HEXIM1 Regulates 17\(^{\beta}\)-Estradiol/Estrogen Receptor-\(\alpha\)–Mediated Expression of Cyclin D1 in Mammary Cells via Modulation of P-TEFb

Ndiya Ogba, Laura J. Chaplin, Yong Qiu Doughman, Koh Fujinaga, and Monica M. Montano

\(^1\)Department of Pharmacology, \(^2\)Division of Infectious Diseases and Department of Molecular Biology and Microbiology, Case Western Reserve University; and \(^3\)Division of Pediatric Cardiology, Department of Pediatrics, Rainbow Babies and Children's Hospital, Cleveland, Ohio

Abstract

Estrogen receptor \(\alpha\) (ER\(\alpha\)) plays a key role in mammary gland development and is implicated in breast cancer through the transcriptional regulation of genes linked to proliferation and apoptosis. We previously reported that hexamethylene bisacetamide inducible protein 1 (HEXIM1) inhibits the activity of ligand-bound ER\(\alpha\) and bridges a functional interaction between ER\(\alpha\) and positive transcription elongation factor b (P-TEFb). To examine the consequences of a functional HEXIM1-ER\(\alpha\)-P-TEFb interaction in vivo, we generated MMTV/HEXIM1 mice that exhibit mammary epithelial-specific and doxycycline-inducible expression of HEXIM1. Increased HEXIM1 expression in the mammary gland decreased estrogen-driven ductal morphogenesis and inhibited the expression of cyclin D1 and serine 2 phosphorylated RNA polymerase II (S2P RNAP II). In addition, increased HEXIM1 expression in MCF-7 cells led to a decrease in estrogen-induced cyclin D1 expression, whereas down-regulation of HEXIM1 expression led to an enhancement of estrogen-induced cyclin D1 expression. Studies on the mechanism of HEXIM1 regulation on estrogen action indicate a decrease in estrogen-stimulated recruitment of ER\(\alpha\), P-TEFb, and S2P RNAP II to promoter and coding regions of ER\(\alpha\)-responsive genes \(pS2\) and \(CCND1\) with increased HEXIM1 expression in MCF-7 cells. Notably, increased HEXIM1 expression decreased only estrogen-induced P-TEFb activity. Whereas there have been previous reports on HEXIM1 inhibition of P-TEFb activity, our studies add a new dimension by showing that E\(_2\)/ER\(\alpha\) is an important regulator of the HEXIM1/P-TEFb functional unit in breast cells. Together, these studies provide novel insight into the role of HEXIM1 and ER\(\alpha\) in mammary epithelial cell function. [Cancer Res 2008;68(17):7015–24]

Introduction

Mammary gland morphogenesis and development requires input from several genetic and epigenetic pathways regulated by hormones and growth factors including estrogens (1, 2). Estrogens mediate their actions through estrogen receptors (ER), ER\(\alpha\) and ER\(\beta\), nuclear steroid receptors that classically regulate transcription either by directly binding to estrogen response elements of target genes (3–5) or indirectly via protein-protein interactions with other transcription factors like SP1 or activator protein 1 (AP-1; ref. 6). In both cases, coregulatory proteins are also recruited to the promoter, and together ERs and these factors elicit changes in mRNA and protein levels of ER target genes and, ultimately, a physiologic response (4–6). Because estrogen signaling controls the balance of growth and apoptosis in normal breast epithelial cells, a disruption of this balance contributes to abnormal cell growth and can lead to tumorigenesis (4, 7). Therefore, it is important to identify and elucidate the mechanism of action of ERs and their coregulators that give better insight into ER-mediated transcriptional regulation (8).

In eukaryotic transcription, the elongation stage is highly regulated and important for the generation of full-length mRNA transcripts (9–11). One of the positive regulators, positive transcription elongation factor b (P-TEFb), has an essential role in RNA polymerase II (RNAP II) transcription elongation (9, 10). In many human cell types, the predominant form of P-TEFb consists of cyclin-dependent kinase 9 (CDK9) and its regulatory partner, cyclin T1 (11). It phosphorylates and thereby inhibits the activity of negative elongation factors, NELF and DSIF (DRB sensitivity inducing factor; ref. 9). It also phosphorylates the COOH-terminal repeat domain (CTD) of the largest subunit of RNAP II (9, 10). The RNAP II CTD consists of multiple repeats of the heptapeptide sequence, YSPTSPS, phosphorylated at serine 5 by general transcription factor TFIIS during initiation and at serine 2 by P-TEFb during elongation (9, 10). These phosphorylation events are crucial for effective transition from an abortive to a productive phase of elongation (11, 12). P-TEFb is essential for productive HIV-1 transcriptional elongation and several studies have shown that various transcription factors bind to and recruit P-TEFb to specific promoters stimulating elongation (12, 13).

