in complete medium for 24 h. Transfections were performed with Fugene 6 (Roche) with various amounts of plasmids as indicated. Luciferase activity was measured as previously described (18, 22).

**Creation of stably transfected cell lines.** LNCaP cells stably transfected with pcDNA3.1 or pcDNA3.1-hAGR2 plasmids (a gift from Dr. C. Young, Mayo Clinic) or C4-2B cells stably transfected with pEGFP-EBP1 (C4-2BE) or vector control (C4-2BG) were established as previously described (22). To generate an AGR2-silenced cell line, C4-2B cells were seeded into 96-well plates and transduced with lentiviral particles corresponding to different short hairpin RNA (shRNA) constructs targeting the *AGR2* gene (Mission shRNA, NM\_006408; Sigma) and the lentiviral transduction particles produced from the sequence-verified lentiviral plasmid, pKLO.1-puro control vector, as a negative control (SHC001V, 03130808MN; Sigma). The derivation of the EBP1 stably transfected C81 cell line and the EBP1-deleted LNCaP C13 and A16 cell lines and their expression of EBP1 have previously been published (25).

**Glutathione S-transferase pull-down assay, immunoprecipitation, DNA affinity precipitation, and western blot analysis.** For studies determining the interaction of EBP1 with Foxa, cell lysates were incubated with glutathione S-transferase (GST)–EBP1, purified as previously described (22), for GST pull-down assay or green fluorescent protein (GFP)–conjugated GFP agarose (Millipore) for immunoprecipitation and analyzed by Western blotting, where indicated, as described previously (22). The interaction of Foxa with the *AGR2* promoter sequence was assessed by DNA affinity precipitation as previously described (28) using LNCaP cell extracts and biotinylated oligonucleotides GTGGGTTACTTGATTGTATTTTTTTCAT (Olig A) containing the Foxa binding consensus elements or ATTCTGAGCTTTTAAAGACTGCACAACT (Olig B) without any potential Foxa binding sites, all derived from the *AGR2* promoter. The polyclonal AGR2 antibody was a gift from Dr. C. Young and the EBP1 antibody was from Millipore. The monoclonal anti-β-actin antibody was from Sigma. Goat polyclonal antibodies Foxa1 (C-20) and Foxa2 (P-19) were from Santa Cruz Biotechnology.

**Boyden chamber assay.** A modified Boyden chamber assay as described previously (29–31) was used to determine cell migration and invasion. Basically, culture plate inserts (8-μm pore size and 12-mm diameter, Millicell-PCF) were coated with 150 μL PBS containing 10 μg collagen and 1 μg fibronectin (BD Biosciences) for 1 h at room temperature. For invasion assays, collagen-coated filters were air

dried, further coated with a total of 30 μL of Matrigel matrix (BD Biosciences) diluted 1:5 in serum-free medium, and allowed to gel in a 37°C incubator for 30 min before adding the cells suspended in 450 μL medium with 5% charcoal-stripped serum. Bottom wells in the system were filled with 600 μL complete medium. After 40 h of incubation, the inserts were fixed in 10% formalin for 20 min and, after washing with water, stained with 0.5% crystal violet in 25% methanol for 30 TO 60 min. Nonmigrating or invading cells on the top of the filters were removed with a cotton swab. Cells that had migrated or invaded to the undersurface of the filter were examined at ×10 or ×20 magnification. Three representative areas were photographed, and the number of cells was counted. Each experiment was performed in triplicate.

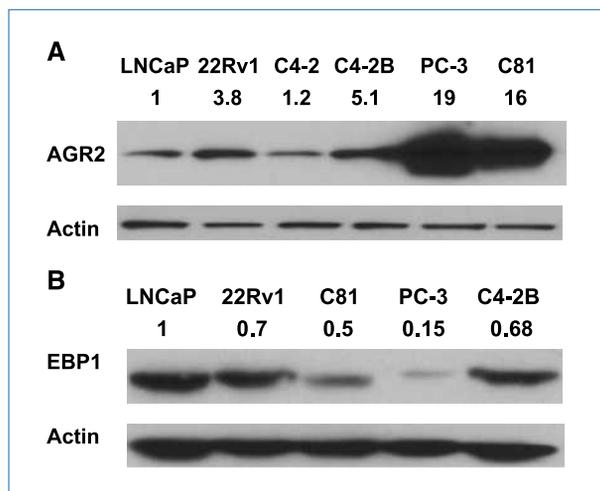
#### **Tissue microarray and immunohistochemical analysis.**

An intermediate density tissue microarray of prostate cancer and benign prostate tissues was received from the Cooperative Prostate Cancer Tissue Resource (National Cancer Institute). Immunohistochemical analysis of AGR2 was performed as previously described (14). Immunostaining was evaluated as in ref. 25.

**Statistical analysis.** Results for luciferase and Boyden chamber assays and correlation between EBP1 and AGR2 protein levels in cell lines were analyzed using a two-tailed Student's *t* test or Mann-Whitney *U* test or one-way ANOVA as appropriate. The association between AGR2 expression and the nature of the prostate tissues (benign, hormone responsive, hormone refractory) was assessed by the  $\chi^2$  test. The Pearson's correlation coefficient test was used to determine the relationship between intensity of AGR2 staining and EBP1 levels.  $P < 0.05$  was considered statistically significant.

