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PRINCIPAL INVESTIGATOR: Donald B. DeFranco, Ph.D.

CONTRACTING ORGANIZATION: University of Pittsburgh  
Pittsburgh, Pennsylvania 15260

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| <b>7. PERFORMING ORGANIZATION NAME(S) AND ADDRESS(ES)</b><br>University of Pittsburgh<br>Pittsburgh, Pennsylvania 15260<br><br><i>E-Mail:</i> dodl@pitt.edu                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               |                                                                     |                                                                    | <b>8. PERFORMING ORGANIZATION<br/>REPORT NUMBER</b>                          |                                  |
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## **Introduction**

Androgens are required for the growth and development of the normal prostate gland (1) but also participate in the progression of prostate cancer, particularly in early stages (2). The principal mediator of androgen action within cells is the androgen receptor (AR), a member of a large super-family of nuclear receptors (3,4). Physiological effects of androgens result primarily from transcriptional regulation of specific target genes brought about by AR interaction with specific DNA sequences, or receptor associations with other transcription factors (5-7) or coactivators (8). Many coactivator proteins have been identified that interact with AR and influence the expression of androgen regulated genes. The mechanism of action of AR coactivators can be diverse and include chromatin modification or direct interactions with the basal transcription machinery (8). One group of coactivators that appear to have a unique mechanism of action includes members of a group III LIM domain family related to paxillin (9,10,11). These proteins (including paxillin and Hic-5/ARA55) reside both within the nucleus and at specialized structures at the plasma membrane termed focal adhesions (10,11) and may exert direct and indirect effects on AR transactivation responding to both androgen signaling and non-steroidal signals that are transmitted through focal adhesions (12). In this proposal, the impact of paxillin effects on AR transactivation has been addressed in order to provide insights into the integration of extracellular signals that initiate at the plasma membrane with AR action in the nucleus to impact androgen regulated gene expression in prostate cells.

## **Body**

In our previous progress report we presented preliminary work that provided support for our hypothesis that paxillin can function as an AR coactivator in prostate cancer cell lines. These studies have now been published (11) and the attached manuscript provides detailed descriptions of these experiments. To summarize the key results of these studies, we have shown that (i) paxillin is a

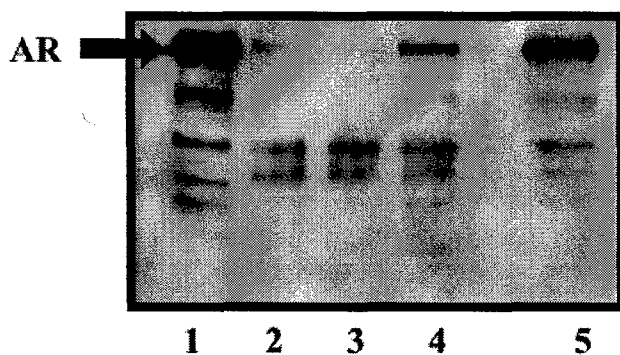
coactivator for AR in prostate cancer cell lines, (ii) paxillin directly interacts with AR in a hormone-independent manner *in vitro* through its carboxyl-terminal LIM domain and (iii) a fraction of paxillin in prostate cancer cell lines and prostate cancer tissue is associated with the nuclear matrix.

While *in vitro* binding assays with a GST-paxillin fusion protein established the direct interaction between paxillin and AR, we set out to show whether this complex could be detected in prostate cancer cells. Initial attempts to isolate paxillin/AR complexes by co-immunoprecipitation using standard conditions were unsuccessful. We therefore adapted our immunoprecipitation assay to include a crosslinking step. As shown in Figure 1, we were able to detect a complex between AR and paxillin in lysates prepared from LNCaP cells. Importantly, AR/paxillin complex formation in the LNCaP cells was hormone-dependent. Therefore, the hormone-independent interaction between paxillin and AR observed in GST-pull down assays with bacterially expressed paxillin protein (see attached manuscript) is not reflective of the hormone-dependence of paxillin-AR interactions in prostate cancer cell lines.



**Fig. 1: Hormone-dependent interaction between paxillin and AR in LNCaP cells.** An immunoprecipitation assay was performed with either control mouse serum (lanes 1,2) or a mouse monoclonal anti-AR antibody (lanes 3,4) using cytosol collected from LNCaP cells that were either untreated (lanes 1, 3) or treated (lanes 2,4) with 100 nM DHT for 2 hours. All extracts were exposed to the crosslinking reagent DSP prior to immunoprecipitation. Following collection of the immunoprecipitates, paxillin was detected by Western blot analysis. Lane 5 shows control LNCaP extract not subjected to immunoprecipitation. Note the hormone-dependent detection of paxillin (lane 4 versus 3) only in precipitates recovered with the anti-AR antibody.

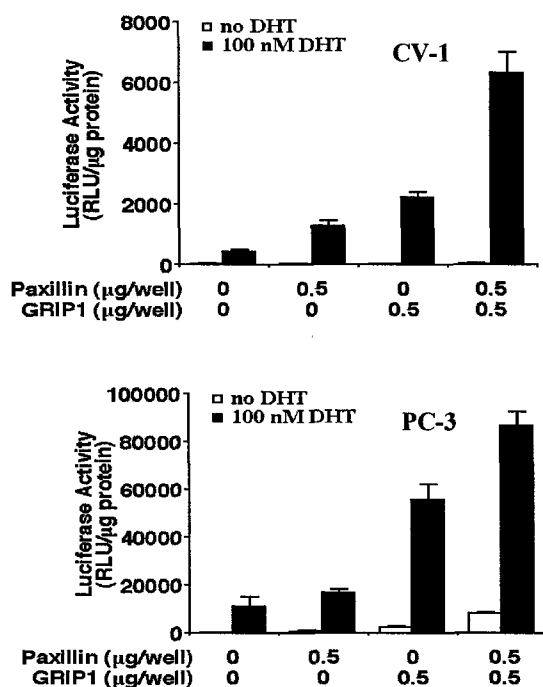
In addition to demonstrating an interaction between AR and paxillin in soluble extracts from LNCaP cells, we have performed co-IP experiments with membrane fractions prepared from these cells. The purpose of these studies was to reveal whether a coupling between paxillin and AR could be initiated at the plasma membrane where paxillin is known to respond to various signaling molecules. As shown in Figure 2, AR and paxillin are recovered in complexes contained in membrane fractions isolated from LNCaP cells. In this case, the association between AR and paxillin is transient as prolonged androgen treatment leads to a loss of AR /paxillin interactions initiated soon after hormone treatment of cells. This time course of AR/paxillin interactions at the plasma membrane suggests that paxillin effects on androgen regulated gene expression may be multi-faceted initiating soon after hormone treatment to promote an interaction between AR and paxillin at the plasma membrane and culminating in direct interactions with the receptor in the nucleus, perhaps at specific AR-regulated target genes.



**Fig. 2: Hormone-dependent interaction between paxillin and AR in membrane fractions from LNCaP cells.** An immunoprecipitation assay was performed with a rabbit polyclonal anti-paxillin antibody (lanes 2-5) using membrane fractions collected from LNCaP cells that were either untreated (lane 5) or treated with 100 nM DHT for 10 min (lane 2), 1 hr (lane 3), or 24 hr (lane 4). Lane 5 shows control LNCaP soluble cytosolic extract not subjected to immunoprecipitation. Following collection of the immunoprecipitates, AR was detected by Western blot analysis. Note the transient association of AR with paxillin in hormone treated cells and the efficient association between paxillin and AR in membrane fractions from non-hormone treated cells.

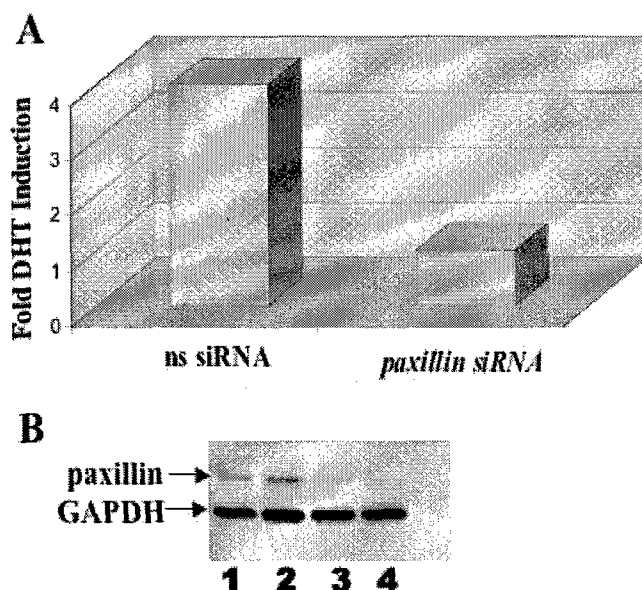
The influence of paxillin effects on AR action in the nucleus was shown in experiments shown in Figure 3 where paxillin was found to cooperate with a member of the p160 family of coactivators (i.e. GRIP1) to synergistically enhance AR transactivation both in CV-1 fibroblast cells (top) and PC-3 prostate cancer cells (bottom). This result could have implications for AR activity in developing prostate cancer as increases in the levels or activity of distinct coactivators (i.e. GRIP1 and paxillin) could act together

to elevate AR transactivation activity beyond some threshold level required to induce or sustain prostate cancer cell survival.



