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Prolactin-induced mouse mammary carcinomas model estrogen resistant luminal breast cancer.

Lisa M Arendt

*Department of Anatomy & Cellular Biology, Sackler School, Tufts University School of Medicine;
Department of Comparative Biosciences, University of Wisconsin–Madison*

Debra E Rugowski

Department of Comparative Biosciences, University of Wisconsin–Madison

Tara A Grafwallner-Huseth

Department of Comparative Biosciences, University of Wisconsin–Madison

Maria Jose Garcia-Barchino

Genetics Department, Navarra University, Irunlarrea St. 1, Pamplona, 31008, Spain; Department of Cancer Biology, Kimmel Cancer Center, Thomas Jefferson University

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Hallgeir Rui

 *Department of Cancer Biology, Kimmel Cancer Center, Thomas Jefferson University,
Department of the Medical Biology, Oncology, and the Oncology Commons,
hallgeir.rui@jefferson.edu*

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Authors

Lisa M Arendt, Debra E Rugowski, Tara A Grafwallner-Huseth, Maria Jose Garcia-Barchino, Hallgeir Rui, and Linda A Schuler

RESEARCH ARTICLE

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Prolactin-induced mouse mammary carcinomas model estrogen resistant luminal breast cancer

Lisa M Arendt^{1,3}, Debra E Rugowski¹, Tara A Grafwallner-Huseth¹, Maria Jose Garcia-Barchino^{2,4}, Hallgeir Rui², Linda A Schuler^{1*}

Abstract

Introduction: Tumors that express estrogen receptor alpha (ER α +) comprise 75% of breast cancers in women. While treatments directed against this receptor have successfully lowered mortality rates, many primary tumors initially or later exhibit resistance. The paucity of murine models of this “luminal” tumor subtype has hindered studies of factors that promote their pathogenesis and modulate responsiveness to estrogen-directed therapeutics. Since epidemiologic studies closely link prolactin and the development of ER α + tumors in women, we examined characteristics of the aggressive ER α + and ER α - carcinomas which develop in response to mammary prolactin in a murine transgenic model (neu-related lipocalin- prolactin (NRL-PRL)). To evaluate their relationship to clinical tumors, we determined phenotypic relationships among these carcinomas, other murine models of breast cancer, and features of luminal tumors in women.

Methods: We examined a panel of prolactin-induced tumors for characteristics relevant to clinical tumors: histotype, ER α /progesterone receptor (PR) expression and estrogen responsiveness, Activating Protein 1 (AP-1) components, and phosphorylation of signal transducer and activator of transcription 5 (Stat5), extracellular signal regulated kinase (ERK) 1/2 and AKT. We compared levels of transcripts in the ER α -associated “luminal” signature that defines this subtype of tumors in women and transcripts enriched in various mammary epithelial lineages to other well-studied genetically modified murine models of breast cancer. Finally, we used microarray analyses to compare prolactin-induced ER α + and ER α - tumors, and examined responsiveness to estrogen and the anti-estrogen, Faslodex, *in vivo*.

Results: Prolactin-induced carcinomas were markedly diverse with respect to histotype, ER α /PR expression, and activated signaling cascades. They constituted a heterogeneous, but distinct group of murine mammary tumors, with molecular features of the luminal subtype of human breast cancer. In contrast to morphologically normal and hyperplastic structures in NRL-PRL females, carcinomas were insensitive to ER α -mediated signals. These tumors were distinct from mouse mammary tumor virus (MMTV)-neu tumors, and contained elevated transcripts for factors associated with luminal/alveolar expansion and differentiation, suggesting that they arose from physiologic targets of prolactin. These features were shared by ER α + and ER α - tumors, suggesting a common origin, although the former exhibited transcript profiles reflecting greater differentiation.

Conclusions: Our studies demonstrate that prolactin can promote diverse carcinomas in mice, many of which resemble luminal breast cancers, providing a novel experimental model to examine the pathogenesis, progression and treatment responsiveness of this tumor subtype.

* Correspondence: schulerl@svm.vetmed.wisc.edu

¹Department of Comparative Biosciences, University of Wisconsin–Madison, 2015 Linden Dr., Madison, WI 53706, USA

Full list of author information is available at the end of the article

Introduction

The hormone, prolactin (PRL), is critical for the development and functional differentiation of the mammary gland [1]. Although its physiologic importance suggests a role in breast carcinogenesis, such activity has been controversial. However, accumulating evidence from a variety of sources now supports a link between this hormone and breast disease. Large prospective epidemiologic studies have correlated circulating levels of PRL with an increased risk of particularly estrogen receptor positive (ER α +) invasive tumors [2]. Although the relationship between circulating PRL and patient survival has been examined only in smaller studies, higher levels have been associated with tumor aggression, higher risk of metastasis and poor long term survival (reviewed in [2,3]). Moreover, the pituitary is not the only source of PRL to the breast. PRL is also expressed within the mammary gland, particularly in humans, permitting autocrine/paracrine actions [4-6]. Recent studies have begun to link genetic variations in the genes for PRL and the prolactin receptor (PRLR) and breast cancer [7-9]. Furthermore, in addition to PRL itself, human growth hormone is also a potent agonist at the PRLR; thus PRLR-transduced signals also may mediate some signals of this other oncogenic hormone [10,11]. Finally, many primary breast tumors, both ER α + and ER α -, express the PRLR, pointing to its potential utility as a therapeutic target and prognostic indicator [4,12,13].

Although epidemiological data support a role for PRL in the development and progression of breast cancer, relatively little is known about its contributions to this disease. In order to investigate the pathogenic actions of PRL, we have developed a transgenic mouse model (NRL-PRL), which mimics the high mammary PRL synthesis observed in women. In this model, the PRL- and estrogen- insensitive NRL promoter drives expression of the rat PRL transgene in mammary epithelia, exposing the gland to locally elevated PRL [14,15]. Non-parous NRL-PRL females develop mammary pathology that exhibits many features of human disease, including early lesions (hyperplasias and intraepithelial neoplasias, similar to ductal carcinoma in situ in women), and eventually, ER α + and ER α - carcinomas. These tumors are locally aggressive, and metastases to local lymph nodes and lungs are occasionally observed [16]. The development of tumors in NRL-PRL females is not dependent on postpubertal ovarian steroids, but is accelerated by supplemental estrogen [17].

Expression of ER α has emerged as the foremost prognostic and therapeutic indicator in primary clinical breast tumors. However, ER α + cancers are themselves very diverse, and many are not susceptible to treatments directed at this pathway. Indeed, 25% of women treated

with tamoxifen will succumb to breast cancer [18]. Large scale transcript profiling of clinical tumors has begun to reveal the basis of ER α + cancer diversity, as well as that of other tumor subtypes defined by pathologic markers (for review, [19]). ER α + tumors share the "luminal" transcript signature, which includes the ER α itself and several other genes linked to ER α expression, as well as defining features such as luminal cytokeratins [19,20]. Efforts to resolve luminal tumors into subtypes that predict therapeutic sensitivity have demonstrated elevated expression of proliferation-related genes in patients at higher risk for relapse, which are useful clinically [21-23]. However, these studies have not revealed vulnerabilities that can be therapeutically targeted.

Mouse models have proven to be useful in elucidating the origins of tumor subtypes, and have revealed complex etiologies and relationships to mammary epithelial lineages [24,25]. However, the paucity of murine models of ER α + tumors has hindered studies of the factors that give rise to this prevalent tumor subtype and modulate responsiveness to estrogen-directed therapeutics. In light of the epidemiologic evidence linking PRL exposure to the development of ER α + cancers in women, we investigated the phenotype of the mammary carcinomas that develop in NRL-PRL females, with regard to features defining clinical luminal tumors, including expression of the "ER α -associated signature", estrogen sensitivity, and activated signaling pathways. Our studies demonstrate that PRL can promote diverse tumor phenotypes, many of which display molecular features of luminal breast cancers, providing insight into the pathogenesis of this cancer subtype.

