Mice lacking progesterone receptor exhibit pleiotropic reproductive abnormalities

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Although progesterone has been recognized as essential for the establishment and maintenance of pregnancy, this steroid hormone has been recently implicated to have a functional role in a number of other reproductive events. The physiological effects of progesterone are mediated by the progesterone receptor (PR), a member of the nuclear receptor superfamily of transcription factors. In most cases the PR is induced by estrogen, implying that many of the in vivo effects attributed to progesterone could also be the result of concomitantly administered estrogen. Therefore, to clearly define those physiological events that are specifically attributable to progesterone in vivo, we have generated a mouse model carrying a null mutation of the PR gene using embryonic stem cell/gene targeting techniques. Male and female embryos homozygous for the PR mutation developed normally to adulthood. However, the adult female PR mutant displayed significant defects in all reproductive tissues. These included an inability to ovulate, uterine hyperplasia and inflammation, severely limited mammary gland development, and an inability to exhibit sexual behavior. Collectively, these results provide direct support for progesterone's role as a pleiotropic coordinator of diverse reproductive events that together ensure species survival.

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Progesterone has been regarded traditionally as the mammalian "pregnancy hormone" (Baulieu 1989; Clarke and Sutherland 1990). Central to progesterone's role in early pregnancy is the ability of this ovarian steroid to coordinate a complex series of interactive steps that begin with the synchronized development of both the preimplantation embryo and the maternal uterus and end with embryo implantation. Although most of the downstream molecular and cellular mechanisms by which progesterone exerts these effects are unclear, the physiological effects of progesterone are known to be mediated initially by a specific intracellular protein termed the progesterone receptor (PR). In most tissues examined, the PR is induced by estrogen, via the estrogen receptor [ER], thereby implying that many of the observed reproductive physiological responses attributed to PR could conceivably be attributable to the combined effects of progesterone and estrogen. One of the current challenges in reproductive endocrinology is to define and characterize those physiological responses that are specifically the result of either progesterone or estrogen in vivo.

The PR has been shown to be composed of two naturally occurring ligand-binding forms, termed PR\(_A\) and PR\(_B\) (Schrader and O'Malley 1972), that arise from the same gene (Conneely et al. 1987; Kastner et al. 1990). The physiological significance of two receptor isoforms for the progesterone ligand is as yet unknown and is currently under intense investigation. The molecular cloning of the avian PR (Conneely et al. 1986) revealed that this receptor was a member of a nuclear receptor superfamily of transcription factors (Evans 1988; Tsai and O'Malley 1994) that includes receptors for a number of potent effector molecules for steroids, thyroids, retinoids, and vitamin D. The PR, as with all members of this superfamily of genes, is defined by a common structural motif that is organized into defined domains in terms of structure and function (Carson-Jurica et al. 1990b). The most conserved region, the DNA-binding domain (DBD), located centrally in the molecule, is composed of two type II zinc fingers which facilitate DNA binding to response elements (Umesono and Evans 1989). The moderately conserved ligand binding domain (LBD) is located in the carboxy-terminal half of the receptor. In addition to its ligand-binding functions, this domain also contains sequences required for receptor dimerization (Fawell et al. 1990) and target gene trans-activation (Dobson et al. 1989; Gronemeyer 1991). The short variable hinge region is located between the LBD
and the DBD and contains additional trans-activation functions together with sequences required for stabilization of interaction of inactive receptors with heat shock proteins (Dobson et al. 1989; Carson-Jurica et al. 1990a). The amino-terminal region of the PR, which is the most hypervariable region in terms of both size and sequence among members of this superfamily, contains trans-activation functions that modulate both the level and promoter specificity of target gene activation (Tora et al. 1988; Sartorius et al. 1994).

Apart from the established role of progesterone in early gestation, a number of studies have implicated a physiological function for progesterone in reproductive events other than in the establishment and maintenance of pregnancy. A variety of experimental approaches, including PR localization studies and the use of antiprogestins and progesterone replacement studies involving ovariectomized animals, have suggested possible developmental and functional roles for progesterone and its receptor in such reproductive systems as the ovary, mammary gland, and brain. However, because of the close temporal and spatial overlap in functional activities attributed to estrogen and progesterone, it has been difficult to directly interpret progesterone’s influence in many of these physiological systems without also considering the extent of estrogen’s role.

