The permissive role of prolactin as a regulator of luteinizing hormone action in the female mouse ovary and extragonadal tumorigenesis

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Bachelot A, Carré N, Mialon O, Matelot M, Servel N, Monget P, Ahtiainen P, Huhtaniemi I, Binart N. The permissive role of prolactin as a regulator of luteinizing hormone action in the female mouse ovary and extragonadal tumorigenesis. Am J Physiol Endocrinol Metab 305: E845–E852, 2013. First published August 6, 2013; doi:10.1152/ajpendo.00243.2013.—Transgenic female mice overexpressing the hCG subunit (hCGβ⁺) and producing elevated levels of luteinizing hormone (LH)/hCG bioactivity present as young adults with enhanced ovarian steroidogenesis, precocious puberty, and infertility. They subsequently develop pituitary prolactinomas, high circulating prolactin (PRL) levels, and marked mammary gland lobuloalveolar development followed by adenocarcinomas. None of these phenotypes appear in gonadectomized mice, indicating that the hCGβ⁺/PRLR⁻/⁻ females remained sterile with an ovarian phenotype similar to PRLR⁻/⁻ females, indicating that LH action, Lhcg expression, and consequent luteinization are not possible without simultaneous PRL signaling. The high frequency of pituitary prolactinomas in PRLR⁻/⁻ mice was not affected by transgenic hCGβ⁺ expression. In contrast, none of the hCGβ⁺/PRLR⁻/⁻ females showed either mammary gland lobuloalveolar development or tumors, and the increased mammary gland Wnt-5b expression, possibly responsible for the tumorigenesis in hCGβ⁺ mice, was absent in double-mutant mice. Hence, high LH/hCG stimulation is unable to compensate for missing PRL signaling in the maintenance of luteal function. PRL thus appears to be a major permissive regulator of LH action in the ovary and of its secondary extragonadal effects. Prolactin receptor-deficient (PRLR⁻/⁻) female mice are sterile due to the failure of embryo implantation, which is, for a large part, due to the failure of corpus luteum (CL) maintenance, the consequence of decreased Lhcg mRNA in the corpora lutea compared with those of wild-type (WT) ovaries, and that the LHR in hCG-treated PRLR⁻/⁻ animals was functional (5). The expression of Lhcg was accompanied by increased expression of the Wnt-5b/Wnt signaling pathway, resulting in β-catenin-stabilizing mammary tumorigenesis (24).

Transgenic (TG) mouse models have confirmed the key role of gonadotropins in normal reproductive function and as oncogenic factors in gonadal and extragonadal tumorigenesis (1, 21). To assess the consequences of prolonged exposure to elevated levels of luteinizing hormone (LH)/choriongonadotropin (hCG) bioactivity, TG mice expressing the hCG subunit under the ubiquitin C promoter (hCGβ⁺/hCGβ⁻ mice) have been developed (38). Female TG mice present initially with precocious puberty, infertility, enhanced ovarian steroidogenesis, and elevated prolactin (PRL) levels. Several extragonadal phenotypes develop in the mice upon aging, including pituitary hyperplasia at the age of 2 mo, followed by gradual growth of pituitaries to macroprolactinomas at the age of 6 mo (2). In addition, marked mammary gland lobuloalveolar development, followed by adenocarcinomas, is observed at the age of 9–12 mo (38). Although Lhcg expression has been detected in pituitary, adrenal, and breast tissues (3), it is apparent in this model that all extragonadal phenotypes are induced through hCG-stimulated ovarian endocrine activity because they all failed to occur if the mice were gonadectomized before puberty (38). The endocrine aberration of the hCGβ⁺/hCGβ⁻ mice with persistently elevated levels of PRL and progesterone, and transient peripherally elevated estradiol, was most probably crucial for the mammary gland tumorigenesis, although direct extragonadal hCG effects cannot be excluded (20). A tumor promoter role for LH has also been proposed on the basis of other TG mouse models, including inhibin-deficient mice (26) and those expressing the Simian virus 40 T-antigen under inhibin α-subunit promoter (23, 36). Transdifferentiation is observed in the mammary tumors of the hCGβ⁺/hCGβ⁻ mice, accompanied by abnormal expression of the Wnt-5b and -7b genes in the tumors and nontumorous mammary glands (24). Importantly, hCG was found to upregulate these Wnt ligands in the mouse mammary gland, independent of the changes in ovarian steroidogenesis (24). Thus, the hCGβ⁺/hCGβ⁻ mice represent a novel model that links enhanced hCG action to dysregulated Wnt signaling in the mammary gland, resulting in β-catenin-stabilizing mammary tumorigenesis (24).