Materials and methods

Reagents

5-bromo-2-deoxyuridine (BrdU) was obtained from Sigma Chemical Co. (St. Louis, MO, USA), and 17 β -estradiol (E2) was purchased from Steraloids, Inc. (Newport, RI, USA). The following antibodies were used for immunohistochemical analyses: BrdU (MAS-250) from Accurate Scientific (Westbury, NY, USA), proliferating cell nuclear antigen (PCNA; PC 10) and progesterone receptor (PR; A0098) from DAKO Cytomation (Carpinteria, CA, USA), c-Fos (SC-52) and estrogen receptor alpha (ER α ; SC-542) from Santa Cruz Biotechnology, Inc. (Santa Cruz, CA, USA), pERK 1/2 (Thr202/Tyr204; 9102), pAkt (S473; 3787), and c-Jun (9162) from Cell Signaling Technology (Beverly, MA, USA), pStat5 (AX1) from Advantex BioReagents, LLP (El Paso, TX, USA).

Mouse models

NRL-PRL mice (line 1655-8, TgN(Nrl-Prl)24EPS) were generated and maintained in the FVB/N strain

background as described [14]. Mice were housed and handled in accordance with the Guide for Care and Use of Laboratory Animals in AAALAC-accredited facilities. All procedures were approved by the University of Wisconsin-Madison Animal Care and Use Committee. p53^{+/-} heterozygotes [26] that were backcrossed more than 10 generations into the FVB/N strain [27] were used to generate p53^{-/-} donors. p53^{-/-} mammary epithelial cells (MECs) were transplanted into FVB/N nontransgenic female recipients, and the resulting tumors harvested (manuscript in preparation, O'Leary and Schuler). Mammary tumors from other genetically modified murine models (FVB/N) were generously provided by Drs. Kim and Alexander (University of Wisconsin) (MMTV-neu [28], and Drs. Green and Zi-Yao Liu (NIH-NCI) (C3(1)-SV40 Tag tumors [29].

Histological examination of mammary tissue

For some studies, mice were injected with 200 mg/kg body weight BrdU 1 h prior to sacrifice to label cells undergoing DNA synthesis. For immunohistochemistry (IHC), deparaffinized slides were exposed to 0.5% H₂O₂ in methanol to block endogenous peroxidase activity, boiled for 15 minutes in 0.1 M citrate (pH = 6.0) or 0.1 M Tris (pH = 9.0) buffer for antigen retrieval, then blocked, and incubated with the primary antibody overnight (for individual antibodies, see Table S1 in Additional File 1). Slides were incubated with the secondary antibody (BioGenex, San Ramon, CA, USA), followed by peroxidase streptavidin and 3,3' diaminobenzidine, and counterstained with hematoxylin. Cells were labeled for PCNA using Vector M.O.M. Basic kit (BMK-2202; Vector Laboratories, Burlingame, CA, USA). pStat 5 was detected as described [30]. Serial sections of tumors were examined for the different endpoints. For all IHC, control sections were stained in parallel (negative controls: omission of primary antibody, irrelevant antibody; positive controls, appropriate for the antibody) as described [14]. To determine labeling indices, 1,000 total cells in at least five randomly chosen microscopic fields in divergent regions of each carcinoma were counted by an investigator blind to the experimental status.

Microarray experiments

In a preliminary study to compare the tumors that develop in NRL-PRL to other mouse models of mammary cancer, cDNA prepared from tumors was hybridized to Agilent Mouse Oligo Microarrays (Agilent Technologies Inc., Santa Clara, CA, USA) using total RNA from equal numbers of C57Bl6/J and 129 male and female pups as the reference sample, as previously described [31]. Data were processed and hierarchical clustering performed, as described [31].

To compare NRL-PRL tumors identified as ER α positive or negative by IHC, total RNA was purified using RNeasy Midi Kits (Qiagen Inc., Valencia, CA, USA) according to the manufacturer's instructions. RNA integrity was assessed using the RNA 6000 Nano Assay and Agilent 2100 Bioanalyzer (Agilent Technologies Inc., Hatfield, PA, USA). MECs were isolated as described [32] from 12-week-old nontransgenic nonparous FVB/N females, for common reference RNA. Reference RNA pooled from 15 mice was reverse transcribed and labeled with Cy3. Tumor and reference samples were co-hybridized overnight to Agilent Mouse cDNA Microarrays (G4104A, 8,614 features), washed, and scanned using Agilent's dual-laser Microarray Scanner (G2565BA) with Feature Extraction Software.

Genes with Cy3 and Cy5 intensity values higher than 100 were considered as expressed genes and were normalized with lowess smoother followed by log base 2 transformation. Values that changed at least 1.5-fold in either direction from the gene's median value were included in the class comparison. Genes differentially expressed between ER α positive and negative classes were determined with a significance threshold of univariate tests at $P < 0.01$. Average linkage hierarchical clustering on both the genes and samples was performed using the gene list generated by class comparison. Analyses were performed using BRB ArrayTools (version 3.6.0) developed by Dr. Richard Simon and Amy Peng Lam (Biometric Research Branch, National Cancer Institute). These data have been deposited in the public database, ArrayExpress (E-MEXP-3013).

Real-time PCR

RNA from tumors and MECs was examined using quantitative real time PCR (qRT-PCR). cDNA was synthesized from 1 μ g RNA using random hexamers (Amersham Biosciences, Piscataway, NJ, USA) and MMLV Reverse Transcriptase (Promega, Madison, WI, USA), and qRT-PCR performed as described [17]. Negative (no cDNA) and positive controls were included with each plate. Specific primer sequences are listed in Table S2 in Additional File 1. Results were calculated using the comparative C_T method and normalized to 18S RNA, and data were analyzed for statistical significance as described in the figure legends.

Tumor sensitivity to ovarian steroids

To examine the responsiveness of tumors to circulating ovarian steroids, tumors were allowed to develop until they reached 1.5 cm in diameter. About one-third of the tumor mass was then removed, concomitant with ovx or sham surgery, or ovx and subcutaneous treatment

with 0.1 mg pellets of 17 β -estradiol (E2; Innovative Research of America, Sarasota, FL, USA), which result in plasma levels of 50 to 75 pg/ml [33]. Tumors were measured twice weekly with calipers, and tumor volume was calculated (the largest diameter \times (the smallest diameter)² \times 0.4). Mice were humanely euthanized and tissue collected when the tumors reached 1.5 cm in diameter. Histological comparisons of the tumor biopsies and end stage tumors were carried out as above.

For some experiments, 1 mm³ NRL-PRL tumor fragments were transplanted bilaterally into fourth inguinal mammary glands of 12-week-old nontransgenic syngeneic FVB/N hosts. After the tumors grew to 0.75 cm in diameter, recipient females were treated with ovx, sham surgery, or sham surgery and weekly subcutaneous injections of 5 mg Faslodex (ICI 182,780; AstraZeneca, Wilmington, DE, USA). Tumors were measured and collected as described above.

Statistical analyses

Statistical analyses were performed using Prism v.4.03 (GraphPad Software, Inc., San Diego, CA, USA). Differences were considered significant at $P < 0.05$.

Results and discussion

Locally elevated PRL leads to histologically diverse ER α positive and ER α negative carcinomas

NRL-PRL nonparous females develop diverse invasive mammary carcinomas with a long latency [14]. In order to examine the breadth of tumor phenotypes promoted by this prolonged PRL exposure, we examined a panel of 39 of these tumors (mean latency \pm standard deviation (S.D.), 18.2 \pm 3.6 months). The majority were adenocarcinomas of several histotypes; carcinosarcomas constituted a minor population (Figure 1, Table 1). These carcinomas varied widely in the proportion of cells that expressed detectable ER α (0 to 79%). Nineteen of the 39 tumors examined exhibited greater than 10% ER α + cells, a common clinically used threshold. The majority did not express detectable progesterone receptor (PR); only four tumors demonstrated PR immunostaining in more than 5% of the tumor cells. The morphological heterogeneity suggests cooperation between PRL and other oncogenic factors during the long period of development of these tumors, similar to that which may occur in women.