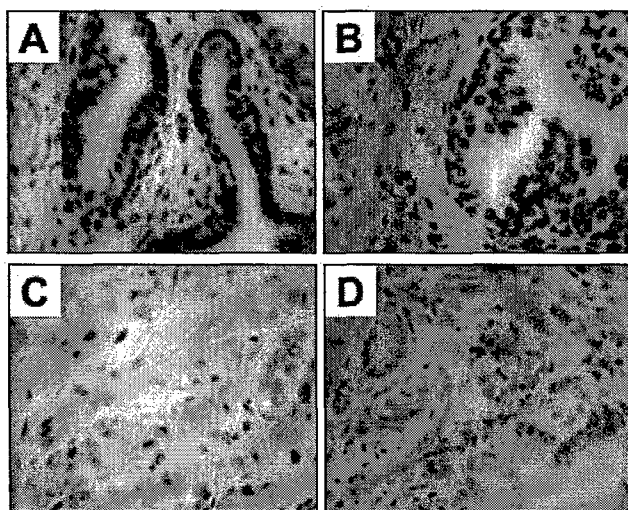
**Fig. 3. Enhancement of hormone-induced AR transactivation by paxillin and GRIP1.** CV-1 cells (top) or PC-3 cells (bottom) were transiently transfected with the indicated amounts of paxillin or GRIP1 expression plasmids, a MMTV promoter-linked luciferase reporter plasmid and an AR expression plasmid. Cells were incubated in the presence of absence of 100 nM DHT for 24 hours. Histogram shows the mean  $\pm$  SD (N=4) relative luciferase activity normalized to total protein levels in each sample.

Since our initial attempts to use anti-sense RNA to ablate paxillin expression were unsuccessful, we decided to alter our strategy and use siRNA against paxillin. In a recent report, Chang's group has used an siRNA approach to reduce expression of an AR coactivator closely related to paxillin, Hic-5/ARA55 (13). Using an analogous strategy focusing on a region of paxillin that has little homology with Hic-5/ARA55 we have been able to selectively ablate paxillin in LNCaP cells (Figure 4B). AR-mediated transactivation in LNCaP cells is reduced upon paxillin ablation (Figure 4A). Thus, paxillin is required for maximal AR-induced transcription in a PC cell line. Examination of the impact of paxillin on HER-2/Neu effects on AR transactivation have not been successful but may be addressed more clearly with the newly developed siRNA approach to ablate paxillin expression in PC cells lines.

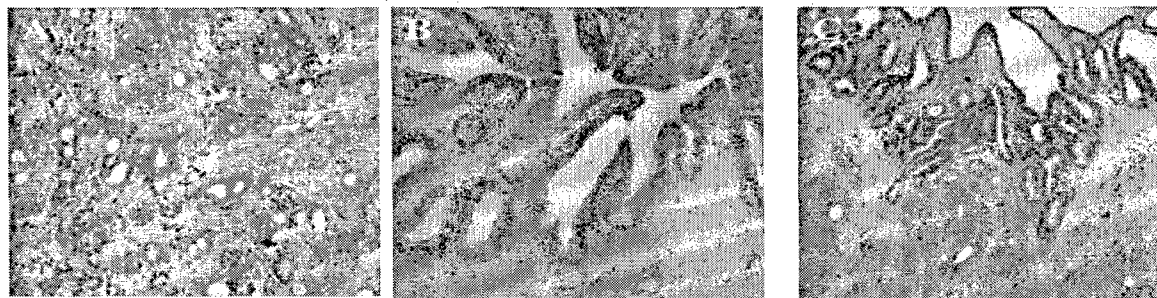


**Fig. 4. Inhibition of AR-mediated transactivation in LNCaP cells upon paxillin ablation.** LNCaP cells were transiently transfected with a MMTV promoter-linked luciferase reporter plasmid and an AR expression plasmid and either a non-specific (ns) siRNA or siRNA directed against endogenous human paxillin. Cells were incubated in the presence or absence of 100 nM DHT for 24 hours. Histogram in A shows the mean of DHT induction of relative luciferase activity normalized to total protein levels in each sample. Note the specific inhibition of DHT induction of transcription in cells treated with paxillin siRNA. Western blot in B shows levels of endogenous paxillin and GAPDH that results in non-hormone (lanes 1, 3) or DHT-treated (lanes 2, 4) cells following treatment with control, nonspecific siRNA (lanes 1,2) or paxillin siRNA (lanes 3,4).

Since paxillin and Hic-5/ARA55 both function as AR coactivators in PC cell lines in culture, we have initiated experiments to examine whether these two closely related coactivators have distinguishable activities *in vivo*. Interestingly, paxillin and Hic-5/ARA55 expression in human prostate tissue is not coincident. Specifically, Hic-5/ARA55 is predominantly expressed in prostate stromal cells both in normal and cancerous tissue (Figure 5) while paxillin is found mainly in the epithelial cells (Figure 6). These results suggest that these two closely related AR coactivators could exert compartment-specific effects to differentially regulate AR action in intact tissue.



**Figure 5. Hic-5/ARA55 expression in human prostate tissue.** Immunohistochemical analysis of Hic-5/ARA55 expression in prostate tissue from normal human donor (A,B) or PC patient (C,D). Control non-immune serum was used in A,C while a mouse anti-Hic-5/ARA55 antibody was used in B,D. Note the restriction of Hic-5/ARA55 staining to stromal compartments in both normal and tumor tissue.



**Figure 6. Paxillin expression in human prostate tissue.** Immunohistochemical analysis of paxillin expression in prostate tissue from normal human donor (B) or PC patient (A,C). Control non-immune serum was used in A, while a mouse anti-paxillin antibody was used in B,C. Note the epithelial cell staining of paxillin in both normal and tumor tissue.

## Key Research Accomplishments

### Task 1:

1. Identification of paxillin as a coactivator for androgen receptor in prostate cancer cell lines
2. Identification of paxillin domain required for androgen receptor coactivation

### Task 2:

1. Demonstration that the interaction between androgen receptor and paxillin in prostate cancer cell lines is hormone-dependent
2. Demonstration of transient interaction between paxillin and AR in membrane preparations following androgen treatment of prostate cancer cell lines
3. Demonstration of synergistic effects on AR transactivation with overexpression of paxillin and a member of the p160 family of coactivators (i.e. GRIP1) in prostate cancer cell lines.
4. Demonstration of role of endogenous paxillin in androgen regulated transcription in prostate cancer cell lines.

5. Demonstration of distinct compartment-specific expression of paxillin and its closely related family member Hic-5/ARA55 in human prostate tissue.

## Reportable Outcomes

### Manuscripts

- Kasai M, Guerrero-Santoro J, Friedman R, Leman ES, Getzenberg RH, and DeFranco DB (2004). The group 3 LIM domain protein paxillin potentiates androgen receptor transactivation in prostate cancer cell lines. *Cancer Research* 63: 4927-4935.
- Guzey, M. & DeFranco D.B. Differential interactions between androgen receptor and paxillin within the nucleus and plasma membrane. *Manuscript in preparation*
- Heitzer M. & DeFranco D.B. Hic-5/ARA55 is a stromal-specific androgen receptor coactivator in prostate and participates in androgen regulation of KGF-1 gene expression in a prostate stromal cell line. *Manuscript in preparation*

## Conclusions

Our results establish that paxillin as well as its closely related LIM domain family member, Hic-5/ARA55 function within the nucleus to potentiate the transactivation activity of AR. For paxillin, distinct effects on AR action may be initiated early through direct interactions with paxillin and the receptor at the plasma membrane and culminate in synergistic actions at target sites within the nucleus with other coactivators. When endogenous paxillin levels are ablated in PC cell lines, AR regulated gene expression is reduced suggesting that paxillin is required for effective androgen action on target genes. Paxillin and its closely related family member Hic-5/ARA55 may exert distinct effects on AR action in intact prostate tissue as they are predominantly expressed in distinct tissue compartments. Hic-5/ARA55 is mainly localized to prostate stromal cells and may influence androgen-regulated expression of growth factors (e.g. KGF-1) that stimulate prostate epithelial cell proliferation. In contrast, paxillin may exert a more direct effect on AR action in prostate epithelium in both normal and cancerous tissues acting at both the plasma membrane and the nucleus with associated AR.