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hydroxysteroid dehydrogenase (20α-HSD), a PRL repressed luteal enzyme that catabolizes and inactivates progesterone.

To dissect out further the specific contribution of the different hormones, i.e., LH/hCG, PRL, and steroids, in the gonadal and extragonadal phenotypes displayed by the hCGβ+ and PRLR−/− mice, we intercrossed in the present study the TG hCGβ+ and PRLR knockout mice.

MATERIALS AND METHODS

Animals. TG female mice of the FVB/N strain expressing heterozygously the hCGβ-subunit cDNA, driven by the human ubiquitin C promoter, and their genotyping and housing have been described before (38). Non-TG littermates were used as controls. WT and PRLR+/- female mice (129/SvJ) were generated as previously described (31) and then backcrossed by successive 12 breedings to a pure genetic FVB/N background. hCGβ+ male and PRLR−/− mice were intercrossed to obtain hCGβ+ and hCGβ+/PRLR−/- female mice. PCR analysis of tail DNA was used to determine the genotypes of the offspring as described previously (31, 38). Animals were housed in a 12:12-h day-night cycle at 22°C and relative humidity of 50% with food and water ad libitum. Mice were bred according to the guide for the Care and Use of Laboratory Animals published by the United States National Institute of Health (NIH Publication No. 85-23, revised 1996). The animal facility was granted approval (No. C94-043-12), given by the French Administration. All procedures were approved by the local ethic committee CAPSud (No. 2012-021).

Fertility studies. For testing the fertility, hCGβ+ and hCGβ+/PRLR−/− female mice were bred with WT males for up to 6 mo, and vaginal plugs were checked. The duration of estrous cycles was determined in six hCGβ+ and hCGβ+/PRLR−/− female mice and WT mice by daily analysis of the vaginal smears for a period of 4 wk. For implantation study, the morning a vaginal plug was found was designated day 0.5 of pregnancy. On day 5.5, the mice were killed, and the number of implantation sites was recorded by monitoring the localized uterine vascular permeability at the sites of blastocysts after intravenous injection of Chicago blue dye solution in saline (Sigma, St. Louis, MO).

Histological analysis. Tissues (ovaries, pituitaries, and adrenal glands) were fixed overnight in 4% paraformaldehyde, dehydrated, and embedded in paraffin, and 5-μm-thick sections were prepared. For histological studies, sections were stained with hematoxylin and eosin.

Mammary gland whole mounts. The fourth inguinal mammary glands were dissected out, spread on glass slides, and fixed overnight in a 1:3 mixture of glacial acetic acid-100% ethanol at room temperature. The tissues were washed in 70% ethanol, rinsed in water, and stained overnight in 0.2% carmine (Sigma) at room temperature. The stained tissues were dehydrated through graded series of ethanol, cleared in toluene, and mounted.