PRL-induced tumors display inverse Stat5 and AP-1 activation and elevated epithelial pAKT

PRL can activate multiple signaling cascades, which may play distinct roles in its physiologic and pathogenic actions in the mammary gland. In order to determine the relationship of these signals to tumor characteristics, we examined the activation of these pathways in the

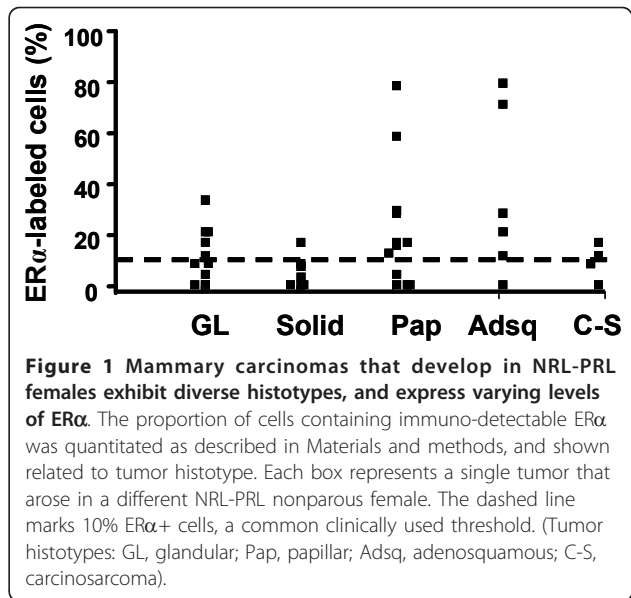


Figure 1 Mammary carcinomas that develop in NRL-PRL females exhibit diverse histotypes, and express varying levels of ER α . The proportion of cells containing immuno-detectable ER α was quantitated as described in Materials and methods, and shown related to tumor histotype. Each box represents a single tumor that arose in a different NRL-PRL nonparous female. The dashed line marks 10% ER α + cells, a common clinically used threshold. (Tumor histotypes: GL, glandular; Pap, papillary; Adsq, adenosquamous; C-S, carcinosarcoma).

panel of PRL-induced tumors by IHC. Signal transducer and activator of transcription 5 (Stat5) is one of the best characterized mediators of PRL activity, and is critical for PRL-induced alveolar development during pregnancy (for review, [1,34]). The tumors that developed in NRL-PRL females contained variable proportions of cells with nuclear pStat5 (range: 4.2 to 93.3%), which was not related to ER α levels (Figure 2a, d, $P > 0.05$). Although many glandular and solid tumors exhibited relatively high proportions of cells containing nuclear pStat5, evidence for activation of this pathway varied substantially even among these histotypes. Many adenocarcinomas demonstrated low nuclear pStat5, as did all of the poorly differentiated carcinosarcomas (Table 1).

PRL can also initiate other signals, including several MAP kinases, which are elevated in mammary lysates of NRL-PRL females [17], and are particularly strongly activated in some breast cancer cells lines [35]. PRL-activated MAP kinases can increase synthesis and phosphorylation of multiple Activating Protein-1 (AP-1) components, activating the AP-1 transcriptional enhancer [36]. AP-1 target genes have been shown to enhance cellular proliferation, survival, and invasion (for review, [37-39]). Both c-Fos and c-Jun were variably expressed in PRL-induced tumors (Figure 2b, c, e). Their expression was directly related to each other (data not shown, $P = 0.005$, $R = 0.83$). Although adenosquamous carcinomas and carcinosarcomas had the highest levels of c-Fos expression, a large subset of other histotypes also displayed significant staining (Table 1). Interestingly, nuclear pStat 5 levels were negatively correlated with both c-Fos (Figure 2e; $P < 0.0001$, $R = -0.63$) and c-Jun (data not shown; $P = 0.0058$, $R = -0.82$). The inverse

Table 1 Characteristics of PRL-induced carcinomas by histotype

N	Glandular 10	Solid 3	Papillary 10	Adenosquamous 3	Carcinosarcoma 4
ERα ¹	10.0 ± 3.6 ^a	4.4 ± 2.9 ^b	21.1 ± 8.1	32.2 ± 10.2 ^{a,b}	8.2 ± 3.0
PR ¹	1.5 ± 1.0	0.3 ± 0.2	13.8 ± 6.3	2.4 ± 1.4	0
pStat5 ¹	49.0 ± 8.13 ^{a,b}	66.6 ± 12.2 ^{c,d,e}	29.4 ± 18.7 ^c	23.9 ± 5.25 ^{a,d}	5.7 ± 1.9 ^{b,e}
c-Fos ¹	18.3 ± 2.8 ^a	15.0 ± 2.5 ^{b,c}	17.6 ± 3.7 ^d	30.5 ± 4.4 ^{a,b}	25.2 ± 3.2 ^{c,d}
S phase ¹	15.6 ± 1.7	15.0 ± 2.5	18.0 ± 2.0	14.7 ± 1.3	15.5 ± 2.2
Apoptotic ¹	2.1 ± 0.3	3.6 ± 1.1	2.2 ± 0.3	1.9 ± 0.4	1.1 ± 0.2
pERK1/2 ²	70% strong, 10% weak stromal; 30% epithelial	100% strong stromal	90% strong stromal; 10% epithelial	100% strong stromal	100% strong stromal & epithelial
pAKT ²	50% epithelial; 20% stromal; 30% negative	67% epithelial; 33% negative	70% epithelial; 10% stromal; 20% negative	33% epithelial & stromal; 67% negative	75% epithelial; 25% negative

¹% tumor epithelial cells that are positive, mean ± s.e.m (see Materials and methods). Same lower case letters indicate significant differences between histotypes ($P < 0.05$; Students *t*-test).

²% of tumors of each histotype that displayed strong staining in different locations (stroma, epithelium, or both) as indicated.

ERα, estrogen receptor alpha; ERK, extracellular signal regulated kinase; PR, progesterone receptor; PRL, prolactin; Stat5, signal transducer and activator of transcription 5.

relationship of these signals in primary NRL-PRL tumors is consistent with our previous observation of PRL signals to Stat5 and AP-1 in several breast cancer cell lines [35].

The higher level of nuclear pStat5 in well-differentiated PRL-induced carcinoma histotypes resembles the association of pStat5 with features of more differentiated clinical breast tumors (for review, [34]). In order to examine the association of this pathway with molecular phenotype in adenocarcinomas, we examined levels of two transcripts associated with distinct mammary behaviors. The 5 PRL-induced adenocarcinomas with highest nuclear pStat5 (70 to 93%) and low c-Fos had significantly elevated transcripts for the proteinase inhibitor, Expi, compared to those adenocarcinomas with lowest pStat5 (4 to 10%, Figure 2f, $P = 0.003$). Expi expression increases in early pregnancy [40], and is reduced in PRLR^{-/-} mammary glands [41]. In contrast, the adenocarcinomas with lowest nuclear pStat5 contained elevated mRNA for the metalloproteinase, *Mmp9* (Figure 2g; $P = 0.03$), which in women is associated with high tumor grade, metastasis, and reduced survival [42], consistent with a more aggressive phenotype.

Elevation of phosphorylated extracellular signal regulated kinases 1/2 (pERK1/2) is found in some clinical tumors in women [43], and is increased in lysates of NRL-PRL mammary glands [17]. The location of pERK1/2 varied substantially among PRL-induced tumors. The majority of the adenocarcinomas (68%) demonstrated strong stromal staining with rare nuclear staining in epithelial cells (Table 1, Figure 3a). In contrast, all of the carcinosarcomas in the panel displayed both strong epithelial nuclear and cytoplasmic labeling (Figure 3b, c). This histotype had significantly higher levels of *Vim* (Figure 3h), but similar levels of *Krt8* transcripts compared to the adenocarcinomas (Figure 3i).

The spindle shaped cell morphology, loss of cell polarity, and expression of *Vim* mRNA within these tumors suggest the epithelial to mesenchymal transition (EMT, for review, [44,45]). Significantly enhanced levels of *Mmp9* transcripts were also detected in these tumors (Figure 3j).