In previous studies, we identified an ER\(\alpha\)-interacting protein, estrogen down-regulated gene-1 [also known as hexamethylene inducible protein 1 (HEXIM1)], and found that it is an inhibitor of ER\(\alpha\) transcription and breast cell growth (14). Additionally, we showed that HEXIM1 expression was lower in human breast tumors when compared with adjacent normal tissue, suggesting a role for HEXIM1 in breast tumorigenesis (14). Concurrent studies identified HEXIM1 as a P-TEFb–interacting factor that also inhibits P-TEFb activity (15, 16). Studies have also shown that HEXIM2, a parologue of HEXIM1, has the same inhibitory effect on P-TEFb (17, 18). We showed that HEXIM1 modulates a functional interaction between ER\(\alpha\) and cyclin T1 in breast epithelial cells and inhibits the recruitment of ligand-bound ER\(\alpha\) (E\(_2\)/ER\(\alpha\)) to the pS2 gene promoter (19). We also found that cyclin T1 seemed to be
necessary for E2 induction of cyclin D1 protein expression (19). Cyclin D1 (CCND1) is a D-type cyclin that regulates G1-S cell cycle progression during cell proliferation (20). CCND1 is also E2/ERα responsive and is thought to play major roles in mammary gland development and breast cancer (1, 21). However, the contribution of E2/ERα to HEXIM1 action in breast cells is not well defined. In addition, the precise mechanism by which P-TEFb regulates CCND1 transcription and how this can be linked to mammary gland development and tumorigenesis need to be further defined.

To address these questions, we developed a transgenic mouse model in our laboratory that is doxycycline inducible and selectively expresses HEXIM1 in the mammary gland under the control of the mouse mammary tumor virus long terminal repeat (MMTV-LTR) promoter previously described (22). Using this model, we show that increased HEXIM1 expression decreased E2-induced CCND1 and serine 2 phosphorylated (S2P) RNAPII expression in modulating HEXIM1 expression on E2/ERα activity. These results elucidate the functional consequences of breast cancer.

Cyclin D1 (CCND1) is a D-type cyclin that regulates G1-S cell cycle progression and breast cancer (1, 21). However, the contribution of E2/ERα to HEXIM1 action in breast cells is not well defined. In addition, the precise mechanism by which P-TEFb regulates CCND1 transcription and how this can be linked to mammary gland development and tumorigenesis need to be further defined.

To address these questions, we developed a transgenic mouse model in our laboratory that is doxycycline inducible and selectively expresses HEXIM1 in the mammary gland under the control of the mouse mammary tumor virus long terminal repeat (MMTV-LTR) promoter previously described (22). Using this model, we show that increased HEXIM1 expression decreased E2-induced CCND1 and serine 2 phosphorylated (S2P) RNAPII expression in modulating HEXIM1 expression on E2/ERα activity. These results elucidate the functional consequences of breast cancer.

**Materials and Methods**

**Materials and antibodies.** 17β-Estradiol (E2) and CDK9 inhibitor 5,6-dichloro-1-β-ribonanosylbenzimidazole (DRB) were obtained from Sigma Chemical Co. Commercially available antibodies used for immunoprecipitation and Western blot analysis are described in Supplementary data.

**Transgenic mouse (MMTV/HEXIM1) generation.** MMTV-rtTA mice were generated as described by Gunther and colleagues (22). To generate pTET-HEXIM1 mice, a plasmid construct was made by subcloning the coding sequence of human HEXIM1 gene downstream of the tetracycline-dependent minimal promoter in the pTet-splice transgene construct. After purification, the resulting plasmid was used for pronuclear injection into FVB oocytes (Case Western Reserve University Transgenic and Targeting Facility). To achieve mammary gland–specific expression of HEXIM1 in a doxycycline-dependent manner, pTET-HEXIM1 mice were crossed with the MTB line, which expressed the reverse tetracycline-dependent transcriptional activator (rtTA) under the control of the MMTV-LTR. From these matings the bigenic mice, MMTV/HEXIM1, were created. Transgene expression was induced by adding 2 mg/mL doxycycline to the drinking water. Bigenic mice were identified by screening genomic DNA from tail biopsies for the presence of the transgenes using PCR and verified by Western blot analysis. See Supplementary Table S1 for the list and sequences of primers used.

**Whole-mount histology and immunodetection.** Mice were induced with doxycycline at 9 wk of age, ovariec-tomized (O VX), and treated with E2 at 12 wk. E2 was administered as daily s.c. injections of sesame oil solution containing 1 mg of E2. Mammary glands from the MMTV/HEXIM1 mice were collected 25 d after start of E2 treatment for whole-mount staining via the Carmine-alum technique and Western blots analyses. Immunohistochem-istry using sections from mammary glands are described in Supplementary data.

**Reverse transcription-PCR analyses.** Human breast epithelial cells, MCF-7, were maintained as described (19) and were transiently transfected with pCMV-TAG2B or pCMV-TAG2B-HEXIM1 using FuGENE HD reagent (Roche) according to the manufacturer’s instructions. Forty-eight hours later, cells were treated with ethanol vehicle or 100 nmol/L E2 for 3 h. Total RNA was extracted using the TRIzol reagent (Invitrogen) and subjected to reverse transcription-PCR (RT-PCR) analyses as described in Supplementary data.

**Western blot analyses.** Western blot analyses using extracts from mammary glands and MCF-7 cells are described in Supplementary data.

**RNA interference.** A Pol II promoter–driven miRNA expression vector system (Invitrogen) was used. To make pDNA-HEXIM1 miR, miRNA oligos (see Supplementary Table S3 for list and sequences) were annealed and cloned into the pCDNA 6.2 GW/EmGFP vector (Invitrogen) according to the manufacturer’s instructions. MCF-7 cells were transfected with pDNA 6.2-GW/EmGFP-miR expression vectors containing either the HEXIM1 miRNA insert or a control LacZ miRNA insert. Following blastidicin selection, cells expressing the highest level of green fluorescent protein (GFP) were flow-sorted and expanded. During experiments, cells were treated with ethanol vehicle or 100 nmol/L E2 for 3 h and harvested as described above for Western blot analyses.