Binding assays on histological sections. Ovaries were collected and coated with cryoprotectant embedding medium (Tissue-Tek; Miles, Elkhart, IN), frozen in cold isopentane, and then stored at −20°C. The binding of 125I-labeled hCG to ovarian frozen sections was studied by an autoradiographic method, as described previously (17). Briefly, ovaries were serially sectioned at a thickness of 5 μm with a cryostat. After fixation for 10 min at 4°C in picric acid-formaldehyde, sections were stored at −20°C and then circled with Depex (Gurr; BDH, Poole, UK). hCG was iodinated by the Iodogen method (Sigma, St. Quentin, France) and purified by Sephadex G-50 chromatography. Sections were incubated in a drop of PBS (0.1% BSA, pH 7.4) containing 125I-labeled hCG (4 × 105 cpm/50 μl). To determine nonspecific binding, for the tested ligand an adjacent serial section of ovary was incubated with an excess of unlabeled ligand (500 ng/50 μl hCG). At the end of the incubation period, the sections were washed two times in PBS, postfixed in 3% glutaraldehyde-PBS, washed in PBS, air-dried, and stained with Feulgen. For autoradiography, they were then dipped in Kodak NTB2 emulsion diluted 1:1 with distilled water, air-dried, exposed for 2 wk at 4°C, and then developed and fixed by classical procedures.

RNA isolation and quantitative RT-PCR. The fourth mammary gland or mammary gland tumor was excised and snap-frozen in liquid nitrogen. Total RNA was extracted from the tissues using Trizol reagent (Invitrogen). Quantitative real-time PCR was performed as described previously (44). After DNase I treatment, RNA was reverse-transcribed and used for quantitative RT-PCR using the Power SYBR Green PCR Master Mix (Applied Biosystems). All primer pairs used were intron spanning: Wnt 5b sense, TGGAGAACCGTG-GAGTACC; Wnt 5b antisense, GGCGACATCGCCATCTTAT; product size 166 bp; T annealing 60°C. The presence of a single, correct-sized PCR product was confirmed by running the samples on a 2% agarose gel. Final primer concentrations were 300 nM. Reaction parameters were carried out on a StepOne Real-Time PCR System (Applied Biosystems). Relative expression within a given sample was calculated as a ratio (amol of specific gene/μmol of 18S).

Statistical analysis. Descriptive statistics were performed for each variable, and parameters were compared using the Mann-Whitney test. Proportions for the two groups were compared using the χ2-test. All values are expressed as means ± SE of combined data from the replicate experiments. Tumor-free survival curves were carried out by the Kaplan-Meier method and compared with the log-rank test; all data were expressed as means ± 95% confidence intervals. Analyses were processed with GraphPad Prism 5 (GraphPad Software, La Jolla, CA). Values of P < 0.05 were considered statistically significant.

RESULTS

Fertility, estrous cycles, and ovarian histology. hCGβ+ females presented, at 3 mo of age, with irregular estrous cycles, with only one out of four mice exhibiting estrus during the 4-wk period of observation and the remaining mice showing a continuous diestrous pattern. Next, at 6 mo of age, all hCGβ+ females (n = 10) presented with a diestrus-type pattern, and no pregnancies were observed during the 6-mo breeding period with WT males. hCGβ+/PRLR−/− mice exhibited regular estrous cycles [duration: 6.8 ± 1.5 days (n = 4) vs. 6.7 ± 0.5 days in WT mice (n = 8), P not significant] at 3–10 mo of age. However, despite the regular estrous cycles, hCGβ+/PRLR−/− females could not achieve pregnancy following breeding with WT males (observation for up to 6 mo), despite the presence of periodic vaginal plugs. We examined the number of implantation sites in four hCGβ+/PRLR−/− females on day 5.5 using the blue dye method. No implantation sites were observed, in compliance with the phenotype already described in the PRLR−/− model mice (9). At diestrus, histological analysis of the ovaries in all genotypes at 3 and 6 mo of age is shown in Fig. 1. The hCGβ+ mice presented with progressive luteinization of their ovaries, with presence of few follicles at various stages of maturation and multiple large luteomas. The hCGβ−/PRLR−/− ovaries showed all classes of follicles but absence of corpora lutea or luteomas, a phenotype similar to PRLR−/− females already described (9).