The PI3K/AKT pathway also has been associated with proliferation, survival, and EMT in human breast tumors, as well as resistance to endocrine therapy [46-48]. pAKT is also elevated in mammary glands of NRL-PRL females [17]. The majority of the carcinomas in this panel displayed strong pAKT labeling in both the nuclei and cytoplasm of tumor epithelium (Table 1, Figure 3d). Other tumors exhibited pAKT labeled cells in both the stroma and epithelium, and a minority in the stromal compartment alone (Figure 3e). Unlike pERK1/2, pAKT expression was not correlated with a specific histotype; carcinosarcomas demonstrated variable expression levels of nuclear and cytoplasmic pAKT (Figure 3f, g).

Together, these observations suggest that a broad spectrum of signals may contribute to progression of these carcinomas. Nuclear pStat5 is elevated in hyperplastic lesions in NRL-PRL glands [17], and early ablation of Jak2 protects against PRL-induced tumors [49], consistent with an important role for the Jak2-Stat5 pathway in their genesis. However, the data presented here indicate that high pStat5 persists in only a minor subset of established PRL-induced carcinomas, suggesting that other signals drive progression of most of these tumors. Elevated activities of AP-1 [50,51], AKT [52] and MAP kinases, including ERK1/2 [43], have been implicated in resistance to conventional chemotherapies and anti-estrogens. PRL can activate these signals alone, as well as potentially cooperate with growth factors to enhance ERK1/2 and AKT activation *in vitro* and *in vivo* [53-55]. In light of the variable levels of *Prlr*

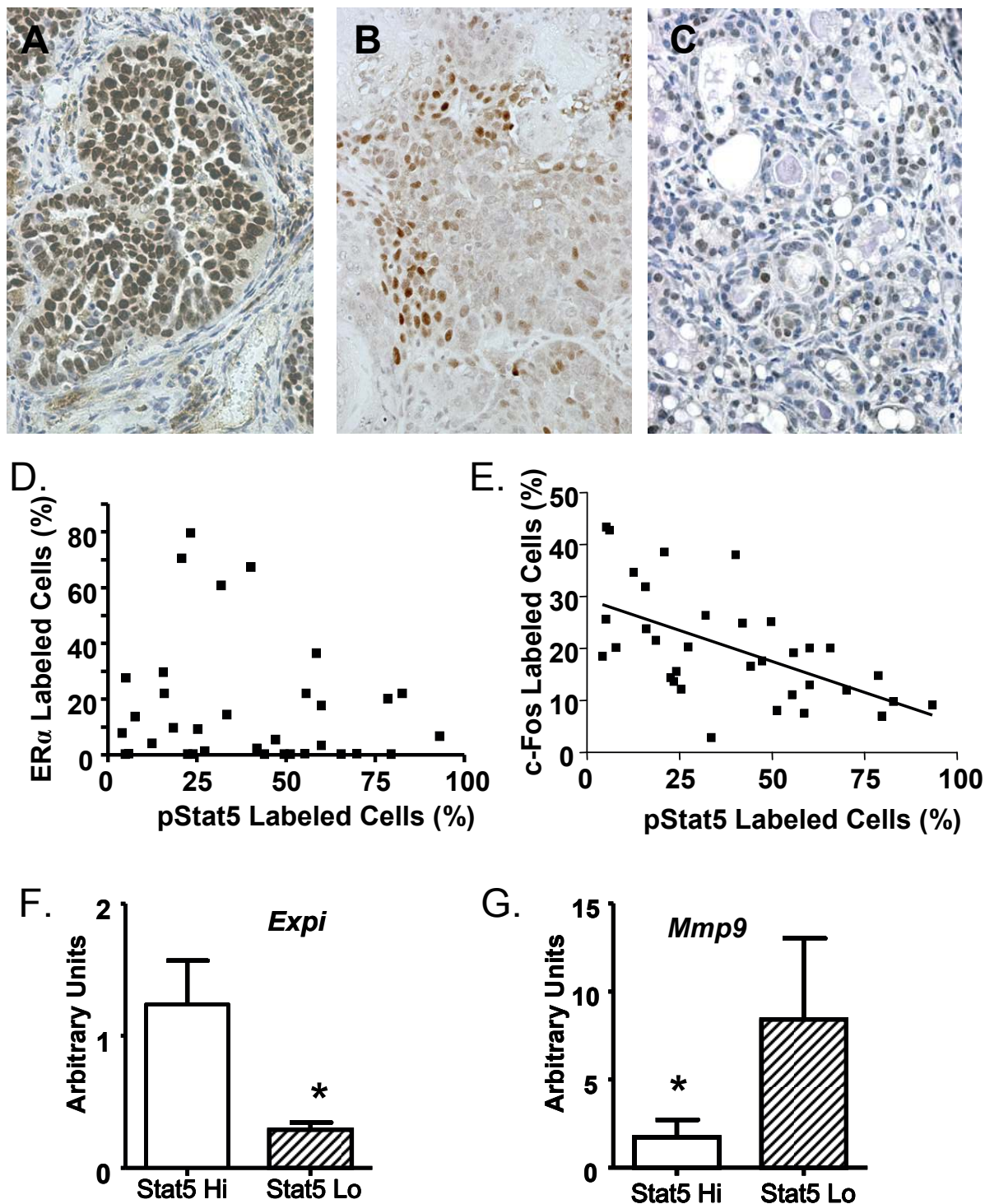
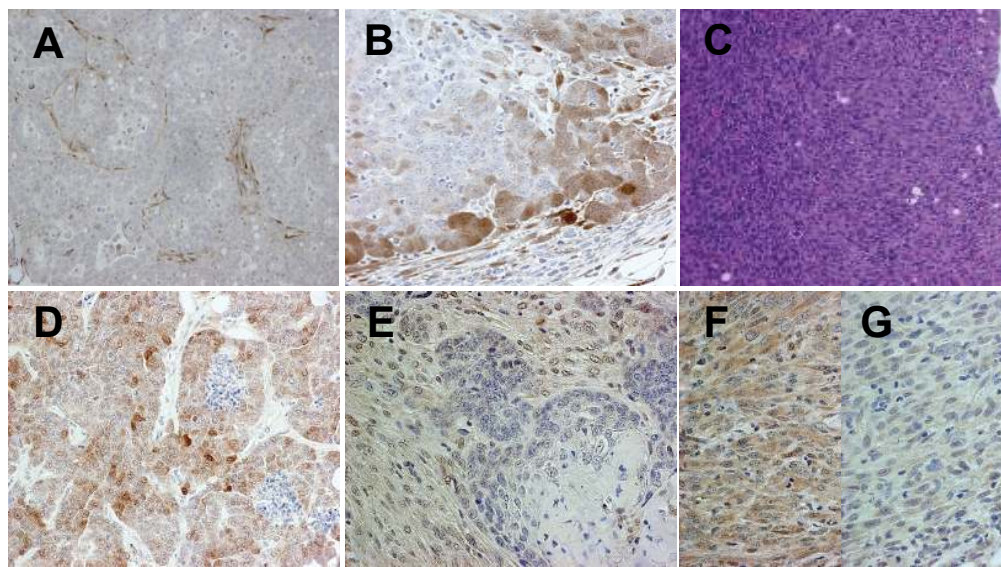
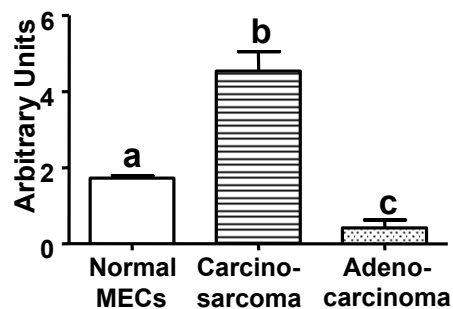