**Chromatin immunoprecipitation assays.** MCF-7 cells were plated on 150-mm plates and transfected as described with pCMV-TAG2B or pCMV-TAG2B-HEXIM1. Forty-eight hours later, cells were treated with ethanol vehicle or 100 nmol/L E2 for 45 and 90 min. Chromatin immunoprecipitation assays were carried out as previously described (19). See Supplementary data for further details of immunoprecipitation and analyses.

**CTD kinase assays.** Kinase assays were done according to previously described protocols with some modifications (23, 24). See Supplementary data for complete description of assay.

**Data analyses.** Statistical significance was determined using Student’s t test comparison for unpaired data and was indicated as follows: *P < 0.05; **P < 0.005.

**Results**

Increase in HEXIM1 expression inhibits E2-driven mammary gland development by decreasing cell proliferation and increasing apoptosis. To examine the functional consequences of an interaction between HEXIM1, ERα, and P-TEFb in the mammary gland, we generated a double transgenic mouse model, MMTV/HEXIM1, which inducibly overexpress HEXIM1 in the mammary gland when the mice are treated with doxycycline (+DOX). We first examined the effects of elevated levels of HEXIM1 on E2-driven mammary gland development by having the MMTV/HEXIM1 mice ovariectomized and treated with E2 as described in Materials and Methods. In comparing whole mounts of mammary glands from −DOX and +DOX animal groups, we observed decreased ductal branching in the mammary glands of +DOX mice when compared with −DOX mice, as well as compared with single transgenics (MMTV alone; Fig. 1A, −DOX/+DOX, insets). We quantified the level of increase of HEXIM1 expression in the mammary gland as ~24% over endogenous levels (data not shown; see Fig. 2A for immunoblot), so we do not foresee that an overwhelming amount of HEXIM1 is needed to dictate these physiologic changes. Because ductal elongation and branching in the mammary gland have been shown to be E2/ERα dependent (1, 25), these data suggest that HEXIM1 inhibits E2/ERα–driven mammary gland morphogenesis.

Previously, we found that HEXIM1 inhibits ERα transcription (14, 19), so this physiologic effect could be due to a dysregulation of ER-responsive genes involved in proliferation and apoptosis. To investigate this, MMTV/HEXIM1 mice were injected with bromodeoxyuridine (BrdUrd) 2 hours before being sacrificed. BrdUrd-labeled nuclei in the mammary gland were detected by immunostaining and apoptotic nuclei were stained by terminal deoxynucleotidyl transferase–mediated dUTP nick end labeling (TUNEL). Quantitation of positively labeled mouse mammary gland development by decreasing cell proliferation and increasing apoptosis. To examine the functional consequences of an interaction between HEXIM1, ERα, and P-TEFb in the mammary gland, we generated a double transgenic mouse model, MMTV/HEXIM1, which inducibly overexpress HEXIM1 in the mammary gland when the mice are treated with doxycycline (+DOX). We first examined the effects of elevated levels of HEXIM1 on E2-driven mammary gland development by having the MMTV/HEXIM1 mice ovariectomized and treated with E2 as described in Materials and Methods. In comparing whole mounts of mammary glands from −DOX and +DOX animal groups, we observed decreased ductal branching in the mammary glands of +DOX mice when compared with −DOX mice, as well as compared with single transgenics (MMTV alone; Fig. 1A, −DOX/+DOX, insets). We quantified the level of increase of HEXIM1 expression in the mammary gland as ~24% over endogenous levels (data not shown; see Fig. 2A for immunoblot), so we do not foresee that an overwhelming amount of HEXIM1 is needed to dictate these physiologic changes. Because ductal elongation and branching in the mammary gland have been shown to be E2/ERα dependent (1, 25), these data suggest that HEXIM1 inhibits E2/ERα–driven mammary gland morphogenesis.

Previously, we found that HEXIM1 inhibits ERα transcription (14, 19), so this physiologic effect could be due to a dysregulation of ER-responsive genes involved in proliferation and apoptosis. To investigate this, MMTV/HEXIM1 mice were injected with bromodeoxyuridine (BrdUrd) 2 hours before being sacrificed. BrdUrd-labeled nuclei in the mammary gland were detected by immunostaining and apoptotic nuclei were stained by terminal deoxynucleotidyl transferase–mediated dUTP nick end labeling (TUNEL). Quantitation of positively labeled mouse mammary gland.
E2/ER prompted us to investigate the effect of HEXIM1 overexpression on ductal branching and cell proliferation in the mammary gland in vivo. Expression levels and serine 2 phosphorylation of RNAP II processes in the mammary gland.

Increased HEXIM1 expression inhibits estrogen-regulated mammary gland morphogenesis due to changes in proliferation and apoptosis. A, MMTV/HEXIM1 mice were treated as described in Materials and Methods. Representative whole mounts were obtained from MMTV and MMTV/HEXIM1 mice with Carminé alum stain. Original magnification, ×40. OVX, ovariectomized. B, BrdUrd-labeled nuclei were detected by immunostaining and apoptotic nuclei were stained by TUNEL. Quantitation of positively labeled epithelial cell nuclei from at least 1,000 nuclei each from five mice per group (–DOX/+DOX). Original magnification, ×40.

epithelial cell nuclei revealed both a decrease in epithelial cell proliferation and an increase in apoptosis (Fig. 1B). Taken together, our data suggest a critical role for HEXIM1 in E2/ERα-driven processes in the mammary gland.