To examine whether the LHR protein was synthesized and able to bind the ligand hormone, we performed 125I-labeled hCG binding analysis on ovarian sections (Fig. 2). The results showed high levels of hCG binding activity in the whole hCGβ+ ovaries (apparently in luteoma tissue) compared with WT mice where LH-binding activity was mostly fond in
Corpora lutea. None of the hCGβ⁺/PRLR⁻⁻ and PRLR⁻⁻ mice showed corpora lutea or luteomas with clear LHR expression. High TG LH/hCG bioactivity was therefore not sufficient to reverse the female PRLR⁻⁻ reproductive phenotype, indicating that LH action, Lhcgr expression, and consequent luteinization did not occur in the absence of PRL signaling.

**Mammary gland development.** Whole mount analysis of the mature mammary glands from virgin mice of all genotypes, performed at the ages of 3, 6, and 9 mo, is shown in Fig. 3. There was a substantial and dramatic lobuloalveolar development from the age of 6 mo in the hCGβ⁺ compared with WT mice, reminiscent of the phenotype of the pregnant mammary gland. In contrast, none was observed in the hCGβ⁺/PRLR⁻⁻ and PRLR⁻⁻ female lobuloalveolar development; moreover, the side branching appears very poor compared with those of the WT mice. hCGβ⁺ mice developed aggressive mammary gland tumors after the age of 9 mo, with rapidly growing tumor mass and necrosis found in the center of the tumors (Fig. 4A). Based on histological criteria, the tumors were mammary infiltrating adenocarcinomas (Fig. 4B) or papillary pilar carcinomas (Fig. 4C) as the main subtypes. These tumors had a high prevalence compared with WT mice (P < 0.001, Fig. 5A), and they were never found in PRLR⁻⁻ or in hCGβ⁺/PRLR⁻⁻ mice during the observation period (Fig. 5A).

Wnt signaling in the mammary gland is dysregulated in the hCGβ⁺ mice, resulting in β-catenin-stabilizing mammary tumorigenesis (23). Results of the qPCR analysis of mammary glands and mammary tumors of the hCGβ⁺, hCGβ⁺/PRLR⁻⁻, and WT females are shown in Fig. 5B. Increased expression of Wnt-5b was present even in the absence of macroscopic tumors in the hCGβ⁺ mammary glands compared with control and hCGβ⁺/PRLR⁻⁻ and PRLR⁻⁻ mice (Fig. 5B). Similar results were observed for Wnt-7b expression (data not shown).

**Pituitary adenomas.** The pituitary glands of PRLR⁻⁻ mice contained massive, multifocal tumors, all monohormonal lactotroph adenomas, as well as the pituitary of hCGβ⁺ mice (3). Next, we studied the prevalence of pituitary adenomas in all genotypes. hCGβ⁺/PRLR⁻⁻ and PRLR⁻⁻ females had slightly enlarged pituitary glands at 6 mo of age and massively enlarged at 14 mo, leading to death of the animals (Fig. 6). hCGβ⁺/PRLR⁻⁻ and PRLR⁻⁻ mice exhibited tumors more frequently than did hCGβ⁺ or WT mice.

**Adrenal glands.** Chronically elevated serum LH, augmented by enhanced PRL production, has been shown to induce
functional *Lhcgr* expression in the adrenal cortex of specific strains of mice (8). Histological analysis of the adrenal glands, performed at the ages of 3, 6, and 9–12 mo, showed no differences between any of the genotypes, with an absence of hyperplasia or tumor development (data not shown). By LH-binding analysis, we were not able to evidence LHR in adrenal gland, whatever the genotype or the age of the mice (result not shown). We confirm here the earlier finding (8) that the FVB/N mouse strain is resistant to postgonadectomy/high gonadotropin-induced adrenal tumorigenesis.

Fig. 3. Whole mount analysis of mammary gland from WT, hCGβ⁺, PRLR⁻⁻, and hCGβ⁺/PRLR⁻⁻ virgin female mice performed at the age of 3, 6, and 9 mo. A representative mammary gland is shown (n = 5 genotype). There was substantial lobuloalveolar development in 6-mo-old hCGβ⁺ mice, compared with WT mice, reminiscent of the phenotype of the pregnant mammary gland. None of the PRLR⁻⁻ and hCGβ⁺/PRLR⁻⁻ females showed lobuloalveolar development. Scale bar = 500 μm.