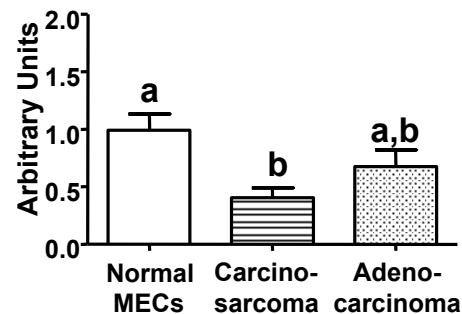
Figure 2 PRL-induced carcinomas demonstrate an inverse relationship between nuclear phosphorylated Stat 5 (pStat 5) and AP-1 proteins. **(a)** Nuclear pStat 5 staining in a glandular adenocarcinoma. **(b)** c-Fos expression in a glandular adenocarcinoma. **(c)** c-Jun staining in a glandular adenocarcinoma. **(d)** The proportion of tumor cells that express ER α and nuclear pStat5 is not correlated. **(e)** Carcinomas exhibit a significantly negative correlation between the proportion of cells that contain nuclear pStat 5 and c-Fos ($P < 0.0001$, $R = -0.63$). ER α , pStat 5, c-Fos and c-Jun expression were evaluated and quantitated as described in Materials and Methods. Each box represents a single carcinoma that arose in a different NRL-PRL nonparous female. Correlations were determined using Spearman's non-parametric test. **(f)** *Expi* mRNA is significantly increased in the adenocarcinomas expressing highest pStat5, compared to the lowest 5 (mean \pm s.e.m., $N = 5$; *, $P = 0.003$). **(g)** *Mmp9* mRNA is significantly decreased in the adenocarcinomas expressing highest pStat5, compared to the lowest 5 (mean \pm s.e.m., $N = 5$; *, $P = 0.03$). (f, g) Transcript levels were measured using qRT-PCR as described in Materials and Methods. Original magnification: a, b, c, 400x.



H. *Vimentin*



I. *Keratin 8*



J. *Mmp9*

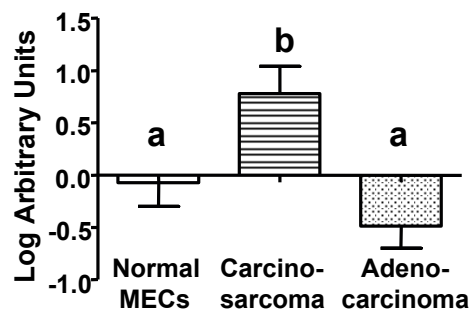


Figure 3 Adenocarcinomas exhibit activated ERK1/2 in stroma and AKT in epithelia; carcinosarcomas display strong epithelial pERK1/2. (a) pERK1/2 localized in stromal cells of differentiated glandular adenocarcinomas. (b) Mammary carcinosarcomas demonstrated nuclear and cytoplasmic staining for pERK1/2. (c) Carcinosarcoma from gland of NRL-PRL female (hematoxylin and eosin stain). (d) Glandular adenocarcinoma from NRL-PRL female demonstrating nuclear and cytoplasmic staining for pAKT. (e) pAKT localized primarily in the stroma of a squamous adenocarcinoma. (f, g) Carcinosarcomas demonstrated variable levels of pAKT expression. (h) Carcinosarcomas demonstrated significantly higher levels of vimentin mRNA compared to either morphologically normal MECs or more differentiated adenocarcinomas from NRL-PRL females. (i) Carcinosarcomas had similar levels of keratin 8 transcripts compared to adenocarcinomas. (j) *Mmp9* mRNA was significantly increased in carcinosarcomas. Transcript levels were measured using qRT-PCR as described in Materials and Methods and represented as mean \pm s.e.m. (MECs, $N = 3$ samples, each containing RNA pooled from 5 mice; carcinosarcomas, $N = 4$; adenocarcinomas with high pStat5, $N = 5$). Letters denote significant differences using ANOVA, followed by Newman-Keuls post test ($P < 0.05$). Original magnification: a, c, 200x; b, d, e, f, g, 400x.

transcripts in these tumors (see below), it is clear that PRL itself may play different roles in ongoing activation of these pathways as they influence lesion progression, tumor phenotype, and treatment sensitivity.

PRL-induced carcinomas share molecular features of luminal tumors in women

To evaluate shared characteristics of PRL-induced carcinomas, in a preliminary study we examined the transcript profiles of six NRL-PRL carcinomas, including three each of the most prevalent histotypes (glandular and papillary), expressing variable levels of ER α (three positive, three negative by IHC, not associated with histotype). Unsupervised hierarchical clustering based on an intrinsic gene list developed for murine models [31] showed that the NRL-PRL tumors formed a distinct subgroup neighboring MMTV-neu tumors.

To compare the phenotype of NRL-PRL tumors more closely to other well-studied genetically modified murine models of breast cancer subtypes in the FVB/N strain background, we compared levels of transcripts for ER α -associated genes in the “luminal” signature [20] and those enriched in various mammary epithelial lineages. MMTV-neu tumors model clinical HER2+ tumors. Like both human HER2+ tumors [25] and other murine tumors that develop in response to MMTV-driven oncogenes [31], MMTV-neu tumors display some characteristics of the “luminal” tumor subtype in women. Tumors in this extensively studied mouse model do not express ER α or FoxA1, a transcription factor which co-regulates many targets [56,57]. However, they do express other ER α -associated genes, including Xbp1 and Gata 3 [58-60]. In contrast, tumors that develop in both C3(1)-SV40 Tag females and transplanted p53^{-/-} models have many features of “basal” tumors in women ([31,61], O’Leary and Schuler, in prep).

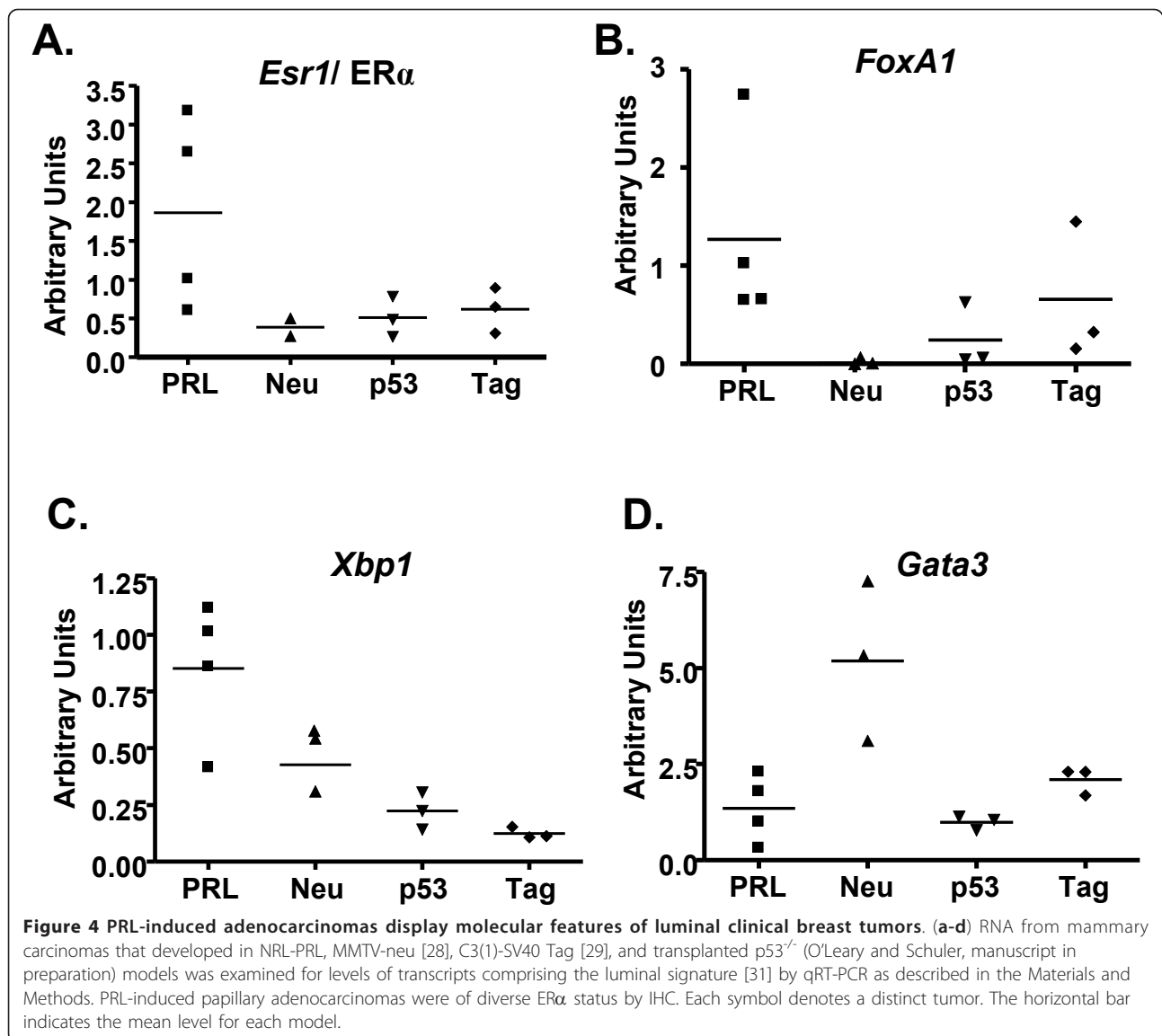
Unlike the morphologically homogeneous MMTV-neu tumors, levels of transcripts that mark luminal breast cancers varied in NRL-PRL carcinomas, consistent with the diverse tumors in this model (Figure 4, Figure S1 in Additional File 1). In contrast to the majority of murine models, PRL-induced adenocarcinomas expressed relatively high levels of *Esr1* mRNA, although as predicted from the IHC, levels varied considerably (Figure 4a). PRL-induced papillary carcinomas also contained comparatively high levels of the ER α -associated transcripts in the luminal signature, including *FoxA1* and *Xbp1* (Figure 4b, c). *Gata3* mRNA was readily detectable in NRL-PRL tumors, but levels were lower than in MMTV-neu tumors (Figure 4d). Although *Gata3* specifies ductal/alveolar cell fate and is markedly reduced in *Prlr*^{-/-} glands [1], this transcription factor also maintains luminal differentiation and is reduced in invasive breast