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# The Group 3 LIM Domain Protein Paxillin Potentiates Androgen Receptor Transactivation in Prostate Cancer Cell Lines<sup>1</sup>

Masaaki Kasai, Jennifer Guerrero-Santoro, Robert Friedman, Eddy S. Leman, Robert H. Getzenberg, and Donald B. DeFranco<sup>2</sup>

Departments of Pharmacology [M. K., J. G.-S., R. F., D. B. D.] and Urology [E. S. L., R. H. G.], University of Pittsburgh Cancer Institute [M. K., E. S. L., R. H. G., D. B. D.], University of Pittsburgh School of Medicine, Pittsburgh, Pennsylvania 15261

## ABSTRACT

Paxillin, a member of the group 3 subfamily of LIM domain proteins, is localized within focal adhesions and participates in a number of signal transduction pathways mobilized upon activation of cell surface receptors. In recent years, a number of group 3 LIM domain proteins have been found to also localize within the nucleus and exert direct effects on transcription. We show here that paxillin is present within nuclei and can target the nuclear matrix of CV-1 cells, cultured prostate cancer cell lines, and human prostate tissue. The increased targeting of androgen receptor to the nuclear matrix upon overexpression of paxillin may be brought about by direct interactions between paxillin and the receptor, which were detected *in vitro*. Paxillin functions as a coactivator for androgen receptor and glucocorticoid receptor, but not estrogen receptor  $\alpha$ , similar to its close relative Hic-5/ARA55. Both paxillin and Hic-5/ARA55 use their COOH-terminal LIM domain to interact with steroid receptors. However, paxillin is distinguished from Hic-5/ARA55 by both the location of its receptor coactivation domain (*i.e.*, COOH-terminal LIM domain) and by the dominant-negative activity of its NH<sub>2</sub>-terminal domain. Thus, highly related group 3 LIM domain proteins may use distinct mechanisms to modulate steroid hormone receptor transactivation.

## INTRODUCTION

Androgens are required for the growth and development of the normal prostate gland (1) but also participate in the progression of prostate cancer, particularly in early stages (2). The principal mediator of androgen action within cells is the AR,<sup>3</sup> a member of a large superfamily of nuclear receptors (3, 4). Physiological effects of androgens result primarily from transcriptional regulation of specific target genes brought about by AR interaction with specific DNA sequences or receptor associations with other transcription factors (5–7).

The NH<sub>2</sub>- and COOH-terminal regions of nuclear receptors, including ARs (8–11), possess transactivation domains that function to either enhance the association of basal transcription factors with the preinitiation complex (12) or recruit various coactivator or coactivator complexes (13). Many coactivators exert their actions through the modification of histones (14, 15) or transcription factors (16–18). Families of nuclear receptor coactivator proteins have been identified that either directly or indirectly affect protein acetylation (13) or methylation (19, 20). Although the modulation of chromatin structure

at hormonally responsive promoters is a critical feature of nuclear receptor coactivators, it seems likely that distinct coactivators could use different mechanisms to potentiate nuclear receptor transactivation (10, 21, 22).

The AR has proven to be a particularly useful reagent for the identification of novel transcriptional coactivators (23–26). For example, Hic-5/ARA55 was first identified as a coactivator for AR (27) and later shown to potentiate the transactivation activities of a subset of steroid hormone receptors (28) including the GR. Although Hic-5/ARA55 was shown to associate with the nuclear matrix (28), most studies of this protein have focused on its focal adhesion binding (29, 30). Hic-5/ARA55 is a member of the paxillin family of focal adhesion proteins (31, 32) that share a closely related COOH-terminal LIM domain region (33). The LIM domain is a unique double zinc finger motif (33) with a conserved CXXCX-16–23 amino acid-HXXHXX-HXXHX-16–21 amino acid-CXX(D/H/C) sequence and is found in a variety of proteins of diverse functions and localization (34). Paxillin and Hic-5/ARA55 belong to a group 3 subfamily of LIM domain proteins and contain either three or four separate LIM domains in their COOH-terminal region (35).

In addition to their localization within focal adhesions, members of the group 3 LIM domain family are also associated with the nucleus (30, 36). For zyxin and the closely related LPP protein, nuclear export signal sequences have been identified that regulate their shuttling between the nucleus and focal adhesions (36, 37). In addition, transactivation domains have been identified within LPP and another zyxin family member, Trip6 (38, 39). Trip6 was isolated based upon its hormone-dependent interaction with thyroid hormone receptor (40). Finally, although the transcriptional regulatory properties of the zyxin family member Ajuba have not been revealed, nuclear localization of Ajuba LIM domains is required for PC19 embryonal carcinoma cells to undergo differentiation *in vitro* into an endodermal cell type (41). Despite the prevalence of nuclear group 3 LIM domain proteins, the physiologically relevant transcriptional targets regulated by these proteins remain largely undefined.

In this report, we examine the nuclear localization and steroid receptor coactivation activity of paxillin. Paxillin, similar to Hic-5/ARA55, is a selective steroid receptor coactivator that has the capacity to localize to the nuclear matrix in prostate cancer cell lines and prostate tissue. However, paxillin is distinguished from Hic-5/ARA55 in the domain requirements of its nuclear functions. Thus, highly related group 3 LIM domain proteins may use distinct mechanisms to modulate steroid hormone receptor transactivation. Given the alterations in focal adhesion signaling and paxillin activity in prostate cancer cells (42, 43), the ability of paxillin to function as an AR coactivator in prostate cancer cells could have important implications for the progression of AR-dependent tumors.

## MATERIALS AND METHODS

**Plasmids.** Mammalian expression vectors for FL paxillin, its NH<sub>2</sub> terminus, and COOH terminus with an NH<sub>2</sub>-terminal HA epitope tag were constructed by inserting PCR-amplified cDNA fragments into the pSG5.HA expression plasmid (28). The NH<sub>2</sub>-terminal paxillin expression plasmid in-

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<sup>2</sup> To whom requests for reprints should be addressed, at Department of Pharmacology, University of Pittsburgh Cancer Institute, University of Pittsburgh School of Medicine, 200 Lothrop Street, Pittsburgh, PA 15261.

<sup>3</sup> The abbreviations used are: AR, androgen receptor; GR, glucocorticoid receptor; ER, estrogen receptor; LPP, LIM-containing lipoma-preferred partner; HA, hemagglutinin; MMTV, mouse mammary tumor virus; CMV, cytomegalovirus; FL, full length; FBS, fetal bovine serum; DHT, dihydrotestosterone; Dex, dexamethasone; PMSF, phenylmethylsulfonyl fluoride; GST, glutathione *S*-transferase; IIF, indirect immunofluorescence; NMST, nuclear matrix-targeting signal; FAK, focal adhesion kinase; LIMO, LIM only; CREB, cyclic AMP-responsive element binding protein; CREM, cyclic AMP-responsive element modulator; CBP, CREB binding protein.

cludes human paxillin amino acids 1–307, whereas the COOH-terminal expression plasmid includes amino acids 308–557. Mammalian expression vectors for FL Hic-5/ARA55 and GRIP1 containing an HA epitope tag were provided by Dr. M. R. Stallcup (University of Southern California School of Medicine, Los Angeles, CA). pLC-LUC contains a luciferase reporter gene driven by the MMTV long terminal repeat with a bifunctional androgen response element and glucocorticoid response element (28). ERE-LUC contains an estrogen response element-driven luciferase reporter gene and was provided by Dr. M. Nichols (University of Pittsburgh School of Medicine). hCMV5-AR contains human FL human AR cDNA under the control of CMV promoter and was provided by Dr. D. Robins (University of Michigan Medical School, Ann Arbor, MI). 6R-GR contains cDNA encoding FL rat GR under the control of Rous sarcoma virus long terminal repeat promoter and was provided by Dr. K. Yamamoto (University of California School of Medicine, San Francisco, CA). CMV-ER containing human ER cDNA under the control of CMV promoter was provided by Dr. M. Nichols (University of Pittsburgh School of Medicine, Pittsburgh, PA). The vectors for FL paxillin, pGEX-4T FL paxillin, pGEX-4T NH<sub>2</sub>-terminal paxillin, and pGEX-4T COOH-terminal paxillin were provided by Dr. S. Thomas (Harvard Medical School, Boston, MA).

**Transient Transfection.** Cells ( $1 \times 10^5$ ) were passed onto 30-mm cell culture dishes 24 h before transfection. Transient transfections were performed using the Lipofectamine Reagent (Life Technologies, Inc., Gaithersburg, MD). Briefly, cells were incubated in transfection medium for 5 h. The transfection medium was replaced with 5% charcoal-stripped FBS in the absence or presence of 100 nM of DHT, 1  $\mu$ M Dex, or 10 nM 11 $\beta$ -estradiol. Cells were harvested after an additional 24-h incubation at 37°C in humidified air containing 5% CO<sub>2</sub>, and then extracts were prepared for luciferase assays (28) using procedures specified by the manufacturer of the Luciferase Assay kit (Promega Corp., Madison, WI).