Increased HEXIM1 expression regulates cyclin D1 protein expression levels and serine 2 phosphorylation of RNAP II in vivo. Our findings that overexpression of HEXIM1 decreased ductal branching and cell proliferation in the mammary gland prompted us to investigate the effect of HEXIM1 overexpression on E2/ERα signaling. As an output, we examined changes in cyclin D1 (CCND1) and c-Myc because both genes are E2/ERα responsive and involved in proliferation integral to mammary gland development and breast cancer (26, 27). MMTV/HEXIM1 mice were ovariectomized and treated with E2 as described, and we found that increased HEXIM1 expression (+DOX) resulted in decreased CCND1 protein expression levels (~67%) in mouse mammary gland cell extracts (Fig. 2A). However, c-Myc expression levels did not significantly change regardless of HEXIM1 expression (Supplementary Fig. S1A), suggesting a difference in sensitivity of ERα target genes to HEXIM1.

In addition, because HEXIM1 inhibits P-TEFb activity (12), we examined the effect of increased HEXIM1 expression on P-TEFb activity in the mouse mammary gland by investigating the expression levels of the S2P RNAP II. In +DOX mice, we observed a decrease in S2P RNAP II levels (~68%; Fig. 2A). It is important to note here that the antibody used to detect S2P RNAP II recognizes both phosphorylated (RNAP IIo) and unphosphorylated (RNAP IIa) forms of RNAP II. We also blotted with another antibody that detects only the unphosphorylated form (Hypo RNAP II) and found that there was no observable change between –DOX and +DOX mice groups (Fig. 2A). In addition, HEXIM2, a parologue of HEXIM1, inhibits P-TEFb activity (12), but in the mammary gland, HEXIM1 protein expression levels are significantly higher (Supplementary Fig. S1B), suggesting a dominant role for HEXIM1 in the mammary gland.

We also examined the expression levels of the serine 5 phosphorylated form of RNAP II (S5P RNAP II), associated with initiation, using immunofluorescent labeling of epithelial nuclei in mammary gland lumen from MMTV/HEXIM1 mice. We did not observe any changes in the percentage of S5P RNAP II positively stained nuclei (S5P +ve cells) when we compared –DOX and +DOX animal groups (Fig. 2B). However, in the +DOX animal group, the percentage of S2P RNAP II positively stained nuclei (S2P +ve cells) within the mammary gland lumen was significantly decreased by 50% (P < 0.05) when compared with the –DOX group (Fig. 2B), verifying our results in the Western blot (Fig. 2A). Given that E2/ERα regulates CCND1 expression (21, 27) and HEXIM1 inhibits P-TEFb activity and associates with ERα (12, 19), the effect of increased HEXIM1 expression levels on CCND1 and S2P RNAP II expression suggests that these changes reflect a complex interaction of multiple pathways that converge at the level of E2/ERα-mediated transcription.

HEXIM1 regulates cyclin D1 and pS2 expression levels in vitro. The effects of increased HEXIM1 expression on CCND1 and S2P RNAP II expression levels could also be the sum result of the disruption of E2/ERα activity in multiple cell types in the mammary gland. To verify that our observations can be attributed to a more localized event in epithelial cells, we investigated the effects of increased HEXIM1 expression in mammary epithelial MCF-7 cells. MCF-7 cells were transfected with pCMV-Tag2B-HEXIM1 or control vector and treated with ethanol vehicle or E2 for 3 hours. In control cells, we observed an average 1.5- to 2-fold E2 induction of both CCND1 mRNA and protein levels (Fig. 3A and B), consistent with what has been shown in other studies (8, 21). As expected, increased HEXIM1 expression diminished E2 induction of CCND1 mRNA and protein expression (Fig. 3A and B, compare lanes 2 and 4). For CCND1 protein expression levels, this was quantified as a 52% decrease in expression by HEXIM1 when normalized to glyceraldehyde 3-phosphate dehydrogenase (GAPDH) expression (Fig. 3B).
We also investigated the effect of increased HEXIM1 expression on other ER-responsive genes and found that increased HEXIM1 levels decreased E2-induced increases in pS2 mRNA and protein levels (Fig. 3A and B; Supplementary Fig. S2A). The E2-induced mRNA levels of other genes including c-Myc, progesterone receptor, and cathepsin D remained unchanged with increased HEXIM1 expression (Fig. 3A; Supplementary Fig. S2A). In addition, E2 induction of c-Myc protein levels was also unchanged with increased HEXIM1 expression (Fig. 3B, with quantitation in Supplementary Fig. S2B). These data also suggest that increased HEXIM1 expression may not have a similar effect on all ERα target genes.

Having observed that increased HEXIM1 expression decreased E2-induced CCND1 and pS2 expression, it stood to reason that HEXIM1 silencing would result in an increase in the induction of both genes by E2 when compared with control cells. HEXIM1 gene expression knockdown (~50%) using miRNA-mediated RNA interference resulted in a statistically significant (P < 0.01)
enhancement of E2-induced CCND1 protein expression levels (Fig. 3C). We did not observe a similar effect with pS2 (data not shown) and E2-induced c-Myc protein expression levels remained unchanged (Fig. 3C). Because multiple regulatory proteins are involved in CCND1 regulation (28), we cannot rule out possible actions of other factors in vivo. In addition, we cannot rule out the possibility of HEXIM1 regulation of other key E2 responsive genes involved in cell proliferation and mammary carcinogenesis not studied here. Nonetheless, taken together, our data support a novel physiologic role for HEXIM1 in E2-driven mammary gland development via regulation of CCND1 levels and serine 2 phosphorylation of RNAP II.