## **Results**

**EBP1 levels inversely correlate with AGR2 expression in prostate cancer cell lines.** To establish a suitable cellular system to investigate how EBP1 regulates AGR2 expression, EBP1 and AGR2 expression levels were measured in a panel of prostate cancer cell lines. Examination focused on several preclinical models recapitulating the hormone-refractory phenotype. The LNCaP subline C81 has been made androgen independent by continuous long-term passage of LNCaP cells in complete medium (26). The LAPC-4 xenograft grows as an androgen-dependent cancer in male severe combined immunodeficient mice and regresses in response to androgen ablation but eventually regrows as an androgen-independent tumor (32). The LNCaP derivative C4-2B exhibits two cardinal features of human prostate carcinoma progression: androgen independence and osseous metastasis (29). Intraosseous injection of androgen-independent 22Rv1 cells generates osteolytic and osteoblastic responses, a subset of the pathology exhibited in patients with metastasis (33). AGR2 protein expression was much higher in androgen-refractory 22Rv1, C81, metastatic C4-2B, and PC-3 cells compared with androgen-dependent nonmetastatic LNCaP cells (Fig. 1A; refs. 34, 35), suggesting that AGR2 contributes to malignant progression in prostate cancer. Importantly, among these cell lines, there is a significant inverse correlation between EBP1 and AGR2 expression



**Figure 1.** EBP1 levels inversely correlate with AGR2 expression in prostate cancer cell lines. *A* and *B*, immunoblots detecting AGR2 (*A*) and EBP1 (*B*) expression in prostate cancer cell lines. Values on top of the bands represent relative densities normalized to actin.

in prostate cancer cell lines ( $P < 0.05$ , one-way ANOVA; Fig. 1*A* versus *B*), supporting our previous finding that EBP1 might be a negative regulator of AGR2 (25, 36).

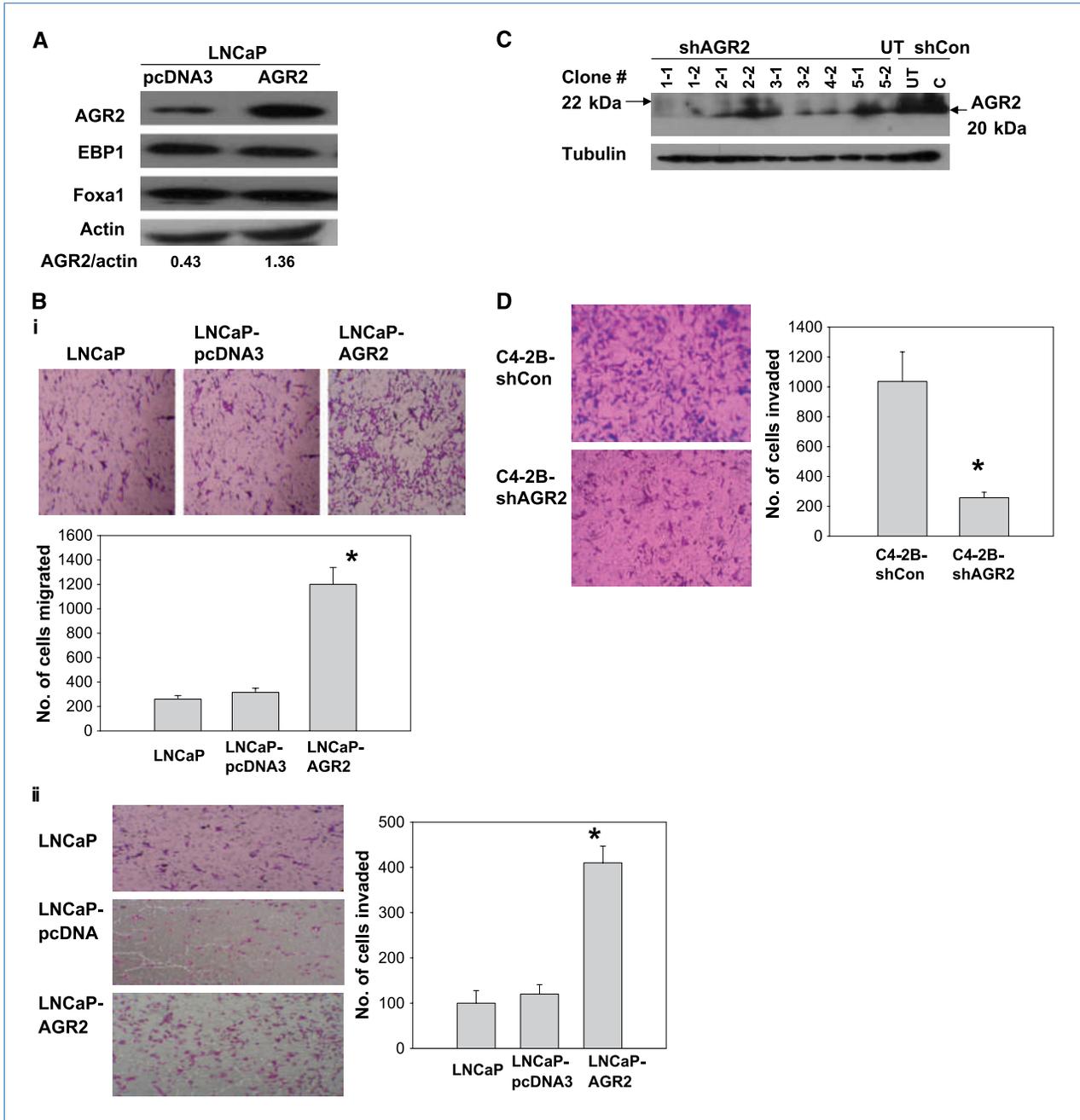
**AGR2 promotes the motility and invasiveness of prostate cancer cells.** Nonmetastatic LNCaP cells expressing low levels of AGR2 (Fig. 1) were selected to test if forced expression of AGR2 cDNA promotes motility and invasiveness. LNCaP cells stably transfected with pcDNA3.1-hAGR2 cDNA have a 3-fold increase in AGR2 expression but no changes in EBP1 and Foxa1 protein levels compared with the vector control-transfected cells (Fig. 2*A*). Forced expression of AGR2 resulted in a marked (>10-fold) increase of motility in LNCaP cells as measured by migration assays ( $P < 0.01$  versus control; Fig. 2*B*, *i*). Increased AGR2 expression also promoted the invasiveness of LNCaP cells through Matrigel, >3-fold over control transfectants ( $P < 0.01$ ; Fig. 2*B*, *ii*). Of note, transfection of pcDNA has no effect on cell migration and invasiveness of LNCaP cells (Fig. 2*B*, *i* and *ii*). In addition, at the 40-hour time point, there was no difference in numbers of LNCaP control and AGR2-transfected cells (data not shown). To further assess the effect of AGR2 in the metastatic process, we transduced metastatic LNCaP derivative C4-2B cells with five individual shRNA lentiviral particles constructs targeting AGR2 mRNA. Only one construct (NM\_006408.2-252s1c1; Sigma) targeting nucleotides 383 to 402 of AGR2 (Genbank NM\_006408.3) silenced AGR2 expression (clones 1-1 and 1-2 in Fig. 2*C*). To ensure that our results hereafter were not specific to any individual subclone of transfected cells, we pooled clones 1-1 and 1-2 for biological analysis. Silencing of endogenous AGR2 in metastatic C4-2B cells significantly reduced invasiveness ( $P < 0.01$  versus control; Fig. 2*D*). No decreases in invasiveness were observed in cells transduced by control lentiviruses (Fig. 2*D*) or by AGR2-targeted lentiviruses that did not decrease AGR2 expression (data not shown), suggesting that these biological changes were unlikely to be due to off-target ef-