Fig. 4. A: representative macroscopic examination of a mammary tumor in 12-mo-old hCGβ⁺ female mouse. B and C: hematoxylin and eosin staining of sections of two mammary gland tumors (infiltrating adenocarcinoma and differentiated carcinoma) in hCGβ⁺ female mouse at 12 mo of age.
DISCUSSION

We have previously shown that hyperprolactinemia is essential for the phenotypic defects in hCG-overexpressing mice (38). Hyperprolactinemia is induced in these mice by combined action of a peripubertal peak in ovarian estradiol production and persistent high progesterone levels produced by the ovarian luteomas. The reproductive phenotype of the mice includes completely disturbed cyclicity and infertility. The crucial role of hyperprolactinemia in this phenotype was demonstrated recently by short-term cabergoline treatment, which prevented the reproductive dysfunction of these mice (35).

Conversely, targeted disruption of the Prlr gene in female mice results in infertility by completely disturbing both the maintenance of CL, decreasing the expression of Lhcgr, and by increasing the expression of 20α-hsd, leading to progesterone insufficiency and blockage of embryo implantation and mammary gland development (5, 19). We observed in this study that chronic overexpression of hCG in PRLR−/− females was not able to rescue their fertility, supporting the permissive role of PRL in the maintenance of reproductive function. We showed also that the absence of PRL signaling protected the hCG+/PRLR−/−/− mice from mammary gland tumorigenesis that occurs in hCG+/− mice in the presence of functional PRLR and high ovarian steroid hormone production.

PRL regulates the expression of a number of genes important for CL function and which may secondarily maintain
progesterone secretion and thereby gestation (42). It has been shown early on that PRL is essential for Lhcg expression in rodent CL (16, 18, 33, 41), and the generation of the Prlrt-deficient mice further substantiated these findings (19). LH is an important stimulus of CL function in a number of species, including humans and rodents (42). Earlier studies have established the timing and the role of PRL and LH in the maintenance of luteal function from day 7 through day 11 of pregnancy (28, 37). Here we showed that no pregnancies were observed in hCGβ+/PRLR−/− females, despite presenting with regular cycles and ovulations, as evidenced by regular plug observation. Instead, the mice had implantation failure, a reproductive phenotype similar to PRLR−/− mice already described (9). Therefore, overexpression of hCG in the absence of PRL signaling was unable to restore luteinization, strengthening the evidence of PRL dependence of the regulation of Lhcg expression, as shown by hCG-binding analysis. Of note, we have previously reported that acute LH/hCG stimulation of PRLR−/− females can induce, if administered early during CL formation, the expression and binding activity of LHR in the CL in the total absence of PRL signaling (5). We also showed that LH/hCG stimulated the expression of steroidogenic enzymes in the absence of PRL signaling. However, despite these positive effects, the level of progesterone secretion remained too low to allow blastocyst implantation. These results suggested that, although the activation of ovarian steroidogenesis is possible without concomitant PRL action, the rapid catabolism of progesterone persisted due to the persistent upregulation of 20α-HSD in the absence of PRL effect.

One of the remarkable features of the hCGβ+ mouse model is the development of the mammary gland phenotype, culminating in formation of malignant mammary tumors in older age (24, 38). We showed in this study that the lack of PRL signaling is able to protect against mammary gland tumors. Indeed, TG and knockout mouse models have proven useful for elucidating their contribution to the regulation of mammary gland growth and differentiation (12, 13, 45, 46). Ovarian steroids and PRL are the likely inducers of mammary tumorigenesis in the hCGβ+ mice because it is completely prevented if the mice are gonadectomized before puberty (38). We have previously shown that PRLR is required for pregnancy-induced lobuloalveolar development. PRLR−/− mice exhibit a phenotype similar to that observed in the progesterone receptor-deficient mice. This was supported by the evidence that progesterone can regulate PRLR expression (30). Even if the side-branching defect observed in PRLR−/− mammary glands can be rescued with progesterone treatment, the lobuloalveolar development still did not occur (9).