HEXIM1 inhibits E2-induced recruitment of ERα and cyclin T1 (P-TEFb) to ERα-responsive genes. Given our observations on the effect of HEXIM1 expression on CCND1 and pS2, it is critical to characterize the molecular events involved in HEXIM1 transcriptional regulation of these ER-responsive genes. We previously reported that HEXIM1 associates with ERα after 90 minutes of E2 stimulation in MCF-7 cells, and increased HEXIM1 expression led to a decrease in E2/ERα recruitment to the pS2 promoter (19). Other reports have shown that P-TEFb associates with the elongating form of RNAP II, S2P RNAP II, which is thought to predominate in the coding regions of genes with increased occupancy toward the 3′ end of genes (29–31). Therefore, we
hypothesized that increased HEXIM1 levels would (a) inhibit the recruitment of ERα to the promoter region of ER target genes and (b) inhibit the recruitment of P-TEFb to regions downstream from the promoter at ER-responsive genes, and this would in turn verify the potential regulation of transcriptional elongation of these genes. To investigate this, we carried out chromatin immunoprecipitation assays in MCF-7 cells and examined the effect of E2 on ERα and P-TEFb occupancy at two ERα-responsive genes, pS2 and CCND1.

We found that an increase in HEXIM1 expression in MCF-7 cells correlated with a 2-fold increase in HEXIM1 occupancy at E2-responsive regions within the pS2 and CCND1 (32) promoters (Supplementary Fig. S3A), which was not significantly affected by E2 treatment. We also found that increased HEXIM1 expression inhibited the recruitment of E2/ERα to the promoter regions of pS2 and CCND1 (Fig. 4B). Now, it is well documented that E2/ERα cycles on and off the promoter of ER-responsive genes, and ERα binding is typically diminished after 90 minutes of E2 stimulation (33, 34). Other studies have also shown that the absolute timing of ERα cycling differs and there can still be significant E2/ERα enrichment at 90 minutes (35, 36). In our studies, it is clear that E2/ERα binding is less at 90 minutes when compared with 45 minutes.
HEXIM1 and P-TEFb Regulation Decreases E2-Driven Proliferation

Figure 4. Increased HEXIM1 expression inhibits E2-induced P-TEFb activity and recruitment of serine 2 (hyperphosphorylated) RNAPII to the coding region of ER-responsive genes. A, increased HEXIM1 expression decreased E2-induced CTD4 peptide phosphorylation. MCF-7 cells were transiently transfected with pCMV-Tag2B-HEXIM1 or empty vector and treated with ethanol vehicle or 100 nmol/L E2 for 90 min. Cell lysates were subjected to immunoprecipitation with antibodies against cyclin T1 and rabbit IgG (as an immunoprecipitation control). The immunoprecipitates were divided into two halves with one half getting 2 μg of the CTD4 peptide added to the reaction (−CTD4/+CTD4). DRB at 50 μmol/L was also added to some immunoprecipitates as a kinase assay control. The kinase reactions were analyzed by SDS-PAGE using autoradiography. Equal volumes of kinase reactions were also analyzed by Western blot to check specificity of anti-cyclin T1 antibody in immunoprecipitation. Representative of at least four independent experiments. B, HEXIM1 inhibits E2-dependent recruitment of S2P RNAPII to pS2 ORF. MCF-7 cells were treated as described in Fig. 4 and subjected to chromatin immunoprecipitation analysis with antibodies against S2P RNAPII, S5P RNAPII, and the unphosphorylated form of RNAPII (8WG16). The regions of pS2 amplified by PCR are as indicated. Columns, mean of three to five independent replicates; bars, SE. **, P < 0.005. C, HEXIM1 inhibits E2-dependent recruitment of S2P RNAPII to CCND1 ORF. MCF-7 cells were treated as described. The regions of CCND1 amplified by PCR are as indicated. Columns, mean of three to five independent replicates; bars, SE. *, P < 0.05.  

HEXIM1 and P-TEFb Regulation Decreases E2-Driven Proliferation

Figure 5. Increased HEXIM1 expression inhibits E2-induced P-TEFb activity and recruitment of serine 2 (hyperphosphorylated) RNAPII to the coding region of ER-responsive genes. A, increased HEXIM1 expression decreased E2-induced CTD4 peptide phosphorylation. MCF-7 cells were transiently transfected with pCMV-Tag2B-HEXIM1 or empty vector and treated with ethanol vehicle or 100 nmol/L E2 for 90 min. Cell lysates were subjected to immunoprecipitation with antibodies against cyclin T1 and rabbit IgG (as an immunoprecipitation control). The immunoprecipitates were divided into two halves with one half getting 2 μg of the CTD4 peptide added to the reaction (−CTD4/+CTD4). DRB at 50 μmol/L was also added to some immunoprecipitates as a kinase assay control. The kinase reactions were analyzed by SDS-PAGE using autoradiography. Equal volumes of kinase reactions were also analyzed by Western blot to check specificity of anti-cyclin T1 antibody in immunoprecipitation. Representative of at least four independent experiments. B, HEXIM1 inhibits E2-dependent recruitment of S2P RNAPII to pS2 ORF. MCF-7 cells were treated as described in Fig. 4 and subjected to chromatin immunoprecipitation analysis with antibodies against S2P RNAPII, S5P RNAPII, and the unphosphorylated form of RNAPII (8WG16). The regions of pS2 amplified by PCR are as indicated. Columns, mean of three to five independent replicates; bars, SE. **, P < 0.005. C, HEXIM1 inhibits E2-dependent recruitment of S2P RNAPII to CCND1 ORF. MCF-7 cells were treated as described. The regions of CCND1 amplified by PCR are as indicated. Columns, mean of three to five independent replicates; bars, SE. *, P < 0.05.  