fects. Silencing of AGR2 has no effect on cell proliferation at the 40-hour time point (data not shown). These results indicated that AGR2 expression is an important determinant of the invasive capacity of human prostate cancer cells.

**EBP1 modulation of the expression of AGR2 is accompanied by changes in motility and invasiveness.** As AGR2 augments invasiveness of nonmetastatic prostate cancer LNCaP cells *in vitro*, we tested if overexpressing or knocking down EBP1, which alters AGR2 expression, could modulate motility and invasive ability. Metastatic C4-2B cells stably transfected with pEGFP-EBP1 cDNA (C4-2BE) expressed a GFP-EBP1 fusion protein (Fig. 3*A*, *left* and *right*). Endogenous EBP1 levels were not changed (data not shown). Ectopic expression of EBP1 in C4-2B cells (C4-2BE) led to a significant reduction of AGR2 protein as compared with pEGFP transfectants (C4-2BG; Fig. 3*A*, *right*). Steady-state levels of AGR2 mRNA were decreased 35% in EBP1 transfectants as compared with vector controls. Expression of Foxa1 remained the same (Fig. 3*A*, *right*), indicating that EBP1 inhibition of AGR2 is not cell type specific (25). C4-2BE cells were less motile (Fig. 3*B*) and invasive (Fig. 3*C*) than C4-2BG controls ( $P < 0.01$  versus control), similar to C4-2B cells in which AGR2 was suppressed by shRNA (Fig. 2*D*). In addition, restoration of EBP1 in androgen-independent C81 cells (C81E), leading to reduced AGR2 expression (25), decreased migration ( $P < 0.01$ ; Fig. 3*D*, *i*). Knocking down EBP1 in LNCaP cells (C13), leading to enhanced expression of AGR2 (25), increased motility compared with A16 control cells ( $P < 0.01$ ; Fig. 3*D*, *ii*). These data support the notion that EBP1 suppresses invasiveness in part via inhibition of expression of the AGR2 gene.

**EBP1 suppresses the AGR2 promoter.** Our previous microarray and quantitative real-time PCR data suggested that EBP1 repression of AGR2 protein levels might be at the transcriptional level (25). To determine if the mechanism of AGR2 repression involves transcriptional regulation, we tested if EBP1 inhibits the promoter activity of the AGR2 gene using luciferase reporter assays. Transient transfection of exogenous EBP1 cDNA in LNCaP cells inhibited AGR2 promoter activity in a dose-dependent manner ( $P < 0.01$ , Mann-Whitney *U* test; Fig. 4*A*, *i*), whereas levels of Foxa1 and Foxa2 protein remain the same (data not shown), suggesting that EBP1 might negatively regulate the expression of the AGR2 gene at the transcriptional level. Zheng and colleagues (18) recently reported that Foxa1 and Foxa2, implicated in prostate carcinogenesis, activate the human AGR2 promoter. We determined that EBP1 diminished Foxa1- and Foxa2-stimulated AGR2 transactivation ( $P < 0.01$ , one-way ANOVA; Fig. 4*A*, *ii*), indicating that EBP1 might interfere with the regulation of AGR2 promoter activity by Foxa1 and Foxa2.