cancer [62], consistent with the less differentiated phenotype of the PRL-induced carcinomas.

In light of the luminal phenotype of many tumors promoted by PRL, we examined levels of transcripts associated with distinct mammary subpopulations and differentiation processes [24,63]. PRL-induced tumors displayed relatively high levels of *Itga6* (encoding integrin $\alpha 6$, CD49f) mRNA, a marker for stem cell and progenitor populations (Figure 5a). They expressed relatively low levels of transcripts for both *Itgb1* (encoding integrin $\beta 1$, CD29), which marks stem cell and basal but not luminal subpopulations (Figure 5b), and also *Itgb3* (encoding integrin $\beta 3$, CD61), a marker for luminal progenitors (Figure 5c). In contrast, three of the four NRL-PRL adenocarcinomas displayed relatively high levels of CD44 (Figure 5d), an adhesion molecule associated with stem cells and metastasis [64,65]. NRL-PRL adenocarcinomas also contained elevated mRNA for several transcription factors, which direct luminal/alveolar cell expansion and differentiation, are co-expressed with PRLR in many normal MECs, and are linked to PRL in genetic models [1,66]. PRL-induced adenocarcinomas contained relatively high levels of mRNA for *Elf5*, a PRL-induced factor that directs the alveolar lineage [67], as well as *CEBP β* , a transcriptional regulator which has been implicated in stem cell and luminal progenitor repopulation [68] (Figure 5e, f). Consistent with their roles in luminal cell commitment, individual genetic ablations of *Gata3*, *Elf5*, *C/EBP β* and also *Stat5* have been shown to alter the size of the MEC subpopulation expressing integrin $\beta 3$ (CD61+), which is enriched in luminal precursors. However, the variable direction and timing of the observed changes reflect their complex actions at more than one stage of this process [24,68,69]. Future studies will elucidate the effect of PRL on mammary epithelial subpopulations and roles of these transcriptional regulators in its actions, and illuminate differences in cellular origin and/or pathways of progression from MMTV-neu tumors and among the different histotypes of PRL-induced carcinomas.

Transcripts for the cdk4/6 inhibitor, *Cdkn2c* (p18^{INK4c}), were very low (Figure 5g). Genetic ablation of this tumor suppressor induces well-differentiated non-invasive luminal tumors in mouse models [70]. Similar results were observed for *Cdkn2a* (p16^{INK4a}, data not shown). These results are consistent with the similar mammary phenotypes of *Prlr*^{-/-} and *Ccnd1*^{-/-} mice [1], and ability of PRL to increase cyclin D1 transcription [71], and support a key role for cdk4/6 in PRL-induced tumorigenesis.

PRL also may contribute to the tumor phenotype of other models; some C3(1)-SV40 Tag as well as MMTV-neu tumors contained appreciable *Prlr* mRNA (Figure 5h).



The CD61+ progenitor population has recently been shown to be the origin of basal tumors [72]; the size of this MEC subpopulation is strongly modulated by Stat5. These observations invite speculation on potential therapeutic or preventative approaches directed at PRL action in multiple tumor types.

These marked differences in the molecular phenotype of PRL-induced tumors compared to other genetically modified mouse models of breast cancer suggest that PRL acts on its physiological target cells to promote diverse luminal-type tumors, but raises questions about the identity of these cells, the role of mediators associated with “normal” function in tumor progression, and factors contributing to tumor heterogeneity. Development of additional markers for the various mammary epithelial subpopulations, and antibodies that recognize

the native conformation of the murine PRLR will assist in determining the characteristics of the precursors of luminal tumors, and identifying the target cells of PRL in this process.

ER α expression is associated with distinct molecular features, but does not confer estrogen responsiveness

In our preliminary studies comparing PRL-induced tumors to other murine models of breast cancer, PRL-induced ER α - and ER α + tumors clustered together, indicating shared features suggesting a common origin. In order to determine characteristics of PRL-induced tumors that are associated with differences in ER α protein expression, we compared transcript profiles from four ER α + and four ER α - adenocarcinomas (determined by IHC) of divergent histotypes that did not segregate