**Nuclear Matrix Preparation.** Subcellular fractionations were performed using procedures described previously (44). Briefly, harvested cells were treated with ice-cold CK buffer (10 mM PIPES, pH 6.8, 100 mM NaCl, 300 mM sucrose, 3 mM MgCl<sub>2</sub>, 1 mM EGTA, 0.5% and Triton X-100) containing 5  $\mu$ g/ml aprotinin, leupeptin, and pepstatin, 1.2 mM PMSF, and 4 mM vanadyl riboside complex. After centrifugation, supernatants were saved (*i.e.*, CSK-extract), and pellets were treated with DNase I digestion buffer (10 mM PIPES, pH 6.8, 50 mM NaCl, 300 mM sucrose, 3 mM MgCl<sub>2</sub>, 1 mM EGTA, and 0.5% Triton X-100) containing 300 units/ml DNase I, 5  $\mu$ g/ml aprotinin, leupeptin, and pepstatin, 1.2 mM PMSF, and 4 mM vanadyl riboside complex. Resultant pellets were further extracted with 0.25 M ammonium sulfate, and the final pellet after centrifugation was designated as the nuclear matrix fraction. The CK buffer extracts, DNase I digestion extracts, and nuclear matrix proteins were subjected to Western blot analysis using a mouse anti-paxillin antibody (Transduction Laboratories, Lexington, KY), goat anti-lamin B antibody (Santa Cruz Biotechnology, Inc., Santa Cruz, CA), or mouse anti-actin antibody (Santa Cruz Biotechnology, Inc.). For *in situ* extractions, analogous extractions were performed with cells that were grown on coverslips. An analogous method (45) was used to prepare nuclear matrix fractions from human prostate tissue obtained from radical prostatectomy samples and prostates obtained

from organ donors. All tissues were collected after appropriate consent under an Institutional Review Board-approved protocol.

**GST Pull-Down Assay.** CV-1 cells were transfected with the pCMV5-AR plasmid and incubated in DMEM containing 10% FBS for 48 h after transfection. CV-1 cells and GRH2 cells were serum starved overnight, followed by an additional 2-h incubation with 100 nM DHT or 1  $\mu$ M Dex, respectively. Whole-cell lysates were then prepared with lysis buffer (10 mM HEPES, pH 7.0, 450 mM NaCl, 5 mM EDTA, 0.05% SDS, 1% Triton X-100 for CV-1 cells; 50 mM Tris-HCl, pH 7.5, 450 mM NaCl, 2.5% sodium deoxycholate, and 1% NP-40 for GRH2 cells) containing 10  $\mu$ g/ml aprotinin, leupeptin, and pepstatin, 1 mM PMSF and sodium orthovanadate, and precleaned with glutathione-Sepharose. Twenty-five  $\mu$ l of glutathione-Sepharose was incubated with 125  $\mu$ g of GST-FL paxillin, GST-NH<sub>2</sub>-terminal paxillin, GST-COOH-terminal paxillin, or GST protein for 1 h at 4°C. Glutathione-Sepharose-bound GST proteins were washed three times with PBS and then added to the CV-1 or GRH2 cell lysates containing 500  $\mu$ g of total protein. Mixtures were incubated for 3 h at 4°C and then washed three times with washing buffer. Proteins bound to GST-protein derived from AR-transfected CV-1 cells or GRH2 cells were subjected to Western blot analysis using the AR (46) anti-AR antibody (Santa Cruz Biotechnology, Inc.) or BuGR2 anti-GR antibody (Affinity Bioreagents, Golden, CO), respectively.

**IIF.** IIF assays were carried out as described previously (28). Briefly, CV-1 cells were fixed with 4% paraformaldehyde and then permeabilized with 0.1% Triton X-100. An anti-paxillin mouse monoclonal antibody (Transduction Laboratories) or an anti-Hic-5/ARA55 rabbit polyclonal antibody (kindly provided by Dr. K. Tachibana, Harvard University, Cambridge, MA) was used to detect endogenous paxillin or Hic-5/ARA55, respectively, for *in situ* assays. A FITC-conjugated goat-anti-mouse IgG F(ab')<sub>2</sub> antibody (Chemicon International, Temecula, CA) or a FITC-conjugated goat-anti-rabbit IgG (H+L) antibody (Jackson ImmunoResearch Laboratories, West Grove, PA) was used as a secondary antibody.

## RESULTS

**Localization of Paxillin within the Nuclear Matrix.** Given the increasing evidence for nuclear localization of group 3 LIM domain proteins (28, 36, 37, 39), we evaluated whether paxillin was likewise present within the nucleus. Biochemical extractions performed either *in situ* or with harvested cells were used to provide a more thorough assessment of subnuclear compartmentalization of paxillin. As shown in Fig. 1, very little paxillin could be detected within the nucleus of CV-1 cells using IIF analysis with paraformaldehyde-fixed and Triton X-100 permeabilized cells. However, paxillin nuclear staining was revealed when cells were extracted with a low-salt detergent buffer (Fig. 1B, *CSK-extracted*). Furthermore, when CV-1 cells were subjected to more extensive biochemical extractions *in situ*, robust staining of paxillin was detected in the nuclear matrix fraction (Fig. 1C).

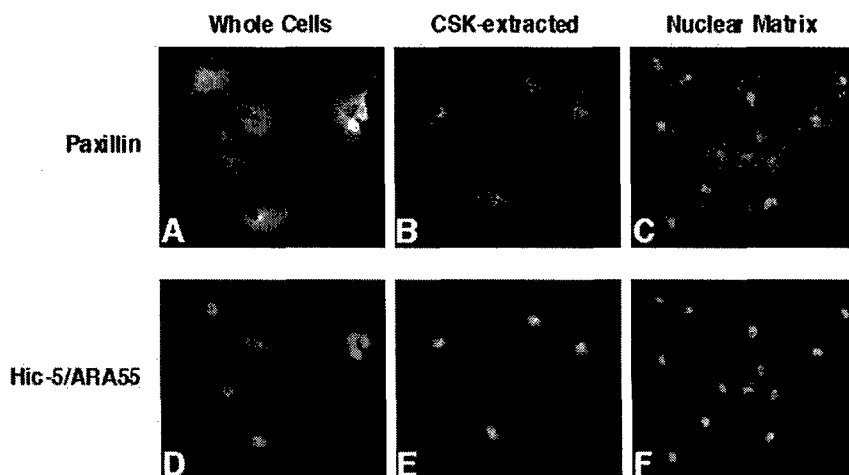


Fig. 1. IIF analysis of endogenous paxillin and Hic-5/ARA55 localization in CV-1 cells. CV-1 cells were fixed with paraformaldehyde and processed for IIF analysis using either an anti-paxillin antibody (A–C) or anti-Hic-5/ARA55 antibody (D–F). Before fixation, cells were either permeabilized with low-salt detergent buffer (*CSK-extracted*; B and E) or further digested with DNase I and then extracted with ammonium sulfate to generate a nuclear matrix fraction (C and F).

The staining pattern of Hic-5/ARA55 in CV-1 was analogous to that observed for paxillin, although nuclear localization of Hic-5/ARA55 was more obvious in whole cells (Fig. 1D).

To confirm the IIF results of paxillin localization and broaden the examination of its subnuclear targeting, CV-1 cells and the PC-3 and LNCaP prostate cancer cell lines were harvested and subjected to differential biochemical extractions that ultimately resulted in the isolation of a nuclear matrix fraction. As shown in Fig. 2A, most endogenous paxillin in all three cell lines examined was recovered from cells by the CSK buffer extraction. Very little paxillin that resists this extraction remains associated with chromatin because DNase I digestion did not lead to recovery of additional paxillin (Fig. 2A). However, paxillin is detected in the nuclear matrix fraction that remains after extensive biochemical extraction in the three cell lines (Fig. 2A). The amount of paxillin recovered in the nuclear matrix fraction is estimated from densitometric analysis of these blots to be ~10% of whole-cell paxillin. This agrees well with previous estimates of the fraction of endogenous Hic-5/ARA55 that targets the matrix of fibroblast cells (28). The integrity of the various biochemical fractions was confirmed upon costaining of blots with either lamin B, to detect nuclear matrix, or actin, to detect CSK buffer-extractable protein (Fig. 2A). Because nuclear matrix binding of ARs is well established (47,

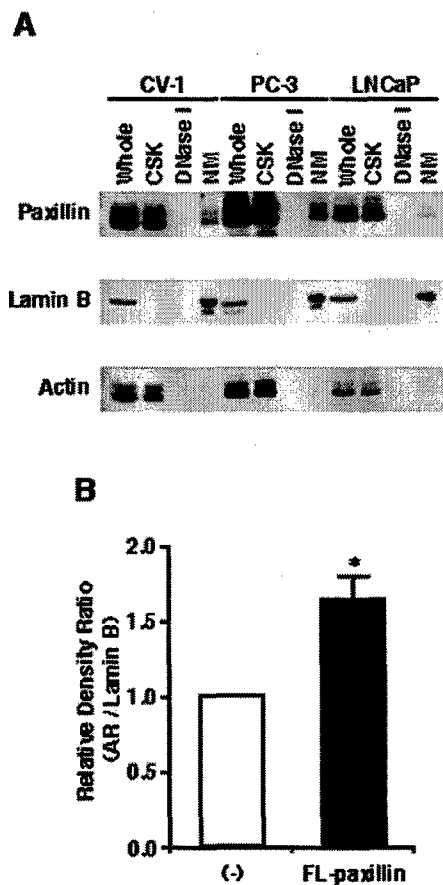


Fig. 2. Subcellular localization of paxillin and effects on AR nuclear matrix binding. A, proteins present in whole-cell lysates (Whole), CSK buffer-extracted supernatants (CSK), DNase I-digested supernatants (DNase I), or nuclear matrix pellets (NM) from CV-1, PC-3, and LNCaP cells were separated by SDS-PAGE and subjected to Western blot analysis using an anti-paxillin antibody. Blots were also stripped and reprobed with an anti-lamin B or anti-actin antibody. B, CV-1 cells transiently transfected with an AR and paxillin expression vector were incubated with 100 nM DHT for 2 h. Nuclear matrix fractions from transfected cells were subjected to Western blot analysis using an anti-AR and anti-lamin B antibody. AR in the nuclear matrix fraction was normalized using lamin B. \*,  $P < 0.05$ , significantly different from the mean value of paxillin-nontransfected control; bars, SE.