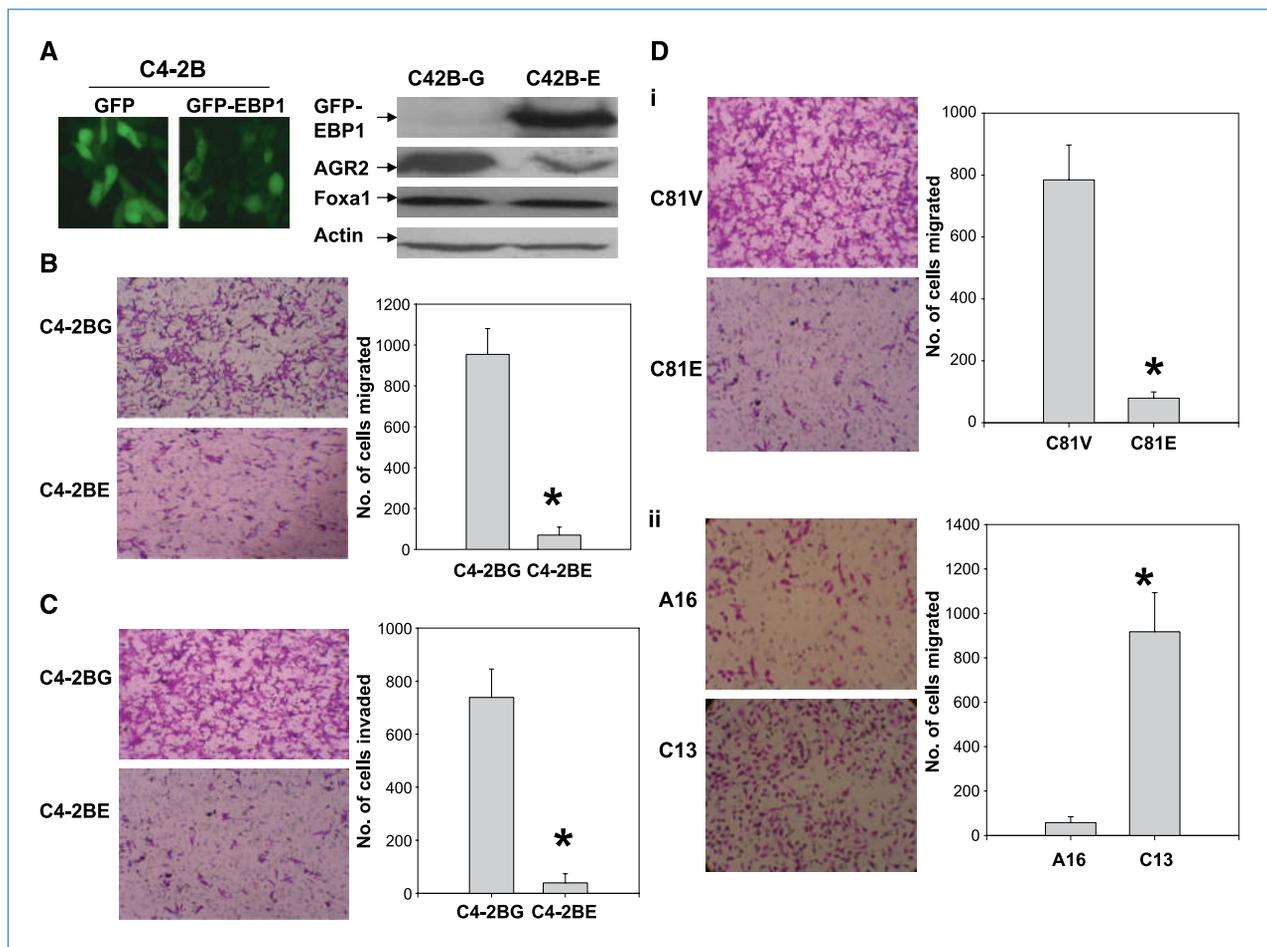
To solidify this initial observation and definitively establish the role of EBP1 in regulating AGR2 activity, we used EBP1-null LNCaP-derived C13 cells, which showed elevated AGR2 expression (25), to examine if AGR2 promoter activity is enhanced by Foxa in an EBP1-depleted environment. Both Foxa1 and Foxa2 induced greater AGR2 promoter activity in C13 cells as compared with control A16 cells ( $P < 0.01$ , one-way ANOVA; Fig. 4*A*, *iii*), suggesting that the interaction of EBP1 and Foxa plays an important role in regulating AGR2 expression.