To circumvent this problem and to directly study PR function in an in vivo context, we have generated a novel mutant mouse strain carrying a germ-line mutation of the PR locus. Although male and female mice deficient in PR underwent apparently normal embryogenesis and developed to the adult state, mice homozygous for this mutation exhibited extensive functional abnormalities in a number of reproductive tissues that included the uterus, ovary, mammary gland, and brain. On the basis of the established role of PR in uterine development an infertility phenotype was expected in the female mouse; however, the number of reproductive systems that were affected by specifically removing PR function was surprising.

These results suggest that during the evolution of the mammalian reproductive system, progesterone’s role has developed as an essential and pleiotypic coordinator of diverse physiological events that ensure the survival of the species. Although progesterone has classically been regarded as the pregnancy hormone, the observations herein suggest that it more accurately may be termed the steroid hormone of reproduction.

Results

Targeted disruption of the mouse PR gene

Using the positive/negative selection approach to enrich for gene-targeted events in AB-1 embryonic stem (ES) cells (Mansour et al. 1988), the overall Southern strategy that was used to identify a mPR gene targeted event is shown in Figure 1A. The gene targeting vector, RV7, was designed to insert the neomycin resistance gene (neo'), PGK-neobpA (see Materials and methods), into the first exon of the mPR gene and downstream from the initiating codons ATGA and ATGB that encode the A and B forms of the PR. This insertion site was chosen to disrupt effectively the transcription of both forms of the PR.

In the case of the normal mPR allele, ES cell genomic DNA digested with HindIII and probed with a 0.5-kb SacI–EcoRI genomic fragment results in a hybridizing band of 5 kb (Fig. 1A). This probe is located just 5’ to the region of homology contained in RV7. In the case of a targeted event at the PR locus, the presence of the neo' cassette in exon 1 introduces an extra HindIII site that results in a shorter hybridizing band of 3.5 kb. A targeted ES cell clone will yield two hybridizing bands, representing the normal intact PR allele (5 kb) and the targeted disruption of the second PR allele (3.5 kb). Using this screening strategy, a targeting frequency of 8% was achieved at the PR locus. To ensure that potential positive clones did not contain 3’ rearrangements, genomic DNA was digested with BamHI and hybridized with a 0.5-kb Neol–Neol 3’ probe (Fig. 1A; data not shown). In addition, to establish the integration of one targeting vector, Southern blots were stripped and hybridized with a neo'-specific probe (data not shown).

To generate chimeric mice, ES cells carrying the targeted disruption of the mPR gene were microinjected into 3.5-day-old C57BL/6 blastocysts that were subsequently transferred to C57BL/6 pseudopregnant recipients. Two male chimeras transferred the PR mutation to the next generation (data not shown). Heterozygous mice were crossed to obtain mice homozygous for the mPR-targeted disruption. Figure 1B shows a typical Southern blot result of a litter derived from such a heterozygote mating. To date, 29 litters derived from heterozygote crosses have generated 204 offspring of which 52 are wild type, 107 are heterozygote, and 45 are homozygote for the PR mutation (26%:52%:22%). Both male and female were equally represented in the litters, and all animals developed to adulthood.

To confirm the loss of PR functional activity in mice that were homozygous for the PR-targeted disruption, uterine cytosolic PR was measured in age-matched ovariectomized wild-type, heterozygous and homozygous female mice (Fig. 1C). All mice were administered 10 μg of estradiol 72 hr prior to assay to further induce PR levels. Using the progestin agonist R5020, both wild-type and heterozygous animals revealed saturable binding of R5020 to endogenous PRs. Interestingly, heterozygotes contained approximately half the number of PRs yet were fertile. In contrast, R5020-binding sites were not detected in the uterus of homozygote animals (Fig. 1C). These results confirmed previous Northern blot analyses (data not shown) indicating lack of expression of functional PR in the homozygote mutant mouse.

Abnormal ovarian function

Adult homozygous mice of both sexes appeared healthy and developed normal external genitalia. Gross anatomical examination did not reveal obvious differences in
organ morphology between the homozygotes and their wild-type and heterozygote littermates. As might be expected, female homozygotes proved to be infertile in crosses with wild-type male mice. In contrast, male homozygotes were found to be as fertile as their wild-type and heterozygous male siblings.