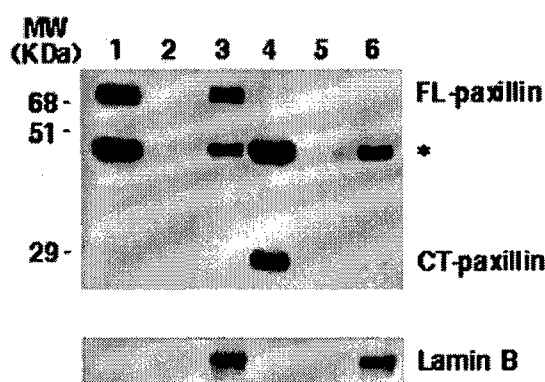


Fig. 3. Subcellular localization of FL COOH-terminal truncated paxillin in CV-1 cells. CV-1 cells were transiently transfected with expression plasmids for either FL HA-tagged paxillin (Lanes 1–3) or a deletion derivative of paxillin containing its COOH-terminal (CT) LIM domains (Lanes 4–6) and subjected to subcellular fractionation. The CSK buffer-extracted fraction (Lanes 1 and 4), DNase I-digested fraction (Lanes 2 and 5), and the nuclear matrix fraction (Lanes 3 and 6) prepared from transfected cells were subjected to Western blot analysis with anti-HA antibody and reprobed with anti-lamin B antibody. \*, nonspecific band that appeared upon anti-HA antibody staining of blot.

48), we examined whether paxillin could function as a nuclear matrix acceptor (47) for AR. As shown in Fig. 2B, overexpression of paxillin in CV-1 cells leads to an increased association of AR with the nuclear matrix. Although paxillin may not be the sole nuclear matrix acceptor protein for AR, these results suggest that it may play a role in the recruitment of AR to the nuclear matrix.

The NMTS of Hic-5/ARA55 is contained within its COOH-terminal LIM domain region (28) and appears to be restricted primarily to its LIM3 domain.<sup>4</sup> The NH<sub>2</sub>-terminal regions of paxillin and Hic-5/ARA55, devoid of LIM domains, are localized exclusively within the cytoplasm when expressed as green fluorescent protein chimera (30). Given the high degree of relatedness of Hic-5/ARA55 and paxillin LIM domains (30), we expected that the paxillin NMTS would likewise be located within its COOH-terminal LIM domain region. However, unlike the COOH-terminal LIM domain region of Hic-5/ARA55 expressed in CV-1 cells, which is indistinguishable from FL Hic-5/ARA55 in its nuclear matrix targeting (28), the COOH-terminal LIM domain region of paxillin is only minimally associated with the nuclear matrix (Fig. 3). Thus, although the NH<sub>2</sub>-terminal region of paxillin does not appear to target on its own to the nucleus, it must contain sequences that cooperate with the LIM domain region to generate a fully functional NMTS.

In addition to examining subnuclear compartmentalization of paxillin in prostate cancer cell lines, we performed an analysis of paxillin localization in prostate tissue from normal, prostate cancer, and benign prostate hyperplasia tissue. As shown in Fig. 4A, paxillin could be detected in the nuclear matrix fraction prepared from prostate, irrespective of the disease state of the tissue. Although our sample size is too limited to make any firm conclusions regarding the relative amount of paxillin in the nuclear matrix of the tissue samples, in just about every case the relative amount of paxillin in the nuclear matrix fractions, expressed as a ratio of paxillin:lamin B, exceeds that detected in LNCaP cells (Fig. 4B). It is unclear whether apparent differences in the range of paxillin nuclear matrix binding between the various tissue samples are significant. Although the analysis of additional prostate tissue samples will be required to reveal whether quantitative differences in nuclear matrix binding exist in different stages of prostate cancer, our studies clearly establish a nuclear matrix association of paxillin in human prostate tissue.

<sup>4</sup> J. Guerrero-Santoro and D. B. DeFranco, unpublished data.

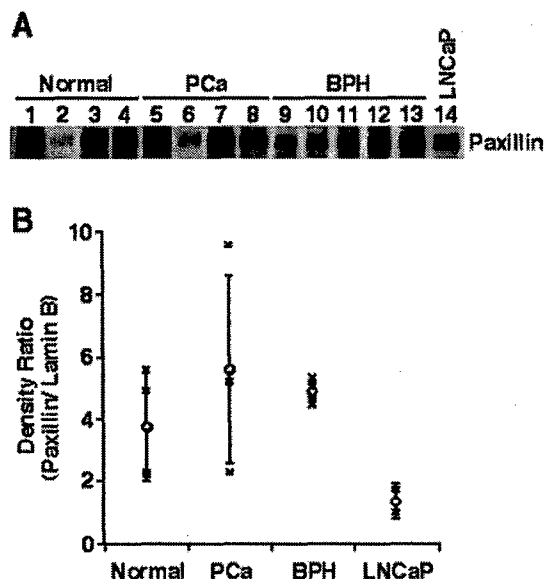


Fig. 4. Presence of paxillin in nuclear matrix fractions of human prostate tissue. Nuclear matrix fractions derived from normal prostates, prostate cancer (PCa), benign prostate hypertrophy (BPH), or LNCaP cells were subjected to Western blot analysis using an anti-paxillin antibody (A) or lamin B antibody (not shown). The ratio of paxillin/lamin B density on individual Western blots is shown (B).

**Paxillin Is a Coactivator for AR.** Although Hic-5/ARA55 and paxillin are closely related in their LIM domain structure (30), they exhibit opposing physiological effects, particularly with regard to their role in cellular proliferation and signal transduction (29, 49, 50). It is assumed that much of this difference relates to their divergent NH<sub>2</sub>-terminal domains (30). Given the ability of paxillin, similar to Hic-5/ARA55 (28), to localize within the nuclear matrix, we examined whether paxillin shared steroid receptor coactivator activity with Hic-5/ARA55. Because paxillin is ubiquitously expressed (51), our studies tested whether overexpression of paxillin was capable of enhancing AR coactivation. As shown in Fig. 5A, the transactivation activity of AR on the androgen-responsive MMTV promoter was potentiated in transiently transfected CV-1 cells upon cotransfection with a paxillin expression vector. A maximum 5-fold effect of paxillin on AR transactivation was observed when 1  $\mu$ g of paxillin expression plasmid was cotransfected with AR and the MMTV reporter (Fig. 5A). Paxillin did not exert any effect on the basal activity of the MMTV promoter. It was not possible to assess whether paxillin effects could be saturated because inclusion of additional DNA in the transfections resulted in reduced cell viability (data not shown). The coactivator activity of paxillin was also exerted on transiently transfected GR (Fig. 5B) but not on ER $\alpha$  (Fig. 5C). This selectivity of paxillin effects is analogous to that observed with Hic-5/ARA55, which was found to act as a coactivator for GR and AR but not ER $\alpha$  (28). As shown in Fig. 6, paxillin is also a coactivator for AR and GR in transiently transfected PC-3 prostate cancer cells.

**Functional Interactions of Paxillin with Other AR Coactivators.** To reveal the relationship between paxillin and other nuclear receptor coactivators, we examined the effects on AR transactivation in cells cotransfected with paxillin and Hic-5/ARA55 or GRIP-1. GRIP-1 is a member of the p160 family of nuclear receptor coactivators (52) and is unrelated in structure and function to Hic-5/ARA55 and paxillin. As shown in Fig. 7, when expressed together in either CV-1 or PC-3 cells, paxillin and Hic-5/ARA55 generally exert additive effects on AR and GR transactivation. In CV-1 cells, paxillin and Hic-5/ARA55 effects on AR transactivation are slightly synergistic (Fig. 7, A and B). Analogous to results obtained with Hic-5/ARA55 (28), cotransfection

of paxillin with GRIP-1 also generated a synergistic effect on AR and GR transactivation in both AR and GR (Fig. 8).