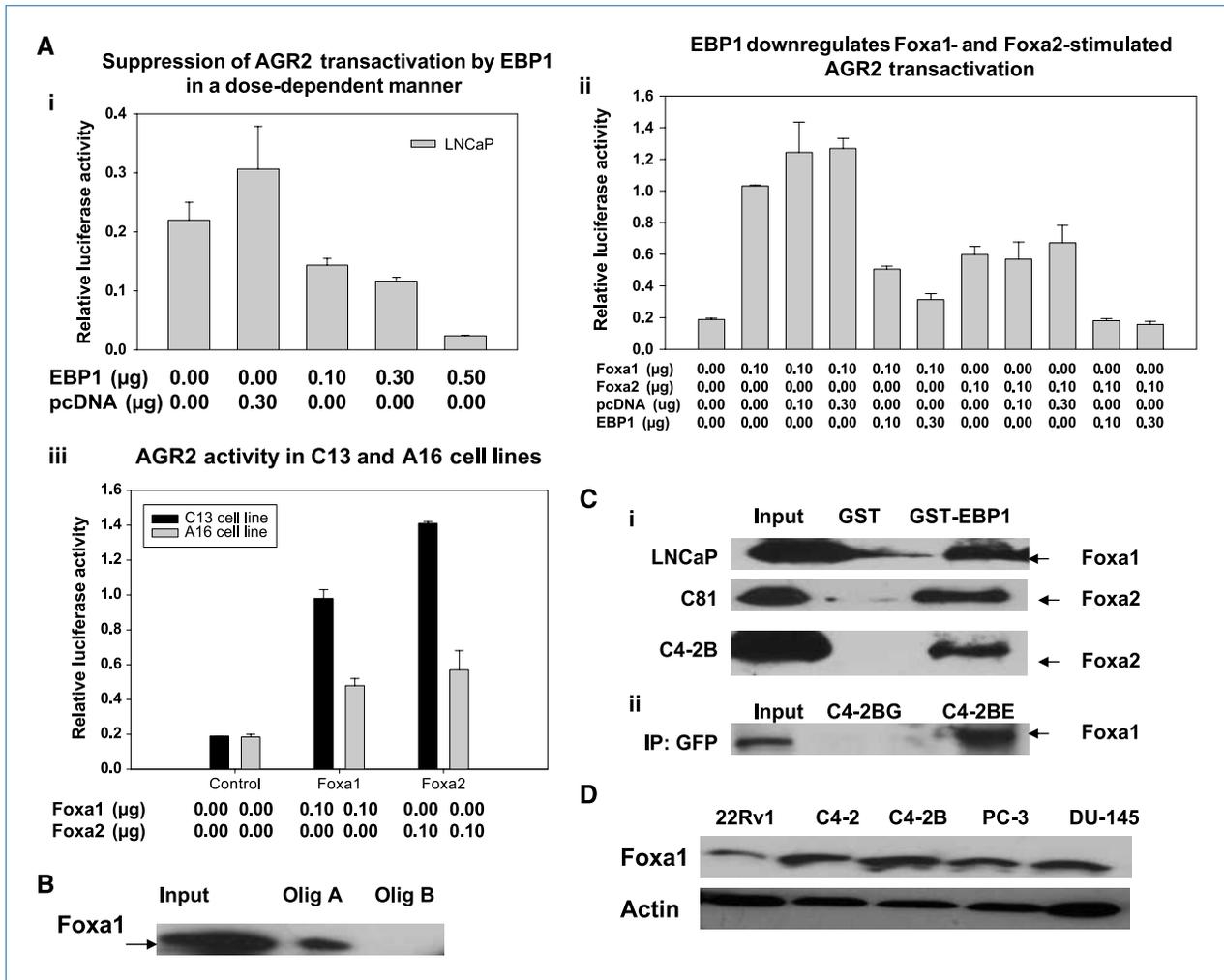
**Figure 2.** AGR2 promotes nonmetastatic prostate cancer cell migration and invasion. *A*, immunoblotting analysis of LNCaP cells stably transfected with pcDNA3 (*LNCaP-pcDNA3*) or pcDNA3-hAGR2 (*LNCaP-AGR2*) using antibodies as indicated. Values under panel represent relative band densities normalized to actin. *B*, *i*, migration ability was estimated using a modified Boyden chamber assay. *Top*, representative image of three independent experiments showing cells that migrated through the Transwell filters coated with collagen and fibronectin. LNCaP is the untransfected parental cell line. *Bottom*, number of migrated cells in representative areas of each membrane that were examined microscopically at  $\times 10$  magnification. *Columns*, mean number of cells that migrated in three representative fields per well; *bars*, SD. \*,  $P < 0.01$ , *t* test, versus control. Shown as one of three independent experiments. *ii*, *left*, representative photomicrograph of invasion assay from three comparable experiments showing cells invading through Matrigel-coated membranes; *right*, cells that had invaded in representative areas of each membrane were examined microscopically at  $\times 20$  magnification. *Columns*, mean number of cells that invaded in three representative fields per well; *bars*, SD. \*,  $P < 0.01$ , *t* test, versus control. Shown as one of three independent experiments. *C*, Western blot analysis of AGR2 expression in cells transduced with lentiviruses targeted to different portions of the AGR2 coding sequence. AGR2 was silenced in stable clones 1-1 and 1-2 using a virus targeting nucleotides 383 to 402 of the coding sequence. *UT*, untransduced cells; *shCon*, empty lentivirus. *D*, representative micrographs of effects of stable silencing of AGR2 on the invasiveness of C4-2B cells as described in *B*. Cells from clones 1-1 and 1-2 were pooled for the migration and invasion assays (*shAGR2*). *Right*, *columns*, mean number of cells that migrated in three representative fields per well; *bars*, SD. \*,  $P < 0.01$ , *t* test, versus control. Shown as one of three independent experiments.

Foxa proteins share four distinct conserved regions (CRI, CRII, CRIII, and CRIV) with an almost identical DNA-binding domain called the winged-helix/forkhead domain (CRI; refs. 16, 17). CRII and CRIII in the COOH terminus and CRIV in the NH<sub>2</sub> terminus are capable of distinct interactions with other transcription factor(s) and/or cofactor(s) for the activity of Foxa (37). The DNA-binding consensus sequence for Foxa proteins is VWWTRTTTRYTY or HWATTGAYTWWD with a 7-bp recognition core motif in common, whereas sequences flanking either side do not share any obvious similarity (16, 17). However, the Foxa-regulated element of the *AGR2* promoter is not known. Based on these criteria, we identified several potential binding sites of Foxa in the *AGR2* promoter (data not shown). Importantly, Foxa1 from LNCaP cell lysates bound specifically to the sequence containing Foxa consensus elements (Olig A, Fig. 4B) but not

the sequence lacking a potential binding motif (Olig B, Fig. 4B). We also examined if EBP1 physically associates with Foxa by *in vitro* GST pull-down assays in several cell lines. GST-EBP1 associated with endogenous Foxa1 and Foxa2 (Fig. 4C, i). Finally, we tested if EBP1 and Foxa1 associate *in vivo* by immunoprecipitation analysis of GFP-EBP1-transfected cells. These results indicated that EBP1 binds Foxa1 *in vivo* (Fig. 4C, ii). Foxa1 expression has been shown to vary in different cell lines (19). To determine if a potential EBP1-Foxa1 interaction may be important in different lines, we measured Foxa1 expression in several cell lines. We found, in agreement with previous reports (19), that Foxa1 was expressed in all cell lines tested at approximately equal levels (Figs. 2A, 3A, and 4D). Foxa2 was selectively expressed in LNCaP derivatives C81 and C4-2B cells (Fig. 4D). We hypothesize that EBP1