As a first approach to determining causes of the infertility in female homozygotes, ovarian function was evaluated. Initial histological studies on ovaries isolated from hormonally untreated wild-type and homozygous mice did not reveal obvious morphological differences [data not shown]. We next examined the response of the homozygous ovary to the exogenous gonadotropins, pregnant mare serum gonadotropin (PMSG), and human chorionic gonadotropin (hCG) [see Materials and methods]. Using this hormonal treatment, wild-type mice were able to undergo superovulation and produce the expected large numbers of oocytes [see Table 1]. Surprisingly, oocytes were never detected in the oviduct or upper uterine horn of similarly treated, age-matched, homozygous females. Based on these data, histological analysis was performed on the ovaries of these animals.

Although the size, weight, and overall external appearance of ovaries isolated from homozygous females did not differ significantly from wild-type mice, histological studies revealed striking developmental defects in response to this hormonal treatment in the homozygous ovary. Figure 2A shows a representative view of an ovarian section stained with hematoxylin and eosin derived from a wild-type female 24 hr after hCG administration. The presence of numerous corpora lutea (CL) and the absence of mature preovulatory follicles is characteristic of an ovary that has recently undergone ovulation. In contrast, ovaries isolated from homozygote females (Fig. 2B) revealed an unexpected presence and an unusual number of mature “preovulatory” follicles [unruptured follicle (UF)] as well as a marked absence of functional CL. In each case, anovulatory follicles underwent cumulus expansion, which is the last step that can be identified at the histological level before a follicle ruptures. Considering that in the mouse, ovulation usually takes place ~12 hr after the administration of hCG, the presence of an intact mature oocyte that has not undergone

Table 1. Oocytes and embryos produced following superovulation

| Group        | Oocytes | One-cell stage | Two-cell stage | No. |
|--------------|---------|----------------|----------------|-----|
| Wild type    | 11 ± 3  | 13 ± 2         | 1              | 6   |
| Heterozygote | 11 ± 2  | 15 ± 2         | 1              | 6   |
| Homozygote   | 0       | 0              | 0              | 6   |

Ovarian function was assayed by determining the ovary’s response to superovulatory doses of the gonadotropins PMSG and hCG [see Materials and methods]. Following hCG treatment, mice were put with sexually experienced wild-type males overnight. Oocytes as well as one- and two-cell stage embryos were flushed from both oviducts of each animal 24 hr after hCG administration and examined and counted using a dissecting microscope. The data are means ± s.d.
A progesterone receptor-deficient mouse model

Figure 2. Ovarian response to PMSG and hCG. (A) A transverse section (4 µm) of a typical ovary isolated from a 6-week-old wild-type mouse (+/ +), treated previously with PMSG and hCG [as described in Materials and methods]. Note the presence of numerous CL. Scale bar 100 µm. (B) A representative cross section of an ovary isolated from an age-matched homozygote female (−/−) that was hormonally treated exactly the same as for the wild type. The unusual presence of a number of UFs is indicated. Scale bar, 100 µm. (C) High magnification (20×) of a CL present in the wild-type ovary. Note the presence of the characteristic hypertrophied luteal cells (LC). Scale bar, 50 µm. (D) High magnification (20×) of a UF present within the PR homozygote ovary. An intact oocyte (O) with a zona pellucida is shown in addition to granulosa cells (GC) that have undergone cumulus expansion. Interestingly, the granulosa cells within this follicle do not show signs of luteinization, cf. their cellular morphology to C. Scale bar, 50 µm. This is a representative result of six animals that were studied per genetic group.

Further cell division or necrosis was unexpected. Furthermore, the granulosa cells located either in the vicinity of the oocyte or positioned near the intact basal lamina do not show signs of luteinization (Fig. 2, cf. C and D).