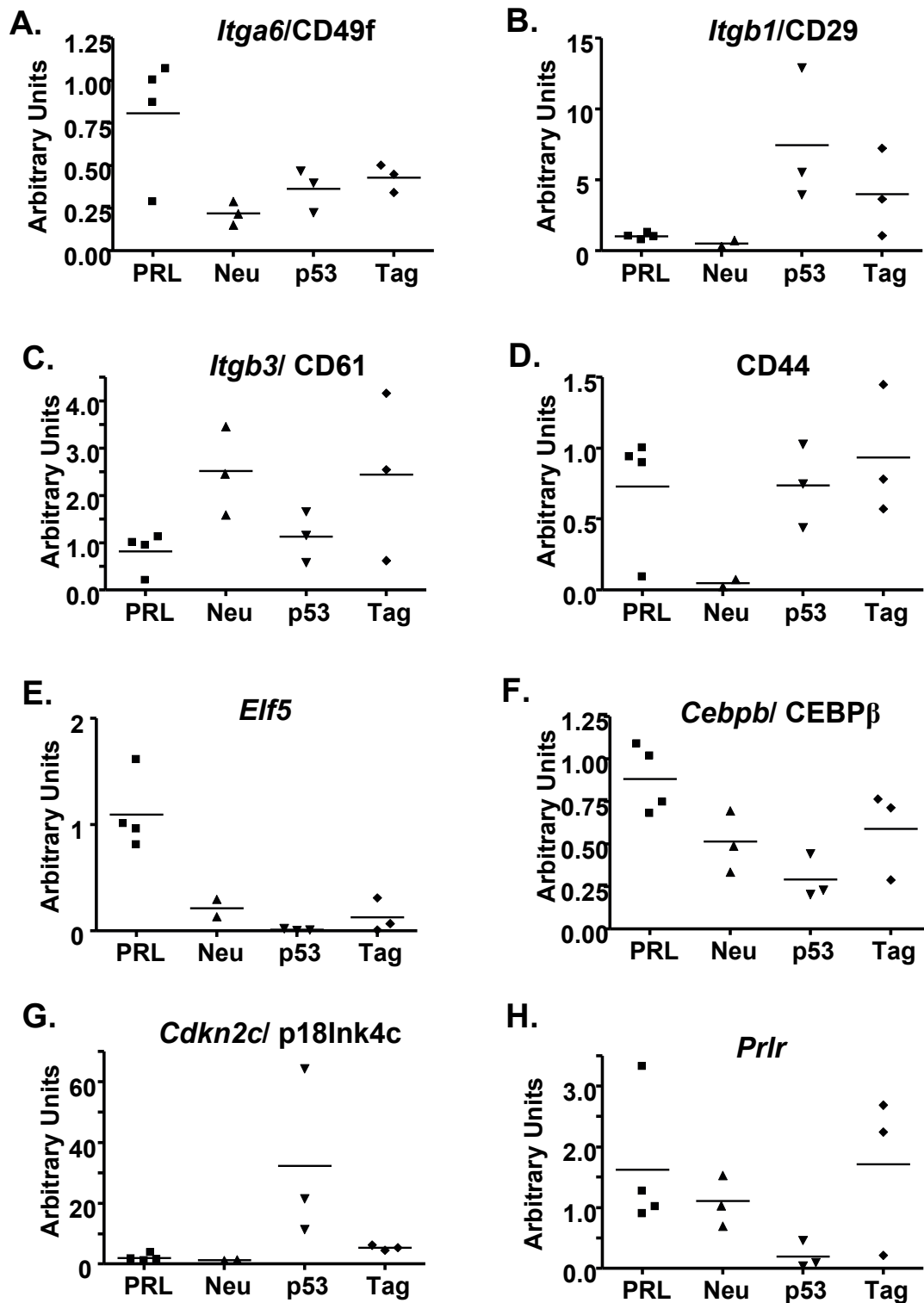


Figure 5 NRL-PRL adenocarcinomas express markers distinct from other well-characterized models of breast cancer. RNA from the set of mammary carcinomas in Figure 4 that developed in NRL-PRL, MMTV-neu [28], C3(1)-SV40 Tag [29], and transplanted p53^{-/-} (O'Leary and Schuler, manuscript in preparation) models was examined for levels of transcripts implicated in mammary lineages by qRT-PCR as described in the Materials and Methods. Each symbol denotes a distinct tumor. The horizontal bar indicates the mean level for each model.

with ER α status (three papillary, two glandular, one adenocarcinoma, two solid). Genes that were differentially expressed among the classes were determined, and hierarchical clustering analyses were performed by comparing expression profiles both across the set of samples and across the set of genes. As shown in Figure 6a, ER α + and ER α - tumors clustered separately. Consistent with the reports of ER α + tumors in women, NRL-PRL ER α + tumors expressed higher levels of *Gata-3* than ER α - tumors, and markers of differentiation, such as the milk proteins, β -casein and whey acidic protein, and CD24a [20,73-75]. Levels of mRNA for the PRL transgene and PRLR varied markedly among both ER α + and ER α - carcinomas, suggesting that PRL activity itself was not responsible for the differences observed (Figure S2 in Additional File 1).

Activity of the NRL-PRL promoter is not altered by estrogen [15], enabling us to examine the role of this steroid in the disease process. Proliferation of both morphologically normal MECs and preneoplastic epithelial hyperplasias in NRL-PRL females is very sensitive to ovariectomy (ovx) and supplemental estrogen [17,54]. In order to examine estrogen responsiveness in established carcinomas, NRL-PRL females bearing primary mammary adenocarcinomas were subjected to sham surgery or ovx. At the time of surgery, a biopsy was removed for examination of tumor histology, rate of proliferation, and steroid hormone receptor expression. The growth of the remaining tumors was monitored until they reached 1.5 cm in diameter (15.3 ± 8.5 days; mean \pm s.d.), as described in the Materials and methods. Like the tumors in Figure 1, histotypes in this experimental panel were diverse, and ER α /PR expression varied widely. They exhibited highly variable rates of growth (Figure 6b); rates of proliferation of ER α - tumors prior to the surgery tended to be higher than ER α + tumors ($P = 0.07$). However, ovx did not alter rates of proliferation of either ER α + or ER α - tumor cells. Consistently, ovx followed by administration of E2 failed to alter proliferation of an independent set of primary tumors (PCNA staining: $30.1\% \pm 12$ vs $31.19\% \pm 7.9$, mean \pm s.d., $P > 0.05$, $N = 5$).

Since primary PRL-induced tumors exhibit highly variable histotype and ER α expression, and ovarian function is likely to be faltering in the aged females, we employed tumor transplantation to further investigate the role of ER α -dependent signals in established tumors. We serially passaged NRL-PRL glandular adenocarcinoma fragments that demonstrated high expression of both ER α and PR (48% and 25% labeled cells, respectively) to bilateral inguinal mammary glands of nonparous 12-week-old syngeneic nontransgenic recipients. Although the tumors that grew from these fragments (about eight weeks per passage) retained the morphology and high

ER α expression of the original tumors, PR was no longer detectable after passage (not shown), suggesting that this process selects for more aggressive tumor cells. After three passages, recipient females were treated with ovx, sham surgery, or sham surgery and the ER α -selective antagonist, Faslodex. None of these treatments altered rates of proliferation of the tumors, as measured by BrdU incorporation (Figure 6c), indicating that relatively well-differentiated ER α + tumors display the estrogen independence observed in the more diverse group of primary tumors.

These findings raise interesting questions regarding the mechanism(s) whereby PRL interacts with ER α -mediated signals in the development and progression of ER α + tumors. The striking estrogen independence of established PRL-induced tumors shown herein contrasts with the cooperation between PRL and estrogen in proliferation of morphologically normal MECs and early lesions [17,54], suggesting acquisition of estrogen insensitivity. Many primary clinical ER α + tumors acquire resistance to estrogen-directed therapies and most display marked therapeutic insensitivity after metastasis. A vast literature implicates many mechanisms, including aberrations not only of ER α itself, but alterations in other molecules and signaling pathways consistent with heightened growth factor activity [76-78]. The ability of PRL to potentiate growth factor signals [53-55,79] and high activation of AKT in the tumors shown here are consistent with a role in the latter mechanism. Clinically, high circulating PRL has been associated with failure of ER α + tumors to respond to tamoxifen and aromatase inhibitors [80-82], and PRLR transcripts were elevated with recurrence after tamoxifen treatment in some studies [13,83]. The histological heterogeneity and variable activation of signaling pathways of PRL-induced tumors observed here suggest that multiple mechanisms may lead to loss of estrogen sensitivity. Together, these features suggest that the NRL-PRL model may provide insight into the origin of the diversity of clinical ER α tumors, and the characteristics of the tumor subpopulation that resists therapy.