**Figure 3.** EBP1 modulates the expression of AGR2 accompanied by changes in motility and invasiveness in prostate cancer cells. *A*, C4-2B cells stably expressing GFP (C4-2BG) or GFP-EBP1 fusion proteins (C4-2BE) were visualized by fluorescence microscopy (left) and confirmed by Western blotting assay using GFP antibody (right, top). Right, middle top, middle bottom, and bottom, AGR2 and Foxa1 levels were also analyzed in these cells and actin was used as a loading control. *B*, left, photomicrographs of Boyden chamber membranes after migration assays showing that many C4-2BG but few C4-2BE cells traversed a coated membrane; right, quantification of the effects of EBP1 and AGR2 expression on migration. Columns, mean number of cells that migrated in three representative fields per well; bars, SD. Shown as one of three independent experiments. *C*, invasion assay was described in Fig. 2. *D*, left, micrographs of Boyden chamber membranes after migration assays with C81V and C81E (i) and A16 and C13 (ii). Right, relative quantification of the effects of EBP1 and AGR2 expression on the migration. Columns, mean number of cells that migrated in three representative fields per well; bars, SD. \*,  $P < 0.01$ , *t* test, versus control. Shown as one of three independent experiments.



**Figure 4.** EBP1 suppresses the promoter activity of the *AGR2* gene by interfering with Foxa transcription factors and binds Foxa proteins. *A, i to iii*, LNCaP or its subline EBP1-null C13 and control A16 cells were transfected with an *AGR2* reporter (pGL3B-*AGR2*, 0.25 µg/well) and other indicated plasmids. Forty-eight hours later, cells were harvested for luciferase measurement. *Columns*, mean of triplicate wells; *bars*, SE. Representative of three independent experiments. *B*, the interaction of Foxa with the *AGR2* promoter sequence was assessed by DNA affinity precipitation as previously described (28) using LNCaP cell extracts and biotinylated Olig A containing a Foxa consensus sequence or Olig B lacking potential Foxa binding sites, all derived from the *AGR2* promoter as described in Materials and Methods. *C*, association of EBP1 and Foxa proteins *in vitro* (*i*, top, middle top, and middle bottom) and *in vivo* (*ii*, bottom). Top, middle top, and middle bottom, equal amounts of GST-EBP1 or GST proteins were incubated with lysates of LNCaP, C81, or C4-2B cells. Associated proteins were analyzed by Western blot analysis with the antibodies indicated. Bottom, lysates of stably expressing GFP (C4-2BG) or GFP-EBP1 (C4-2BE) cells were incubated with GFP agarose and immunoprecipitation was performed using a GFP antibody. Associated proteins were analyzed by Western blotting using Foxa1 and Foxa2 antibodies. *D*, Western blot analysis of Foxa1 expression in a panel of prostate cancer cell lines.

represses Foxa-mediated transactivation of the *AGR2* gene by interference with the functions of Foxa DNA binding and/or transactivation.