Uterine response to progesterone and estrogen

Preliminary histological analysis of uteri isolated from intact untreated homozygote females demonstrated the presence of a normal myometrial and endometrial compartment [data not shown]. To determine the response of the homozygote uterus to the effects of estrogen and progesterone, both age-matched, six-week-old ovariectomized virgin homozygote and wild-type females were treated hormonally for 3 weeks with a daily dose of 1 mg of estradiol and 1 µg of progesterone. Following ovariectomy, uteris isolated from hormonally untreated wild-type (Fig. 3A) and homozygote (Fig. 3B) animals were atrophic and generally indistinguishable morphologically. Administration of estrogen and progesterone induced profound gross anatomical and histological changes in the homozygous female. In the case of the wild-type uterus (Fig. 3C), hormonal treatment resulted in both a moderately enlarged and highly developed uterus, a characteristic phenotype that develops following estrogen and progesterone treatment. In the homozygous animal, estrogen and progesterone treatment resulted in an abnormally enlarged fluid-filled uterus (Fig. 3D). Cytospin preparations of intraluminal uterine fluid taken from homozygotes demonstrated a moderate infiltration of polymorphonuclear (PMN) leukocytes, indicative of an acute inflammatory response. Histologic evaluation of transverse sections of the uterine horn from these animals (Fig. 3F) revealed thickening of the uterine wall by extracellular edema, acute inflammatory cells, and proliferation of mucosal and glandular epithelia. Inflammatory changes were limited to the mucosa, submucosa, and stromal components of the endometrium. PMN leukocytic infiltration was present in both endometrial stroma and the overlying luminal epithelium. This epithelium appeared hyperplastic and disorganized compared to wild-type controls. Endometrial glands were enlarged greatly and lined by hypertrophied epithelia. The above histological changes corresponded to the previously reported phenotype observed in the uterus of the wild-type mouse treated with estrogen alone (Martin et al. 1973). In contrast, uteri from wild-type littermates (Fig. 3E) were lined by normal low cuboidal to low columnar epithelium, small endometrial glands, and a condensed endometrial stroma with no evidence of inflammation.

Uterine response to decidual stimulation

To determine whether the abnormal sensitivity of the homozygote uterus to estrogen and progesterone was also accompanied by an inability to undergo deciduization, the uterine response to an artificial decidual stimulus was measured in age-matched wild-type and mutant mice [see Materials and methods]. To elicit the decidual response artificially, an estrogen and progesterone treatment was used as described previously (Ledford et al. 1976), followed by the stimulation of the left uterine horn of each animal by mechanical traumatization [see Materials and methods]. The right uterine horn was not stimulated and served as the control.

In the case of the wild-type mouse, a decidual response, exhibited as a dramatic increase in uterine horn...
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The conditions used in this study produced a decidual response in the wild-type uterus that resulted in a >20-fold increase in the weight of the stimulated horn as compared to the PR mutant.

Mammary gland development

Initially, mammary gland whole mounts were performed on the inguinal mammary gland (number 4 gland) iso-

Figure 3. Uterine response to estrogen and progesterone treatment. Comparative gross anatomy of uteri isolated from hormonally untreated ovariectomized wild type (+/+) and the PR homozygote (-/-) did not reveal a significant difference in morphology. All mice were 6 weeks old. In situ gross anatomy of wild-type (C) and PR mutant (D) uteri, following estrogen and progesterone treatment (see Materials and methods), is indicated by arrows. Note the marked enlarged fluid filled uterus in the PR homozygote (D). (E) Histological analysis of a representative cross section of the uterine wall of the hormonally treated wild-type mouse shows the presence of a normal uterine architecture, luminal epithelium (LE), glandular epithelium (GE), stromal cell layer (S), and myometrium (M). Scale bar, 50 μm. (F) Morphological analysis of a typical transverse section of the uterine wall of the PR mutant, treated with estrogen and progesterone, reveals an abnormal uterine structure. Note the hyperplastic luminal (LE) and hypertrophic glandular (GE) epithelia, loosely arranged stromal layer (S), and presence of polymorphonuclear leucocytes (indicated by arrowheads). Scale bar, 50 μm. This is a typical result from six animals per genetic group that were studied.

Wild type (+/+) and PR mutant (-/-). The data are represented as means ± S.E.M.

Figure 4. Uterine response to a decidual stimulus. (A) Gross morphology of the wild-type uterus following decidual stimulation (see Materials and methods). In contrast to the unstimulated right (R) uterine horn (control horn), note the dramatic increase in size and extensive vascularization of the left (L) uterine horn following mechanical stimulation. (B) Using the same experimental conditions as the wild type, the PR mutant uterus failed to exhibit the typical decidual response in the stimulated L uterine horn. (C) A histogram shows the ratios of averaged weights of stimulated over unstimulated (control) uterine horns from six wild-type (+/+) and six PR mutant animals (-/-). The data are represented as means ± S.E.M.
lated from intact, hormonally untreated virgin homozygotes (6 weeks old) and their wild-type littermates. Morphological analysis of these glands consistently showed no significant difference in ductal development between the homozygote and wild-type mouse virgin glands [data not shown]. To determine whether the PR mutation had an effect on the development of the mammary gland to the differentiated phenotype that occurs during early pregnancy [Imagawa et al. 1994], mammary gland whole mounts were taken from both estrogen and progesterone-treated and untreated ovariectomized mice as described above. In the case of the hormonally untreated ovariectomized homozygote and wild-type female, whole-mount studies revealed a common rudimentary ductal structure in both genetic groups [Fig. 5, cf. A and B]. However, following estrogen and progesterone treatment, hematoxylin staining of whole mounts derived from the wild-type mammary gland revealed a complex ductal arborization to the periphery of the fat pad [Fig. 5C]. In contrast to wild-type animals, hormonally treated homozygote females exhibited a more basic ductal structure with less extensive dichotomous and lateral side branching [Fig. 5D]. Confocal laser scanning microscopy confirmed the striking absence of lobuloalveolar development at the end of each duct in the PR mutant animal [Fig. 5, cf. E and F].