**Domains of Paxillin That Contribute to AR Coactivation Activity.** The group 3 LIM domain proteins share extensive homology within their COOH-terminal LIM domain region but possess divergent NH<sub>2</sub>-terminal domains (34). To identify the domain responsible for steroid receptor coactivator activity, paxillin NH<sub>2</sub>- and COOH-terminal domains were transfected with AR into CV-1 cells. Western blot analysis showed that both domains of paxillin were equivalently expressed in transfected cells (data not shown). As shown in Fig. 9, A and B, the COOH-terminal LIM domain of paxillin was as effective as FL paxillin in potentiating the transactivation activity of AR and GR

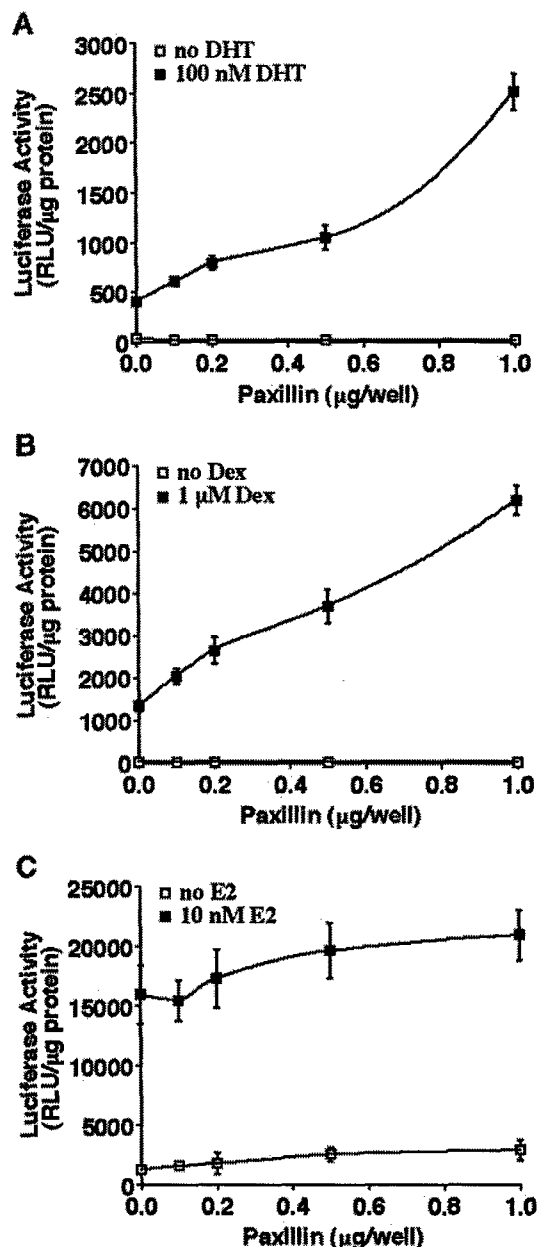


Fig. 5. Paxillin enhances AR and GR but not ER transactivation in CV-1 cells. CV-1 cells were transiently transfected with various amounts of a paxillin expression vector along with the appropriate luciferase reporter plasmid and AR (A), GR (B), or ER (C) expression vectors. After replacement of transfection medium with DMEM containing 5% charcoal-stripped FBS, the cells were incubated in the presence or absence of 10<sup>-7</sup> M DHT (A), 10<sup>-6</sup> M Dex (B), or 10<sup>-8</sup> M 17 $\beta$ -estradiol (E2; C) for 24 h. Each point is the mean (bars, SD; n = 4) normalized to total protein levels in each sample.

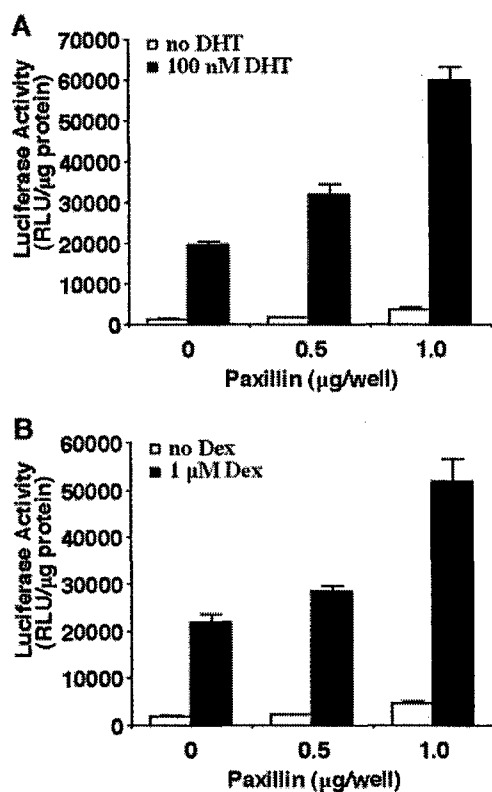


Fig. 6. Paxillin enhances AR and GR transactivation in the PC-3 prostate cancer cell line. PC-3 cells were transiently transfected with indicated amounts of a paxillin expression vector along with a MMTV promoter-linked luciferase reporter plasmid and an AR (A) expression vector or empty expression plasmid (B). A GR-expressing vector was not necessary for the GR transactivation assay because PC-3 cells express functional GR. After replacement of transfection medium with DMEM containing 5% charcoal-stripped FBS, the cells were incubated in the presence or absence of  $10^{-7}$  M DHT (A) or  $10^{-6}$  M Dex (B) for 24 h. The histograms show the means (bars, SD;  $n = 4$ ) of relative luciferase activity normalized to total protein levels in each sample.

in CV-1 cells. Cotransfection of equal amounts of FL and COOH-terminal paxillin generated additive effects on AR and GR transactivation (Fig. 9, A and B). Basal activity of the MMTV promoter (Fig. 9) was not affected by COOH-terminal paxillin. In analogous trans-

fection experiments, the COOH-terminal region of Hic-5/ARA55 was not found to exert any effects on GR transactivation (28).

In contrast to the effect of the COOH-terminal region of paxillin, overexpression of the NH<sub>2</sub>-terminal region of paxillin did not potentiate AR transactivation in CV-1 cells. However, the NH<sub>2</sub>-terminal domain of paxillin was not neutral because its overexpression interfered with the coactivator activity of transfected FL paxillin on AR and GR (Fig. 9, C and D). This "dominant negative" effect of the NH<sub>2</sub>-terminal region of paxillin did not result from reduced expression of transfected FL paxillin (data not shown). Because AR and GR transactivation activity was not reduced by NH<sub>2</sub>-terminal paxillin beyond that observed in the absence of overexpressed paxillin (Fig. 9, C and D), this domain of paxillin may not affect the functioning of other coactivators, at least at the levels expressed under these transfection conditions.

**Interaction between Paxillin and AR.** To assess whether paxillin effects on AR and GR are direct, we used a GST pull-down assay to examine paxillin interactions with these receptors. As shown in Fig. 10A, a FL paxillin-GST fusion protein bound to AR or GR expressed in transfected CV-1 cells or the GrH2 rat hepatoma cell line, respectively. Control pull-down assays with GST alone did not result in the recovery of GR or AR, whereas the FAK protein was recovered with GST-paxillin pull-downs, as expected (Fig. 10, A and B). Importantly, the *in vitro* interaction between the steroid receptors (*i.e.*, AR and GR) and paxillin was equally efficient when receptors were unliganded or hormone bound (Fig. 10, A and B). As shown in Fig. 10, C and D, the interaction between the steroid receptors (*i.e.*, AR and GR) and paxillin occurred through its COOH-terminal LIM domain. This result is consistent with previous domain mapping of Hic-5/ARA55, which delineated its COOH-terminal LIM domain region as a steroid receptor-interaction domain (28).

## DISCUSSION

LIM proteins are classified into three broad groups based largely on their localization, known partners, and apparent biological function (34). Group 1 LIM proteins localize primarily within the nucleus and affect transcription directly through their associated homeodomains or by acting as transcriptional cofactors. LIM proteins in this group are important regulators of development in various tissues (53, 54). The

Fig. 7. Enhancement of hormone-induced AR or GR transactivation by paxillin and Hic-5/ARA55. CV-1 cells (A and B) or PC-3 cells (C and D) were transiently transfected with the indicated amounts of paxillin or Hic-5/ARA55 expression vectors along with a MMTV promoter-linked luciferase reporter plasmid and an AR (A and C) expression plasmid, GR expression plasmid (B), or empty expression plasmid (D). Cells were incubated in the presence or absence of  $10^{-7}$  M DHT (A and C) or  $10^{-6}$  M Dex (B and D) for 24 h. The histograms show the means (bars, SD;  $n = 4$ ) of relative luciferase activity normalized to total protein levels in each sample.