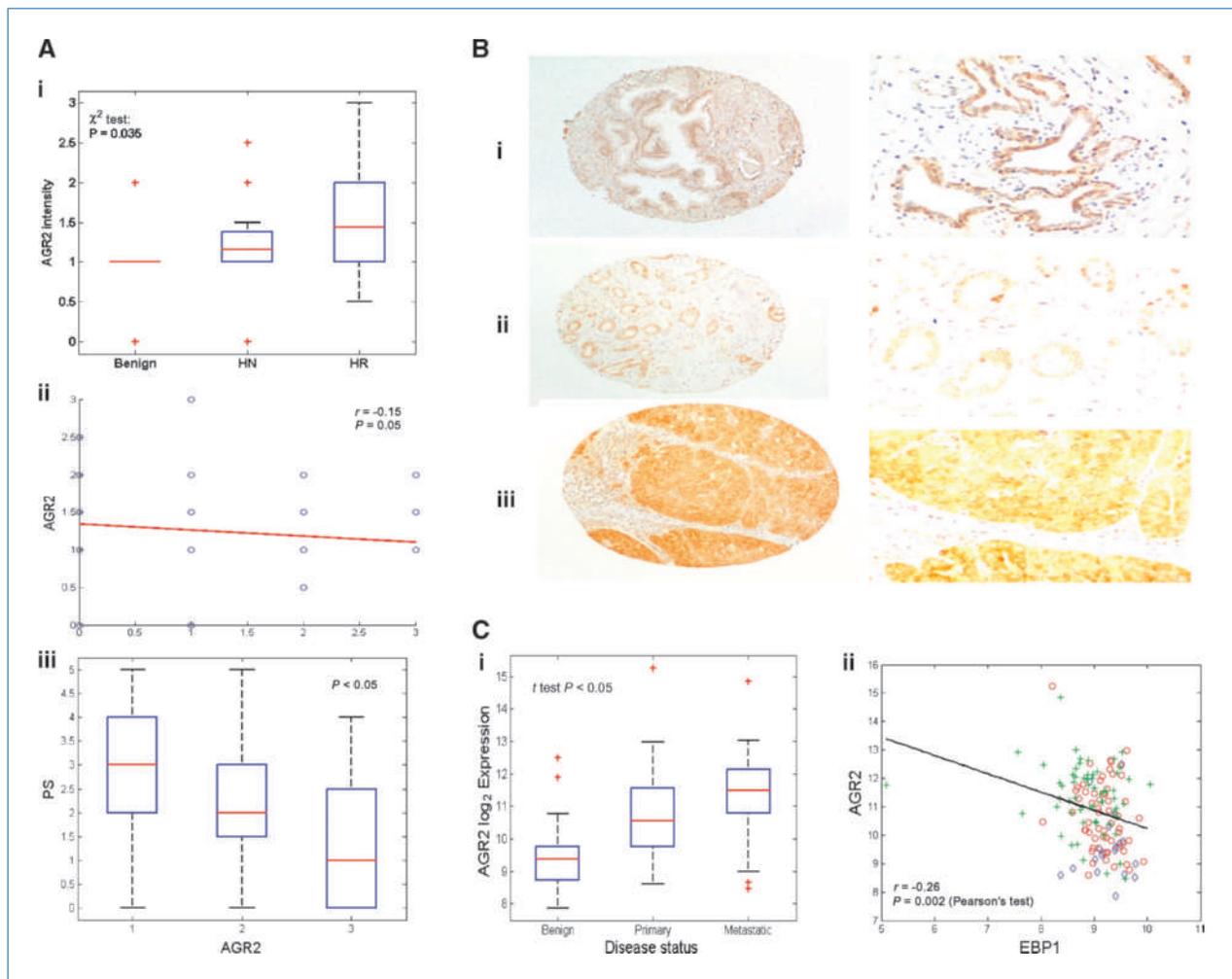
**Expression of *AGR2* increases with prostate cancer progression and inversely correlates with loss of EBP1 expression.** To extend our observations to the clinical setting, we next examined a possible correlation between EBP1 and *AGR2* expression levels in primary tumors. A prostate cancer tissue microarray previously used for EBP1 staining (25) was used for immunohistochemical analysis of *AGR2* (14). Immunohistochemical staining showed that among 17 benign cases, 5.88% were unstained, 52.94% stained weakly positive, and 35.29% stained moderately positive for *AGR2*. Among 51

hormone-responsive carcinoma cases, 60.78% stained weakly positive, 33.33% stained moderately positive, and 5.88% stained strongly positive. Among 36 hormone-refractory cases, 36.11% stained weakly positive, 41.67% stained moderately positive, and 22.22% stained strongly positive. The difference in *AGR2* staining among benign, hormone-responsive, and hormone-refractory cases is statistically significant ( $P = 0.035$ ,  $\chi^2$  test; Fig. 5*A, i*). The differences in staining of hormone-responsive and hormone-refractory versus benign tissues are statistically significant ( $P < 0.05$ ). Although the difference in staining of hormone-refractory versus hormone-responsive tissues is not statistically significant ( $P = 0.12$ ), the overall trend does indicate that *AGR2* expression increases with the progression of

prostate cancer (Fig. 5A, *i*). This is shown by the nonzero correlation coefficient ( $r = 0.17$ ,  $P = 0.038$ ) between AGR2 intensity and the progression status. In the benign tissue, AGR2 immunostaining can be observed mainly in basal cells (Fig. 5B, *i*), whereas in malignant tissues enhanced intensity is seen in both the cytoplasm and nucleus of luminal epithelial cells (Fig. 5B, *ii* and *iii*), similar to previous reports (14, 15). AGR2 was originally presumed to be a secreted protein. A recent analysis indicated that the KTEL sequence in the COOH terminus of AGR2 was recognized by the endoplasmic reticulum (ER) retention receptor and served to localize AGR2 predominantly to the ER of transfected cells (38). Studies on different isoforms or variants of AGR2 and its interaction with different proteins may thus provide some suggestion on what controls the localization of AGR2 protein

across species. We also found that increased AGR2 expression is significantly inversely correlated with EBP1 staining intensity ( $r = -0.15$ ,  $P = 0.05$ ; Fig. 5A, *ii*). In addition, increased AGR2 expression is significantly ( $P < 0.05$ ) correlated with a decrease in EBP1 labeling frequency (Fig. 5A, *iii*).

We further interrogated a publicly available cDNA microarray expression data set (39) that shows a statistically significant decrease of EBP1 expression with prostate cancer progression. We found a significant increase of AGR2 as prostate cancer progressed (Fig. 5C, *i*), indicating that AGR2 had a significant role in distant metastasis ( $P < 0.05$ ). To study the correlation between EBP1 and AGR2 expression, we directly quantified their expression relationship by plotting log-transformed expression units of EBP1 (Fig. 1B in ref. 25) against AGR2 staining and measured their expression similarity using



**Figure 5.** Expression of AGR2 increases with prostate cancer progression and inversely correlates with EBP1 expression. **A, i**, box plots of immunohistochemical staining intensity of AGR2 in normal prostate tissue ( $n = 42$ ) or tissues from patients with prostate cancer ( $n = 153$ ). For all panels, the box represents the SD of the distribution, and the line through that box represents the mean of the distribution. The horizontal lines above and below the box represent the extreme values of the distribution. **ii**, correlation of AGR2 staining intensity with EBP1 staining intensity. **iii**, correlation of AGR2 staining intensity with EBP1 proportion score (PS; percentages of cells stained with EBP1: 1, <5%; 2, 5–30%; 3, 31–70%; 4, >70%). **B**, representative staining of AGR2 in benign tissues (*i*), moderately malignant carcinomas (*ii*, Gleason 3 + 3), and highly malignant carcinomas (*iii*, Gleason 4 + 5). **C, i**, a public data set (39) shows the average gene expression level at three different stages (benign,  $n = 23$ ; primary,  $n = 64$ ; metastatic,  $n = 250$ ) of prostate cancer. **ii**, scatter plots of AGR2 versus EBP1 expression units in all these samples.