**Lordosis response**

To determine whether the PR mutation altered the sexual receptive response [lordosis] of homozygote females to the mounting attempts of sexual experienced wild-type males, the lordosis response was measured in age-matched (60-day-old) ovariectomized wild-type (+/+ ) and homozygous (−/−) females following estrogen and progesterone treatment. In the absence of progesterone, both wild-type and homozygous animals did not display a significant lordosis behavior 48 hr after estrogen priming [Fig. 6A]. However, 6 hr following progesterone treatment, a marked increase in the lordosis quotient (LQ) was observed in estrogen-primed wild-type females [Fig. 6B]. In contrast, in five separate experiments, significant progesterone-facilitated behavioral response was not observed in the estrogen-primed homozygote females [Fig. 6B].

**Discussion**

To gain further insight into the physiological functions of PR and to differentiate clearly between ER- and PR-mediated responses, we have generated a mouse model carrying a null mutation of the PR gene using homologous recombination in mouse ES cells. Although PR has been detected as early as the blastocyst stage of development [Hou and Gorski 1993], heterozygote matings generated PR mutant homozygote embryos at the normal Mendelian frequency with no deviation in the sex ratio, indicating that embryonic-derived PR is not required for embryonic survival or for prenatal development of the female reproductive system. As expected, adult female PR homozygotes were found to be infertile. The basis of this infertility could be found in the extensive developmental abnormalities of the ovary, uterus, and mammary gland as well as an inability to exhibit the sexual behavioral response, lordosis. Based on these observations, we conclude that the PR has evolved as a transcription factor with pleiotropic effects that allows it to coordinate most aspects of female reproduction.

**The PR is essential for ovulation**

Although the ovary has long been established as the primary endocrine tissue for progesterone biosynthesis and release, the involvement of progesterone as a local modulator of ovarian function has been a matter of controversy for many years [Yoshimura et al. 1987]. However, a growing number of studies have recently implicated an intraovarian role for progesterone in the ovulation process. For example, in rats, several inhibitors of proges-
have shown that ovulatory levels of LH, via a cAMP homozygous for this mutation. Morphological analysis of ovaries isolated from rat preovulatory follicles (Natraj and Richards 1993; Chandrasekher et al. 1994), the lack of bona fide CL and the absence of luteinized UFs in ovaries derived from the PR-deficient mouse following gonadotropin treatment provides strong in vivo evidence for an important functional role for PR in the luteinization process. Finally, it is clear from our in vivo studies on the ovary that progesterone and its receptor play a pivotal role in two distinct but coupled intraovarian pathways initiated by the LH surge, namely ovulation and luteinization.

Uterine developmental defects

In the uterus and throughout the female reproductive tract, specialized groups of cells undergo continuous synchronized waves of proliferation and differentiation in response to the cyclical rise and fall of ovarian estrogen and progesterone. Unlike the ovary, the expression of the PR seems to be primarily regulated in the uterus by estrogen (Jordan and Dix 1979). It has been known for some time that in response to a decidualization signal, estrogen and progesterone promote profound morphological changes in the growth and differentiation of the epithelial and stromal cell layers of the endometrial compartment (Clarke and Sutherland 1990). These studies also suggested that estrogen was the main proliferative stim-
ulus during this stage of uterine development whereas progesterone effects were more involved in differentiation. More recently, a mouse model carrying the mutation for the targeted disruption of the ER gene failed to exhibit increased uterine weight or hyperemia in response to estrogen stimulation, providing strong evidence for a proliferative role for estrogen in the uterus [Lubahn et al. 1993]. However, conclusions drawn from this data must take into account that in removing ER function, PR function is also ablated. Therefore the observed uterine phenotype in the ER defective mouse could be interpreted as attributable to the combined loss of ER and PR function. As a result, the generation of a new mouse strain that is defective in PR provides a necessary animal model to study not only progesterone’s role in the uterus but also to define the separate functions of estrogen and progesterone in vivo.