HEXIM1 and P-TEFb Regulation Decreases E2-Driven Proliferation

Figure 6. Increased HEXIM1 expression inhibits E2-induced P-TEFb activity and recruitment of serine 2 (hyperphosphorylated) RNAPII to the coding region of ER-responsive genes. A, increased HEXIM1 expression decreased E2-induced CTD4 peptide phosphorylation. MCF-7 cells were transiently transfected with pCMV-Tag2B-HEXIM1 or empty vector and treated with ethanol vehicle or 100 nmol/L E2 for 90 min. Cell lysates were subjected to immunoprecipitation with antibodies against cyclin T1 and rabbit IgG (as an immunoprecipitation control). The immunoprecipitates were divided into two halves with one half getting 2 μg of the CTD4 peptide added to the reaction (−CTD4/+CTD4). DRB at 50 μmol/L was also added to some immunoprecipitates as a kinase assay control. The kinase reactions were analyzed by SDS-PAGE using autoradiography. Equal volumes of kinase reactions were also analyzed by Western blot to check specificity of anti-cyclin T1 antibody in immunoprecipitation. Representative of at least four independent experiments. B, HEXIM1 inhibits E2-dependent recruitment of S2P RNAPII to pS2 ORF. MCF-7 cells were treated as described in Fig. 4 and subjected to chromatin immunoprecipitation analysis with antibodies against S2P RNAPII, S5P RNAPII, and the unphosphorylated form of RNAPII (8WG16). The regions of pS2 amplified by PCR are as indicated. Columns, mean of three to five independent replicates; bars, SE. **, P < 0.005. C, HEXIM1 inhibits E2-dependent recruitment of S2P RNAPII to CCND1 ORF. MCF-7 cells were treated as described. The regions of CCND1 amplified by PCR are as indicated. Columns, mean of three to five independent replicates; bars, SE. *, P < 0.05.  

HEXIM1 and P-TEFb Regulation Decreases E2-Driven Proliferation

Figure 7. Increased HEXIM1 expression inhibits E2-induced P-TEFb activity and recruitment of serine 2 (hyperphosphorylated) RNAPII to the coding region of ER-responsive genes. A, increased HEXIM1 expression decreased E2-induced CTD4 peptide phosphorylation. MCF-7 cells were transiently transfected with pCMV-Tag2B-HEXIM1 or empty vector and treated with ethanol vehicle or 100 nmol/L E2 for 90 min. Cell lysates were subjected to immunoprecipitation with antibodies against cyclin T1 and rabbit IgG (as an immunoprecipitation control). The immunoprecipitates were divided into two halves with one half getting 2 μg of the CTD4 peptide added to the reaction (−CTD4/+CTD4). DRB at 50 μmol/L was also added to some immunoprecipitates as a kinase assay control. The kinase reactions were analyzed by SDS-PAGE using autoradiography. Equal volumes of kinase reactions were also analyzed by Western blot to check specificity of anti-cyclin T1 antibody in immunoprecipitation. Representative of at least four independent experiments. B, HEXIM1 inhibits E2-dependent recruitment of S2P RNAPII to pS2 ORF. MCF-7 cells were treated as described in Fig. 4 and subjected to chromatin immunoprecipitation analysis with antibodies against S2P RNAPII, S5P RNAPII, and the unphosphorylated form of RNAPII (8WG16). The regions of pS2 amplified by PCR are as indicated. Columns, mean of three to five independent replicates; bars, SE. **, P < 0.005. C, HEXIM1 inhibits E2-dependent recruitment of S2P RNAPII to CCND1 ORF. MCF-7 cells were treated as described. The regions of CCND1 amplified by PCR are as indicated. Columns, mean of three to five independent replicates; bars, SE. *, P < 0.05.
consistent with other studies showing that the recruitment of P-TEFb to promoter-proximal and coding regions stimulates transcriptional elongation (30, 38). In addition, in previous studies, we observed a decrease in the co-recruitment of cyclin T1 with ERα at the pS2 promoter with increased HEXIM1 expression (19), but it seems that the total occupancy of cyclin T1 at the promoter region does not change with increased HEXIM1 expression (Fig. 4C). In this context, our data suggest a dual role for P-TEFb in ERα transcription. On one hand, it acts as a coactivator for ERα-driven transcription by directly associating with E2/ERα (19), but it also serves its general purpose as a transcription elongation factor, with E2 acting as an enhancer of P-TEFb occupancy in the context of ER-responsive genes.