Our results demonstrate that the precocious abnormal lobuloalveolar development of the mammary gland in hCGβ+ females is either dependent on ovarian endocrine hyperfunction or on PRL signaling. Accordingly, cross talk between the steroid hormones and the PRL pathway is able to synergize on mammary epithelium (14). The final way to dissect out the specific contribution of the different hormones would be to cross the hCGβ+ mice individually with mice deficient of the estrogen and progesterone receptors. Although Lhcg is expressed in mouse mammary gland, its direct stimulation by the high circulating hCG levels is unlikely because we have previously shown that Lhcg knockout mice carrying orthotopic transplants of WT ovary can become pregnant, and their mammary glands undergo normal pregnancy-associated lobuloalveolar maturation to enable lactation (32).

In humans, the role of PRL in the development of breast cancer still remains controversial. Some authors reported associations between PRL or hPRLR single-nucleotide polymorphisms and the risk of breast cancer (15, 25, 43), but no functional analysis has been undertaken to provide causality and mechanistic insight into these associations. This question prompts further studies on the involvement of the hPRLR in breast disease pathogenesis. Wnt signaling in the mammary gland was dysregulated in the hCGβ+ mice, as previously described (24), resulting in β-catenin-stabilizing mammary tumorigenesis. This response was not observed in the absence of PRL signaling, suggesting that disturbed Wnt signaling is dependent of PRL action, either directly or via LH/hCG stimulation of ovarian steroidogenesis. Hence, this mouse model provides novel information about hCG action and hormonally induced tumorigenesis of the breast.

We have previously demonstrated an amplifying effect of progesterone on the growth of estrogen-induced pituitary tumors in the hCGβ+ mice (1, 2). Also, the PRLR−/− mice exhibit pituitary hyperplasia and adenoma formation (40). In these mice, two factors contribute to the release of the lactotroph from its usual secretory and proliferative controls: a decrease in the normally inhibitory dopaminergic control and a second, direct effect at the level of the pituitary that is most consistent with an antiproliferative action of PRL on lactotrophs (40). Knowing that PRL is a stimulator of progesterone secretion, we observed pituitary adenomas in PRLR−/− and hCGβ+/PRLR−/− mice. However, no additive effect was shown in the hCGβ+ mice, suggesting that the deletion of PRLR is a stronger stimulus of lactotroph growth than elevated progesterone level.

Finally, we assessed adrenal gland morphology, since chronically elevated serum LH, augmented by enhanced PRL production (22), has been shown to induce functional LHR expression in the adrenal cortex of certain strains of mice, and that chronically elevated LH/hCG levels might stimulate growth and steroidogenesis of the adrenal cortex (7). Indeed, postgonadectomy tumorigenesis is thought to represent metaplasia of the subcapsular adrenocortical stem/progenitor cells between the zona glomerulosa and fasciculate that, under the influence of LH, transform into cells resembling gonadal stroma (6, 8, 27). Histological analysis of the adrenal glands showed no differences between genotypes, and absence of tumor development. This is in accordance with our previous work, which has demonstrated that adrenals of FVB/N mice appear resistant to postgonadectomy alterations (8).

The fertility parameters and eventual tumor formation of male mice was not evaluated since PRLR−/− and and hCGβ+ males (10, 39) were both described as fertile, and that the exposure of male mice to chronically elevated levels of hCG was not sufficient to promote testicular tumor formation.

In conclusion, persistently high LH/hCG stimulation is unable to compensate for missing PRL signaling in the maintenance of mouse luteal function, but the absence of PRLR expression prevents mammary gland tumorigenesis induced by
hCG overexpression. PRL action thus appears to be a major permissive regulator of LH action directly in the ovary and, indirectly, through ovarian steroidogenesis, in the mammary gland. This work paves the way for future studies on the molecular pathways involved in hormone-dependent tumorigenesis of the mammary and pituitary glands.

GRANTS

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DISCLOSURES

No conflicts of interest, financial or otherwise, are declared by the authors.

AUTHOR CONTRIBUTIONS


REFERENCES