In contrast to the wild-type female, ovariectomized female PR mutant mice exhibited extensive gross and histological morphologic abnormalities of the uterus and lower reproductive tract when chronically treated with estrogen and progesterone. These abnormalities appeared to correlate to some degree to estrogen hyper-sensitivity previously observed in ovariectomized wild-type female mice when treated with estrogen for prolonged periods of time in the absence of progesterone [Martin et al. 1973; Quarmby and Korach 1984]. In the case of the PR mutant, the presence of extensive proliferation of the luminal and glandular epithelial cell layers of the endometrium confirms the studies described for the ER-defective mouse model [Lubahn et al. 1993] as well as providing additional in vivo support for a proliferative role for estrogen in the uterus.

More surprising was the presence of a strong local inflammatory reaction in the PR homozygote uterus in response to this hormonal regimen. This inflammatory response was characterized by a general disruption of the uterine architecture and a marked infiltration of PMNs in the edematous endometrial stroma and overlying mucosal epithelium. In addition, analysis of the uterine luminal fluid, contained in distended uteri of these animals [Fig. 3D], revealed the presence of a large number of PMNs. These observations provide compelling support for the proposal that progesterone can exhibit anti-inflammatory effects in this tissue. This proposal can be related to a number of recent reports in the literature. For example, the expression of interleukin 8 (IL-8), a powerful chemotactic cytokine for neutrophils [Matsushima et al. 1992] and T lymphocytes [Larsen et al. 1989], was shown to be inhibited by progesterone in cultured rabbit uterine cervical fibroblasts [Ito et al. 1994] as well as in human choriodecidual cells [Kelly et al. 1992]. In addition, the antibortefacient and antiprogestin RU486 has been shown to stimulate prostaglandin E (PGE) levels in the decidual and chorionic tissue in early pregnancy [Cheung et al. 1993], thereby providing indirect evidence for an inhibitory role for progesterone in prostaglandin synthesis. Furthermore, PGE has been shown to act in concert with IL-8 in triggering neutrophil invasion that subsequently results in the remodeling of the local connective tissue [Colditz 1990]. Although speculative, on the basis of these reports and our in vivo studies, we suggest that to maintain successfully the maternal tissue architecture during pregnancy, progesterone suppresses many local inflammatory responses initiated by prostaglandins and cytokines.

Finally, the abnormal sensitivity of the homozygote uterus to estrogen and progesterone treatment was also accompanied by the failure of this organ to respond to an artificial decidual stimulus, indicating that the endometrial cell layer of the homozygote uterus is refractory to the implantation signals of the preimplantation embryo. Current investigations are under way to examine gene expression under these conditions and to define more clearly the role of PR in the induction of the decidual response.

**Mammary gland lobuloalveolar development requires PR**

As with most female sex organs, the proliferation and differentiation of the mammary gland is dependent on a delicate balance between the actions of estrogen and progesterone. Until recently, it generally was considered that estrogen was the primary ovarian hormone involved in the proliferation of the normal mammary gland as well as being responsible for the onset and progression of mammary tumors, whereas progesterone would serve more of an antiproliferative role [Clarke and Sutherland 1990]. This concept, which was based mainly on analogy to the role of these hormones in uterine development, is currently under intense scrutiny [Horwitz 1992]. However, certain data have accumulated to implicate a mitogenic role rather than an antiproliferative role for progesterone in the development of the mammary gland. For example, previous studies have shown that although estradiol alone can induce DNA synthesis in mammary glands of ovariectomized mice [Bresciani 1968; Shyamala and Ferenczy 1984; Daniel et al. 1987], this mitogenic response can be augmented further by coadministration of progesterone [Bresciani 1968; Shyamala 1987]. In addition, studies have implied that progesterone may initiate proliferative responses in the normal mammary epithelium in virgin animals [Haslam 1988] as well as contributing to the growth and development of the lobular-alveolar system in mammary glands of pregnant animals [Imagawa et al. 1985; Haslam 1988].