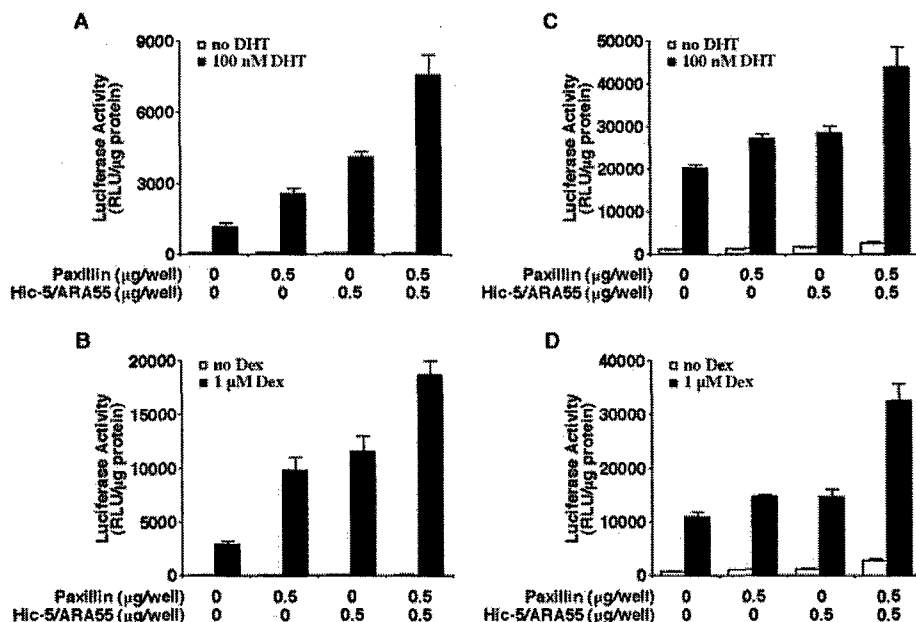
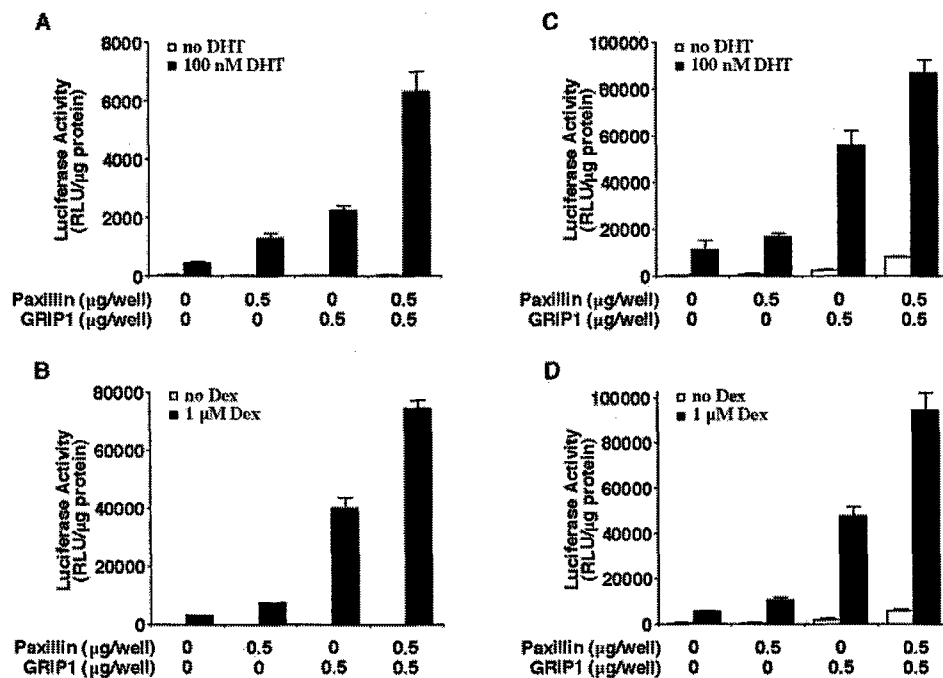


Fig. 8. Enhancement of hormone-induced AR or GR transactivation by paxillin and GRIP1. CV-1 cells (A and B) or PC-3 cells (C and D) were transiently transfected with the indicated amounts of paxillin or GRIP1 expression plasmids along with a MMTV promoter-linked luciferase reporter plasmid and an AR (A and C) or GR expression plasmid (B), or empty expression plasmid (D). Cells were incubated in the presence or absence of  $10^{-7}$  M DHT (A and C) or  $10^{-6}$  M Dex (B and D) for 24 h. The histograms show the means (bars, SD;  $n = 4$ ) of relative luciferase activity normalized to total protein levels in each sample.

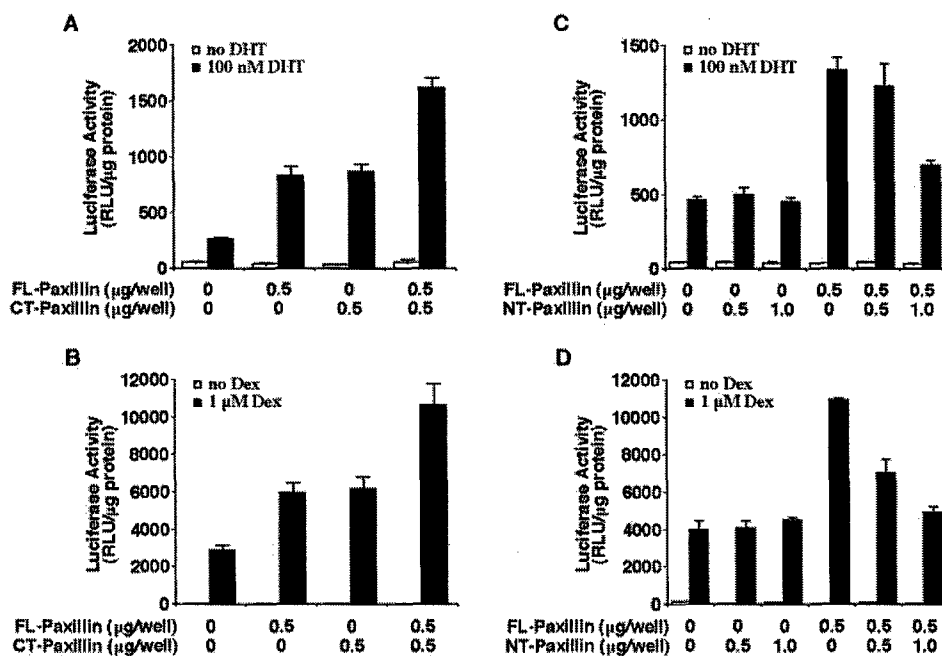


group 3 LIM proteins have been characterized by their association with cytoskeletal elements and predominant cytoplasmic localization (34). However, there is increasing evidence for nuclear localization of these proteins, where they might exert direct or indirect effects on transcription. In this report, we establish that paxillin, the founding member of this family, can be localized within the nucleus and in particular possesses nuclear matrix binding activity. In addition, we also show that paxillin can act as a coactivator for AR and GR, but not ER $\alpha$ . These properties of paxillin are analogous to those revealed previously for the group 3 LIM domain protein, Hic-5/ARA55 (27, 28). Although other members of the group 3 LIM domain family have not been tested for their effects on steroid hormone receptors, Trip6, a member of this family isolated as a thyroid hormone-interacting

protein (40), functions as a relA coactivator (51). Thus, group 3 LIM domain proteins that localize predominantly within the cytoplasm, and particularly at cellular adhesion sites, have the capacity to accumulate within nuclei and exert direct effects on transcription.

Zyxin (36), LPP (37), and Ajuba (41) are members of the group 3 LIM domain protein family, the shuttling of which between sites of cell attachment and the nucleus have been assessed directly. In P19 embryonal carcinoma cells, Ajuba shuttles from sites of cell adhesion to the nucleus in response to retinoic acid-induced differentiation into an endodermal cell type (41). Although paxillin is expressed early in embryogenesis and is essential for early embryonic development (55), its localization in response to differentiation cues *in vitro* and *in vivo* has not been examined. Given the effects of paxillin and Hic-5/

Fig. 9. Effect of paxillin COOH terminus or NH<sub>2</sub> terminus on AR or GR transactivation. CV-1 cells were transiently transfected with the indicated amounts of FL paxillin, NH<sub>2</sub>-terminal (NT) paxillin, or COOH-terminal (CT) paxillin expression plasmids along with a MMTV promoter-linked luciferase reporter plasmid and AR (A and C) or GR (B and D) expression plasmids. Cells were incubated in the presence or absence of  $10^{-7}$  M DHT (A and C) or  $10^{-6}$  M Dex (B and D) for 24 h. The histograms show the means (bars, SD;  $n = 4$ ) of relative luciferase activity normalized to total protein levels in each sample.



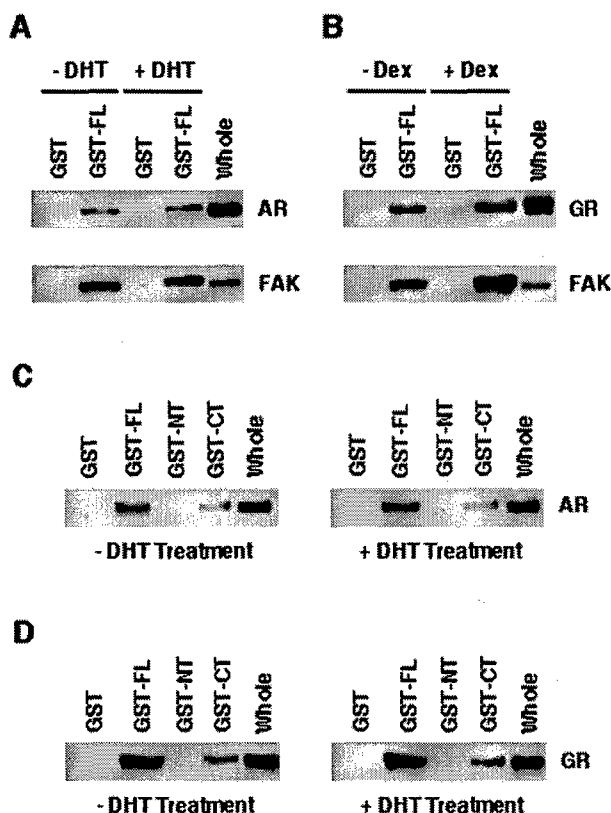


Fig. 10. Hormone-independent interaction of paxillin with steroid receptors *in vitro*. Cell lysates of CV-1 cells transfected with human AR or the GRH2 hepatoma cell line were prepared after either a 2-h hormone treatment or nontreatment and incubated with a glutathione-Sepharose-bound GST-FL-paxillin (A–D), GST-NT-paxillin (C and D), or GST-CT-paxillin (C and D) fusion protein or GST protein for 3 h. Proteins eluted from glutathione-Sepharose were subjected to Western blot analysis with anti-AR antibody (A and C) or anti-GR antibody (B and D), followed by reprobing with anti-FAK antibody (A and B).