HEXIM1 inhibits E2-induced P-TEFb activity and recruitment of S2P RNAP II to coding regions of ER-responsive genes. Because E2 enhanced P-TEFb recruitment to pS2 and CCND1 genes, we investigated if E2 also increased P-TEFb activity as a means of promoting transcriptional elongation of ERα-responsive genes. To do this, we conducted kinase assays using endogenous immunoprecipitates of cyclin T1 from MCF-7 cells and examined the kinase activity of P-TEFb with a synthetic peptide substrate, CTD4 (YSPTSPS4). As shown in Fig. 5A, E2 treatment increased P-TEFb activity in MCF-7 cells (compare lanes 2 and 5). We also observed that increased HEXIM1 expression in MCF-7 cells inhibited E2-induced increase in P-TEFb activity (Fig. 5A, compare lanes 2 and 5 with lanes 8 and 11). We confirmed that this inhibition was selective for P-TEFb by adding the CDK9 inhibitor DRB to an equal half of the kinase immunoprecipitates (Fig. 5A, lanes 3, 6, 9, and 12) and that the input levels of cyclin T1 and HEXIM1 were evenly loaded (Supplementary Fig. S4A). Quantitation of 32P incorporation into the CTD4 peptide verified that HEXIM1 significantly abrogates E2-induced P-TEFb activity in MCF-7 cells (Supplementary Fig. S4B).

We also observed that increased HEXIM1 expression did not inhibit basal P-TEFb activity in HEXIM1-transfected MCF-7 cells compared with nontransfected cells (Fig. 5A, compare lanes 2 and 8). One reason for this could be attributed to the fact that at basal levels, the P-TEFb complex is thought to occur in two states, 50% free and 50% HEXIM1-7SK RNA bound (12), and in a study by He and colleagues (11), gradually decreasing HEXIM1 expression by RNA interference did not initially have much effect on this equilibrium because it was the “free” form of HEXIM1 that was being down-regulated, without any effect on the HEXIM1 protein associated with P-TEFb. In our experiments, it is possible that increasing HEXIM1 expression may not be significantly perturbing the P-TEFb equilibrium initially, but increases the availability of free HEXIM1 populations, which subsequently diminishes any increases in E2-induced P-TEFb activity.

Because HEXIM1 inhibited the recruitment of cyclin T1 to the coding regions of ER-responsive genes and E2-induced P-TEFb activity, we determined the effect of increased HEXIM1 levels on the recruitment of S2P RNAP II as a mark of the modulation of transcription elongation (9). At the pS2 promoter, we found that E2 stimulated the enrichment of all forms of RNAPII, without any significant changes when HEXIM1 expression was increased (Fig. 5B). This result differs from what has been observed in some studies, which show low S2P RNAP II occupancy in comparison with S5P RNAP II at the promoter of transcriptionally active genes (29, 37). However, it is possible for both S2P and S5P RNAP II to occupy similar regions on DNA (39), which likely marks the beginning of the transition to elongation, but it is unclear because we did not study proximal upstream or downstream regions to the pS2 promoter. In addition, there was no change in the recruitment of all forms of RNAPII to the CCND1 promoter in both control and HEXIM1-transfected cells, although it was not as sensitive as the pS2 promoter to E2 stimulation (Fig. 5C).

However, at the pS2 and CCND1 coding regions, we found that increased HEXIM1 expression inhibited the E2-dependent recruitment of S2P RNAP II, without significant changes in the recruitment of S5P and unphosphorylated RNAPII forms (Fig. 5B and C). These data indicate that in mammary epithelial cells, HEXIM1 does not affect transcription initiation because the recruitment patterns of all forms of RNAPII to the promoter regions of pS2 and CCND1 were unchanged regardless of HEXIM1 expression levels in the cell. However, increased HEXIM1 decreases transcription elongation because the recruitment of S2P RNAP II to the coding region was decreased.

Discussion

This study provides novel evidence for a physiologic role of HEXIM1/P-TEFb interaction in attenuating E2/ERα driven transcription in the mammary gland and breast cancer cells. First, we showed that increased HEXIM1 expression in the mammary gland of a transgenic mouse model decreased ductal branching, an E2-driven developmental process, due to changes in proliferation and apoptosis. We correlated these changes with a decrease in CCND1 and S2P RNAPII expression in vivo and in vitro. We also show that overexpression of HEXIM1 diminished E2-induced recruitment of ERα and cyclin T1 to the promoter and coding regions, respectively, of ER-responsive genes. Further, we show that E2 enhances the activity of the P-TEFb kinase CDK9, which is inhibited by increased HEXIM1 expression. Surprisingly, increased HEXIM1 expression inhibited only E2-induced increases in P-TEFb activity and not basal P-TEFb kinase activity. In our studies, this
inhibition of P-TEFb activity translates to a decrease in the recruitment of S2P RNAP II, and not other forms of RNAP II, to the coding regions of the ER-responsive genes pS2 and CCND1. These findings support a role for P-TEFb and transcription elongation in cell proliferation, but more importantly, the data suggest a novel mechanism of action for HEXIM1 that can be recapitulated in vivo and a possible therapeutic role for HEXIM1 in hormone-dependent breast cancer.

Given that the regulation of CCND1 is complex (28), we do not assume that our findings represent a comprehensive explanation about E2-regulation of CCND1. In addition, other sites within CCND1 contribute to the transcriptional output (21). We investigated the recruitment patterns of ERα, P-TEFb, and RNAP II to two sites within the gene: an E2-responsive region of the promoter and the coding region. Perhaps, a more extensive analysis of different sites within the genes pS2 and CCND1 and even other ER-responsive genes will reveal other insights into the mechanism of HEXIM1 regulation of these genes. However, we believe that the information gathered from this study was sufficient to show the regulatory effects of HEXIM1 on ERα and P-TEFb recruitment to pS2 and CCND1, suggesting a role for HEXIM1 and P-TEFb in ERα transcriptional regulation of some, but not all, ERα target genes.