Progesterone also has been implicated in the growth of carcinogen-induced [Welsh 1985] and transplantable rat mammary tumors [Robinson and Jordan 1987] as well as in spontaneous tumorigenesis of the murine mammary gland [Nagasawa et al. 1988]. As further support for the stimulatory effect of progestins on the mammary gland epithelium, antiprogestins were found to inhibit the growth effects of progesterone in certain experimental mammary tumors, suggesting that the proliferative actions of progesterone are mediated through the PR [Michna et al. 1989]. Therefore, the growth stimulatory effects of progesterone in normal mammary gland development and in the progression of mammary tumorgenesis...
esis may have far-reaching implications for the use of progestin agonists in contraception and postmenopausal hormonal replacement as well as for the therapeutic use of antiprogestins in the control of hormone-dependent breast cancers.

In the case of female mice lacking the PR, we observed less extensive ductal development in the mammary gland compared to the wild type as well as a complete absence of interductal lobular–alveolar structures despite treatment with pregnancy levels of estrogen and progesterone. The data provide unequivocal in vivo evidence that progesterone is involved in the proliferation of the mammary ductal epithelium and thus has an obligatory role in the establishment of the lobular–alveolar system that is required for lactation. The animal model described here should provide a unique opportunity to dissect the relative importance of estradiol and progesterone in normal and neoplastic mammary growth and also to access the therapeutic use of antiprogestins in diverse clinical settings.

**The PR is essential for the expression of lordosis**

We have shown that the estrogen-primed ovariectomized female mouse, homozygous for the PR mutation, was unable to exhibit a significant lordosis response in the presence of an experienced male when administered progesterone. This result confirms recent reports that describe the failure of progesterone to induce a sexual behavioral response in estrogen-primed ovariectomized rats that were previously administered antisense oligonucleotides to the PR intracerebroventricularly [Mani et al. 1994b] and into the ventromedial hypothalamus [Ogawa et al. 1994]. Together these results demonstrate that the PR, induced by estrogen in the region of the ventral medial nucleus of the hypothalamus [VMNH] and in the preoptic area, is essential for the expression of this behavioral response. Interestingly, the lack of estrogen-induced PR in the VMNH of the male brain is thought to be a primary reason for the inability of the male to elicit this behavior [Rainbow et al. 1982]. We speculate that the inability of the PR homozygote female to elicit the lordosis response may be associated with a defect in a “PR-induced” developmental event that may have occurred during the laying down of permanent neuronal networks in prenatal life. Alternatively or in addition, it is quite possible that the developmental defect occurs in ongoing transient and reversible organizational events (synaptic plasticity) that are known to occur in the VMNH of the adult [Frankfurt et al. 1990]. Finally, recent in vitro and in vivo studies have shown that the PR can be trans-activated in a ligand-independent manner by the neurotransmitter dopamine [Power et al. 1991; Mani et al. 1994a]. The fact that dopamine also can exert a stimulatory response on this sexual behavior in estrogen-primed rats implies an important physiological role for the PR as a point of convergence for neurotransmitter and progesterone-signaling pathways and further emphasizes the importance that evolution has placed on this receptor as a central coordinator of key regulatory pathways of reproduction.

In summary, the PR-defective mouse model has been useful in defining the essential role that evolution has assigned to progesterone in the development and function of most aspects of the female reproductive system. Collectively, these studies emphasize the importance of the PR as a pleiotropic coordinator of many different reproductive events that function together to establish in turn, fertilization, pregnancy, and lactation. A most important advantage in studying this mouse model is that it provides an essential tool to define clearly those physiological events that are specifically attributable to either estrogen or progesterone. For future studies, this mouse model will provide an ideal system in which to identify potential novel progesterone target genes in the ovary, uterus, and mammary gland through the use of differential cDNA cloning strategies.