ARA55 on steroid receptor transactivation, the subcellular localization of these proteins may be controlled at multiple levels in developing *versus* fully differentiated cells. Although it is unclear whether the subcellular localization of paxillin and/or Hic-5/ARA55 can be regulated by cell adhesion or attachment in steroid hormone-responsive tissue, these dually localized coactivators could provide a direct route for extracellular cues to be transmitted to nuclear hormone receptors.

Paxillin and Hic-5/ARA55 participate in opposing cellular responses. Specifically, paxillin is more often associated with proliferative signals (32), whereas Hic5/ARA55 expression coincides with growth inhibition and cellular senescence (49). Hic-5/ARA55 and paxillin are expressed in a wide variety of vertebrate tissues, yet it is unclear how their effects on similar downstream targets bring about diverse biological responses. Overexpression of Hic-5/ARA55 in 293T cells inhibits integrin-mediated tyrosine phosphorylation of paxillin, an effect most likely mediated by competition for FAK binding (49). However, as shown in our studies, paxillin and Hic-5/ARA55 do not exhibit opposing actions on GR and AR coactivation. Rather, when expressed in limiting amounts Hic-5 and paxillin exert additive effects on steroid receptor coactivation. This observation suggests that Hic-5 and paxillin share some common mechanism in their effects on steroid receptor transactivation.

Although both Hic-5/ARA55 and paxillin use their COOH-terminal LIM domain region for interaction with AR and GR, there appears to be some distinction in their nuclear receptor coactivation. For example, although the COOH-terminal LIM domains of paxillin are as

efficient as FL paxillin in potentiating GR and AR transactivation, this domain of Hic-5/ARA55 does not exhibit coactivator activity on these same receptors in transiently transfected cells (28). The NH<sub>2</sub>-terminal domain of paxillin is likewise distinguished from that of Hic-5/ARA55, given its dominant-negative effect of paxillin coactivation. The LIM domains of paxillin and Hic-5/ARA55 are highly related (30), but it is not unprecedented for highly related LIM domains to differ in their actions on transcription. For example, closely related LIM domains included within different members of the FHL family of LIMO proteins exhibit vastly different transactivation properties when tested within a single cell type (56). Extensive mutagenesis of the ACT protein member of the FHL family revealed that a precise arrangement of specific LIM domains plays a critical role in its ability to act as a transcriptional coactivator for the CREB and CREM proteins (56). This feature of the LIM domain proteins may restrict the utility of domain swaps to delineate the basis for differential effects of paxillin *versus* Hic-5/ARA55 COOH-terminal LIM domains on AR and GR coactivation.

The FHL2 member of the LIMO family has been found to be a very selective AR coactivator and has a restricted tissue distribution (57). These properties distinguish FHL2 from paxillin and Hic-5/ARA55, which do not share AR specificity as coactivators and are expressed in a wide range of tissues and cell types (27, 28, and this report). This highlights the versatility of LIM protein involvement in steroid receptor transactivation.

The interaction that we observed *in vitro* between paxillin is hormone independent. Although this appears to contrast the hormone-dependent *in vitro* interaction between Hic-5/ARA55 and AR (27), we in fact did not observe a hormone dependence of Hic-5/ARA55 interactions *in vitro* with another steroid receptor protein, GR.<sup>5</sup> It is unclear whether the differences in hormone dependence of LIM domain protein interaction with steroid receptors reflect an artifact of extract preparation or is biologically relevant. In this regard, we note that FHL2, another recently identified LIM domain-containing AR coactivator, also did not exhibit hormone dependence in its interaction with AR *in vitro* (57). Analogous to Hic-5/ARA55 and paxillin, FHL2 interactions with AR involve direct contact with its LIM domain (57).

Because we did not observe an increase in basal activity of hormone-responsive promoters in transfected cells with Hic-5/ARA55 or paxillin, functional interactions of these coactivators with steroid receptors *in vivo* are likely to be hormone dependent. Alternatively, unliganded AR and GR may be restricted in their ability to interact with Hic-5/ARA55 and/or paxillin because of associated heat shock proteins (58). This might also explain why AR or GR targeting to focal adhesions has not been detected, although this possibility may need to be reexamined with more sophisticated cell imaging techniques. Finally, the limited amount of Hic-5/ARA55 and paxillin localized within the nucleus could provide another means of restricting interactions of steroid receptors with these coactivators. We did not observe any effects of hormone treatment on nuclear localization or Hic-5/ARA55 or paxillin in transiently transfected cells,<sup>6</sup> but the possibility of hormone- or cell adhesion-dependent changes in Hic-5/ARA55 and paxillin subcellular localization cannot be excluded without more thorough investigation. Overexpression of paxillin did significantly enhance AR nuclear matrix binding in transfected cells, illustrating a potential role for group 3 LIM domain proteins in steroid receptor trafficking within the nucleus.

The mechanism of LIM domain protein coactivation of steroid hormone receptors remains undefined. Within all members of LIM protein groups, the LIM domain is likely to serve as a platform for

<sup>5</sup> J. Guerrero-Santoro and D. B. DeFranco, unpublished data.

<sup>6</sup> J. Guerrero-Santoro, M. Kasai, and D. B. DeFranco, unpublished data.

protein/protein interactions (34). Although LIM proteins that contain a homeodomain, such as Lhx-3/P-LIM, bind directly to specific transcriptional regulatory sites (59), some LIMO proteins, such as LMO2, are recruited to a specific target site via their interaction with DNA-bound transcription factors such as GATA-1 and the E47-TAL1 heterodimer (60). In this context, the LIM domain facilitates assembly of a multiprotein complex that is required for optimal transcription (56). Future studies should reveal whether the group 3 LIM domain proteins Hic-5/ARA55 and paxillin serve an analogous role to stabilize steroid receptor complexes on DNA.

Recruitment of transcriptional cofactors by individual LIM domains has also been reported. For example, transcriptional activation activity of the group 1 LIM protein, Lhx3, is repressed by RLIM, a RING finger protein that interacts with a Lhx3 LIM domain (61). The specific transcriptional repression activity of RLIM is brought about by its recruitment of corepressor complexes (61). RLIM does not interact with group 3 LIM proteins (*i.e.*, paxillin and zyxin) *in vitro* (60). Lhx2, a LIM protein that like Lhx3 is involved in regulation of pituitary-specific gene expression (62), also uses a partner protein to recruit a transcriptional cofactor. In contrast to RLIM, the Lhx2 LIM domain-interacting protein, MRG1, appears to function as a coactivator through its recruitment of p300/CBP (63). However, Hic-5/ARA55 and paxillin do not appear to function in an analogous mechanism to recruit other established coactivators. Although we have only obtained negative results in assays of Hic-5/ARA55 binding to nuclear coactivators of the p300 and p160 families,<sup>5</sup> the synergistic effect of Hic-5/ARA55 and paxillin on GRIP coactivation supports the contention that these proteins use a distinct mechanism for coactivation. Recent independent results also suggest that Hic-5/ARA55 coactivation of AR is CBP independent (10). The LIMO proteins of the FHL family function as CREB/CREM coactivators in a CBP-independent manner (56). The coactivation activity of these LIM domain proteins may be related to the presence within the LIM sequence of autonomous transactivation domains (56). Hic-5/ARA55 also possesses an autonomous transactivation domain, but this is located within its NH<sub>2</sub>-terminal region and not its LIM domains (28). Although paxillin has not been tested for the presence of an autonomous transactivation domain, it differs from Hic-5/ARA55 in the location of its coactivation domain, which is localized within its LIM domain region.

Given the differences in biological functions of paxillin and Hic-5/ARA55, it may be overly simplistic to assume that their actions as coactivators for steroid hormone receptors are completely overlapping. Both proteins are expressed in androgen-responsive tissue (*e.g.*, prostate), but it is well established that they respond in distinct ways to multiple signaling pathways. The divergent nature of their NH<sub>2</sub>-terminal domain may play a large role in differential responsiveness of these coactivators, but the potential impact of minor variations in LIM domain sequences cannot be ignored. Importantly, the alterations in paxillin function associated with the progression of prostate cancer cells to a metastatic state (43) necessitate a more thorough examination of the impact of this coactivator on AR action in prostate cancer cells with alternative cell migration and adhesion properties.

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