Pearson's correlation coefficients. As shown in Fig. 5C (ii), we observed a significant negative correlation between EBP1 and AGR2 across all samples with  $r = -0.26$  and  $P = 0.002$ . All together, these data suggest the possibility that EBP1 suppresses metastasis of prostate cancer cells by inhibiting the expression of the *AGR2* gene; the loss of EBP1 activity may functionally enhance invasiveness of advanced prostate cancer by releasing the brake on *AGR2* promoter activity.

## Discussion

The role of the developmentally expressed gene *AGR2* in metastasis is being increasingly studied in many cancer types. However, its action in prostate cancer specifically has not been analyzed extensively. This is the first report indicating that stable expression of mammalian *AGR2* cDNA in nonmetastatic prostate cancer cells promotes motility and invasiveness, linking increases of *AGR2* expression with a propensity for prostate cancer cells to undergo metastasis. A previous report showed that overexpression of *AGR2* in benign, nonmetastatic, rat mammary cells leads to metastasis (5). *AGR2* expression also promotes tumor growth in esophageal adenocarcinoma cells (10). Silencing *AGR2* significantly reduced cell proliferation and invasion of pancreatic cancer cells (13). The prognostic relevance of aberrant *AGR2* expression has consistently been shown in primary breast (6–8), esophagus (9, 10), pancreas (11–13), and prostate (14, 15) carcinomas. Moreover, in the first global gene expression profiling of circulating tumor cells, the *AGR2* gene was expressed in the majority of the metastatic samples, regardless of the cancer type (40). These results are in keeping with our current findings that *AGR2* serves a proinvasive and prometastatic function. Notably, we identified EBP1 as a novel potent suppressor of *AGR2*, in keeping with the demonstration that EBP1 is implicated in progression of prostate cancer (22, 24, 25). We hypothesize that EBP1 modulates the expression of *AGR2*, leading to changes in motility and invasiveness of prostate cancer cells. In addition, we observed a significant inverse correlation between EBP1 and *AGR2* expression in prostate cancer cell lines and clinical samples of prostate cancer. These results strongly support our concept that EBP1 suppresses the invasive ability of cells by inhibiting the expression of *AGR2*.

The mechanisms underlying the deregulation of *AGR2* in metastatic cancer remain elusive. A recent report suggested that genetic or epigenetic changes are not key to changes in *AGR2* levels. Rather, *AGR2* expression is controlled by a broad set of signaling and metabolic pathways (41). In the prostate, regulation of *AGR2* expression may be dependent or independent of the AR (14), which itself is subject to various upstream signals, such as aberrant growth factors and receptor tyrosine kinases (42). In this context, EBP1 negatively regulates cross-talk between the ErbB network and the AR signaling axis (22–24). On the other hand, forkhead transcription factors regulate highly complex, multigenic processes in development and are directly involved in malignancies (16, 17), including that of prostate (19). Interestingly, *AGR2* is positively modulated by Foxa1 and Foxa2, suggesting a potential Foxa-*AGR2* axis. EBP1 downregulated the expression

of *AGR2* by antagonizing Foxa activation of the *AGR2* promoter, leading to suppression of invasion and metastasis.

The detailed mechanisms underlying EBP1-mediated suppression of *AGR2* is under investigation in our laboratory. The current study indicated a physical interaction between EBP1 and Foxa, which bound to its consensus sequence in the *AGR2* promoter. The crystal structure of EBP1 suggests that the conserved LxxLL motif in EBP1 interacts with structurally diverse binding sites, such as AR, Rb, mSIN3, and HDAC2 (43). This motif might be well involved in the interaction between EBP1 and Foxa transcriptional factors, important in modulating *AGR2* transactivation. Indeed, the three Foxa proteins share four distinct conserved regions (CRI, CRII, CRIII, and CRIV; refs. 37, 44, 45). There is >90% homology among the Foxa proteins at CRI, which functions as the DNA-binding domain. CRII and CRIII in the COOH terminus and CRIV in the NH<sub>2</sub> terminus are capable of distinct interactions with other transcription factor(s) and/or cofactor(s). Further investigation will clarify if EBP1 attenuates the transcriptional activity of Foxa by reducing the binding of Foxa proteins to the *AGR2* promoter via its interaction with the Foxa DNA-binding domain and/or by blocking the Foxa transactivation domain via interaction with other conserved regions. Because Foxa1 and Foxa2 have been found to be expressed in a wide range in prostate cancer and prostate cancer cell lines (ref. 19 and current data), we suggest that EBP1 may associate with Foxa1 and Foxa2 in a unique manner in different stages of prostate cancer.

Nevertheless, deregulation of both EBP1 (25) and Foxa (19) may contribute to invasion and distant spread of cancer cells at least in part by promoting expression of *AGR2*. Further studies focusing on this previously uncharacterized EBP1-Foxa-*AGR2* signaling circuit, especially in *in vivo* animal models, would identify new downstream targets with translational potential for early detection and therapy of prostate cancer.

## Disclosure of Potential Conflicts of Interest

No potential conflicts of interest were disclosed.

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