Conclusions

Despite the prevalence of ER α + tumors clinically, very few mouse models resemble this breast cancer subtype. Our studies demonstrate that elevated local PRL can promote diverse carcinomas, which display varying levels of ER α /PR, nuclear pStat5, pERK1/2 and pAKT, and AP-1 components. Transcript analysis demonstrates that many of these carcinomas express the ER α -associated transcript signature that defines this clinical tumor subtype, and suggests that they arise from the physiologic target cells of PRL. In contrast to morphologically normal structures and hyperplastic lesions,

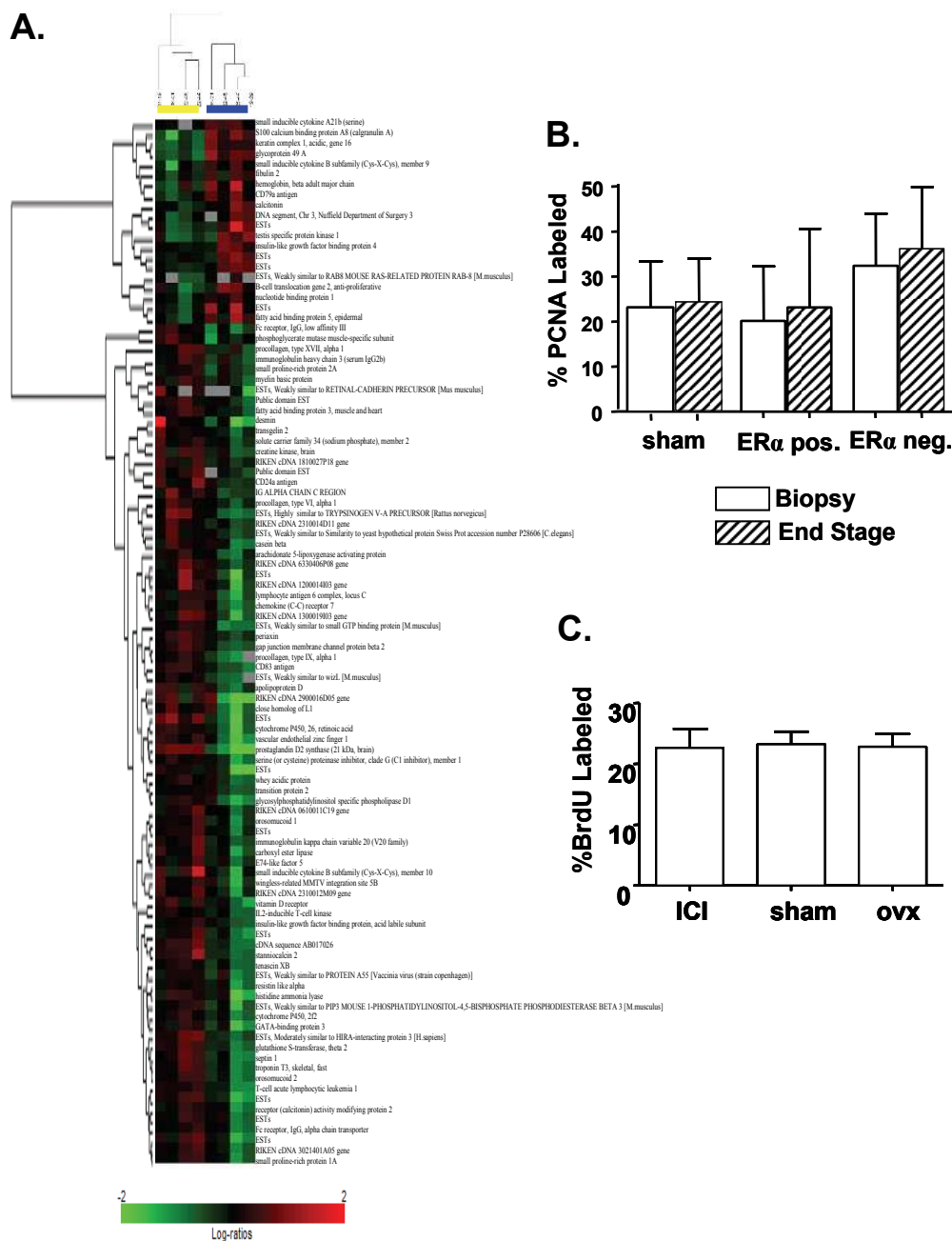


Figure 6 ERα+ and ERα- carcinomas demonstrate distinct transcript profiles, and are insensitive to ovarian steroids. (a) Hierarchical clustering of genes and tumor samples demonstrates differential expression between ERα+ (yellow) and ERα- (blue) adenocarcinomas at the $P < 0.01$ significance level ($N = 4$ each). ERα+ was defined as $\geq 10\%$ of the cells contained detectable ERα by IHC; ERα- contained $< 10\%$. Red indicates high relative expression in ERα positive adenocarcinomas; green, low relative expression; black, no difference. Each column represents an independent tumor, and each row is a gene. **(b)** Growth of PRL-induced tumors is not diminished by ovariectomy, regardless of ERα status. Mammary carcinomas of distributed histotypes in NRL-PRL nonparous females were biopsied, and the mice subjected to sham surgery or ovariectomy. Subsequent tumor growth was monitored until tumors reached 1.5 cm in diameter (15.3 ± 8.5 days; mean \pm s.d.), and proliferation determined by PCNA IHC as described in the Materials and Methods. Sham, $N = 12$; ovx ERα+, $N = 10$; ovx ERα-, $N = 9$. Treatment did not alter the rate of growth (Student's paired t -test, $P > 0.05$). **(c)** Treatment with the ER inhibitor and downregulator, Faslodex (ICI 182,780), does not inhibit growth of ERα positive tumor fragments. Fragments of well-differentiated ERα+ tumors were transplanted into fat pads of nontransgenic recipients and treated with ovariectomy, sham surgery, or Faslodex as described in Materials and Methods. Ovariectomized and Faslodex-treated females had significantly reduced uterine weights (27 ± 5 mg, 30 ± 8 mg, mean \pm s.d., respectively), compared to sham-treated females (83 ± 5 mg). Treatment did not alter the rate of proliferation ($P > 0.05$).

established carcinomas are strikingly insensitive to estrogen, suggesting acquisition of independence. The role of ongoing PRL signals in established tumors will require additional investigation. Recent studies suggest that PRL also can antagonize traditional chemotherapeutics in breast cancer cells *in vitro* [84,85], which is supported by small studies of PRL on therapeutic resistance *in vivo* [86]. Together, our findings suggest that the NRL-PRL mammary tumor model will be helpful to understand the pathogenesis and heterogeneity of luminal tumors and identify factors determining susceptibility to anti-estrogen and traditional chemotherapies, to examine the role of PRL in these responses, and to test novel therapeutic strategies, including combinatorial regimens including therapies directed at PRL.

Additional material

Additional file 1: Supplementary tables and figures. Table S1: Immunohistochemistry conditions. Description of retrieval methods, blocking conditions and antibody dilutions used for each antigen examined. Table S2: Primers employed for RT-PCR analyses. Sequences of the primers used to quantify transcripts of interest. Figure S1: PRL-induced carcinomas of different histotypes display variable levels of transcripts for ER α -associated genes. Levels of mRNA for various transcripts of interest in individual PRL-induced tumors of different histotypes, determined by qRT-PCR. Figure S2: Levels of transgene and Prlr transcripts are variable in NRL-PRL adenocarcinomas, and are not associated with ER α status. Levels of mRNA for rPRL transgene and cytokeratin 8 (Krt8), and PRL receptor in individual tumors, determined by qRT-PCR.

Abbreviations

AP-1: Activating Protein 1; BrdU: bromodeoxyuridine; EMT: epithelial mesenchymal transition; ER α : estrogen receptor alpha; ERK: extracellular signal regulated kinase; IHC: immunohistochemistry; MECs: mammary epithelial cells; MMTV: mouse mammary tumor virus; PCNA: proliferating cell nuclear antigen; PR: progesterone receptor; PRL: prolactin; PRLR: prolactin receptor; qRT-PCR: quantitative reverse-transcriptase polymerase chain reaction; NRL: neu-related lipocalin; Stat5: signal transducer and activator of transcription 5.

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Author details

¹Department of Comparative Biosciences, University of Wisconsin-Madison, 2015 Linden Dr., Madison, WI 53706, USA. ²Department of Cancer Biology, Kimmel Cancer Center, Thomas Jefferson University, 233 South 10th St, 332 BLSB, Philadelphia, PA 19107, USA. ³Department of Anatomy & Cellular Biology, Sackler School, Tufts University School of Medicine, 136 Harrison

Ave, Boston, MA 02111, USA. ⁴Genetics Department, Navarra University, Irunlarrea St. 1, Pamplona, 31008, Spain.

Authors' contributions

LMA and LAS contributed to concept design, data analysis and interpretation, and manuscript writing. DER carried out the transcript analyses and contributed to interpretation. TAG-H contributed to the generation of experimental data and analysis. MJG-B and HR analyzed the pStat5 expression of the tumors. All authors read and approved the final manuscript.

Competing interests

The authors declare that they have no competing interests.

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