Based on our previous (19) and current studies, we speculate that E2 enhances an ERα-P-TEFb interaction, and this increases the population of active P-TEFb at the gene locus of ER-responsive genes, which in turn phosphorylates the CTD of RNAP II. This phosphorylation event positions the gene in an active elongation state with increased S2P RNAP II occupancy at the coding regions and enhanced recruitment of other forms of RNAP II marking the gene in an active transcription state (31, 39). However, increased HEXIM1 expression inhibits ERα and P-TEFb enrichment at the promoter and coding regions, respectively, of ER-responsive genes, thus decreasing the population of P-TEFb available to phosphorylate RNAP II to the serine 2 phosphorylated form associated with transcriptional elongation. In addition, we observed that increased HEXIM1 inhibits E2-induced P-TEFb activity and postulated that this was due to an increase in the “free” form of HEXIM1, which serves to diminish any subsequent increases in P-TEFb activity. Taken together, this scenario could represent the mechanism by which HEXIM1 modulates ERα-mediated transcription in the context of some ER-responsive genes (see Fig. 6 for proposed model).

The understanding of general mechanisms that control elongation stems from studies showing that HIV-1 harnesses P-TEFb as a cofactor to promote efficient mRNA transcript synthesis (12). These and other studies have raised questions about whether P-TEFb acts as a general transcription elongation factor or serves in a gene-selective or context-dependent manner. The P-TEFb complex components, cyclin T1 and CDK9, have not been shown to have sequence-specific DNA binding activity, but transcription factors interact with and recruit P-TEFb to their respective promoter targets (12, 40, 41). In addition, DNA microarray analyses of hearts from cyclin T1 transgenic mice indicate selective increases in subsets of genes, rather than a global increase in mRNA expression, when compared with nontransgenic mice (42). Our studies suggest that in breast epithelial cells, P-TEFb can be modulated by E2/ERα and HEXIM1 in the context of some ERα target genes, although identical E2-induced recruitment patterns for P-TEFb to pS2 and CCND1 genes suggest that a general transcription elongation mechanism is also involved. In addition, the interaction of P-TEFb and E2/ERα supports a positive aspect of ERα transcription elongation regulation, but E2/ERα also interacts with negative elongation factor (NELF) and this interaction inhibits ERα-mediated transcription (43).

HEXIM1 has been shown to have P-TEFb–independent action as seen with the glucocorticoid receptor (44). We have also reported on P-TEFb–independent action of HEXIM1 in cardiovascular development (45). Given this evidence, it is clear that HEXIM1 can inhibit transcription in a P-TEFb–dependent and P-TEFb–independent manner. Therefore, we cannot assume that the effect of HEXIM1 in the mammary gland is solely on ERα/ P-TEFb because other factors are involved in mammary gland development. However, we were able to show a specific inhibition pattern that HEXIM1 exerted on E2-induced events at ER-responsive genes, pS2 and CCND1, and increased HEXIM1 levels inhibited E2-induced P-TEFb activity. Thus, based on our data, the HEXIM1 inhibition patterns observed in this study are largely P-TEFb dependent in both our cell culture and animal models. Several studies also support an emerging role for HEXIM1 as a regulator of cell growth and differentiation (12, 46). Conversely, deletion of CLP-1, the mouse HEXIM1 gene, leads to pathologic cardiac hypertrophy and perinatal death (47).

In this study, a targeted increase in HEXIM1 expression in the mouse mammary gland driven by a mammary epithelial cell promoter (MMTV-LTR) led to a decrease in ductal branching (Fig. 1A), an E2/ERα–driven mammary gland developmental process (25). This observation was attributed to a decrease in proliferation and an increase in apoptosis (Fig. 1B). The decrease in proliferation was linked to a concurrent decrease in CCND1 expression, further showing that HEXIM1 regulates ER-responsive genes in vivo. Future studies will aim to address not only HEXIM1 regulation of other E2/ERα target genes but also HEXIM1 regulation of other nuclear receptors relevant in mammary cell function and tumorigenesis.

Disclosure of Potential Conflicts of Interest
No potential conflicts of interest were disclosed.

Acknowledgments
Received 3/3/2008; revised 6/24/2008; accepted 6/26/2008.

Grant support: NIH grant CA92440 (M.M. Montano) and Department of Defense Predoctoral Fellowship W81XWH-06-1-0126 (N. Ogba).

The costs of publication of this article were defrayed in part by the payment of page charges. This article must therefore be hereby marked advertisement in accordance with 18 U.S.C. Section 1734 solely to indicate this fact.

We thank Dr. Huayun Deng for his help with some of the animal work and advice on immunohistochemistry, and Dr. Lewis Chodosh (University of Pennsylvania) for the MMTB mice and the pTet-Splice vector.

References
3. Bardin A, Bouille N, Lazenec G, Vignon F, Pujol P. Loss of ERα expression as a common step in
Cancer Research


19. Wittmann BM, Fujinaga K, Deng H, Ogha N, Montano MM. The breast cell growth inhibitor, estrogen down-regulated gene I, modulates a novel functional interac-


30. Ahn SH, Kim M, Buratowski S. Phosphorylation of serine 2 within the RNA polymerase II C-terminal domain affects transcription factor IIH and RNA polymerase II from serine 2 within the RNA polymerase II C-terminal domain couples transcription and 3' end processing. Mol Cell 2004;13:867–76.