**Materials and methods**

**Design and construction of the gene targeting vector RV7**

The mPR genomic DNA fragment (7 kb) that comprised the gene-targeting vector RV7 was isolated from a mouse 129Sv × Dash II genomic library [Stratagene, La Jolla, CA] using the full-length mPR cDNA as a probe [Schott et al. 1991]. This mPR genomic fragment consisted of the first two exons of the mPR gene and the first intron as well as part of the second intron. The remainder of the genomic DNA consisted of a region that was located 5’ to exon 1. Exon 1 encodes the nonconserved amino-terminal domain of the receptor that contains the two initiating methionine residues, ATG<sub>B</sub> and ATG<sub>A</sub>, that define the amino termini of two forms of the receptor, namely B and A, respectively [Connelly et al. 1987]. Exon 2 encodes the first zinc finger of the conserved DNA-binding domain of the receptor [Huckaby et al. 1987]. The neo<sup>r</sup> gene PGKNEObpA [Soriano et al. 1991] was inserted into a unique Xhol restriction site in exon 1 that was located 225 and 721 bp downstream from the initiator codons ATG<sub>A</sub> and ATG<sub>B</sub>, respectively, in the mPR gene. This site was chosen as the location to insert the neo<sup>r</sup> gene because this insertion would effectively disrupt exon 1 by prematurely terminating transcripts initiated from codons ATG<sub>A</sub> and ATG<sub>B</sub>. The insertion of the neo<sup>r</sup> gene into exon 1 divides the 7-kb mPR genomic fragment into 5’ and 3’ arms of mPR homology that are 1.5 and 5.5 kb in size, respectively. The herpes simplex virus thymidine kinase (HSV–TK) gene [Mansour et al. 1988] was attached 5’ to exon 1 and inserted with a transcriptional orientation opposite to both the neo<sup>r</sup> and mPR genes. The cloning plasmid used in this vector construction was pSP72 [Promega Biotec, Madison, WI]. Finally, prior to electroporation into ES cells, RV7 was linearized at the 3’ end of the long arm of homology by the restriction enzyme Asp718 that was introduced into a naturally occurring Ncol site.

To identify the PR mutation in ES cells and mice, Southern blot analysis was performed on genomic DNA isolated from actively growing ES cells and mouse tail biopsies, respectively. In the case of ES cells, cell lysis, DNA precipitation, and restriction enzyme digestion were performed in 96-well microtiter plates as described previously [Ramirez-Solis et al. 1992]. For mouse tail biopsies, samples were extracted with phenol chloroform following an overnight proteinase K digestion step, and the resulting DNA was subsequently precipitated with ethanol. DNA samples were digested with HindIII overnight, resolved by
Introduction of RV7 into mouse ES cells

The general procedures for the culturing and manipulation of ES cells prior to and after the electroproporation step were followed as described (Robertson 1987). Briefly, 10^7 ES cells were electroporated with 25 μg of linearized RV7 in 0.9 ml of PBS at 230 V and 500 μF with a Bio-Rad Gene Pulsar. Electroporations were performed routinely using the actively growing ES cell line AB-1 (McMahon and Bradley 1990) at passage 13. Following electroporation, ES cells were plated (10^7 cells/100-mm plate) on a monolayer of the mouse embryonic fibroblast cell line SNL76/7 (McMahon and Bradley 1990). ES cells were cultured in the presence of G418 (350 μg/ml) and FIAU (0.2 μM) 24 hr postelectroporation. After ~10 days of G418 and FIAU drug selection, drug-resistant ES cell colonies were picked and expanded in 96-well SNL76/7 feeder plates (master plates). A duplicate gelatinized 96-well plate (no feeder layer) of each master plate was also prepared to identify targeted events by Southern analysis. The master plates containing the ES cell clones for blastocyst microinjection were frozen at -70°C until identification of those ES cells scoring positive for the targeted event.

Generation of chimeric mice and Germ-line transmission of the PR mutation

Six targeted ES cell clones were tested for germ-line transmission of the PR mutation. ES cells (12–15 cells) were microinjected into the blastocoeel of 3.5-day-old blastocyst stage embryos derived from C57BL/6 females. Embryos were transferred unilaterally into the uterine horn (six to seven embryos per horn) of pseudopregnant F1 [CBA×C57BL/6] foster mothers. Approximately 10 days after birth, the sex of the offspring was determined and the extent of agouti coat color was evaluated. Male chimeras with 60%–100% agouti coat color were backcrossed to C57BL/6 females, and germ-line transmission was determined by the presence of agouti offspring. The PR mutation was established also in a 129SvEv genetic background by crossing male chimeras with 129SvEv females.

Histological studies

Prior to postmortem examination, all animals were euthanized by sedation with methoxyflurane (Metafane, Pitman Moore, Mundelein, IL) followed by cervical dislocation. A gross necropsy was performed on each animal and the following tissues were taken for histopathologic assessment: heart, lungs, trachea, kidneys, urinary bladder, liver, spleen, brain, adrenal gland, thyroid gland, parathyroid gland, pancreas, salivary glands, esophagus, stomach, small and large intestine, cervical and mesenteric lymph nodes, ovaries, uterus, vagina, testes, epididymis, seminal vesicles, and preputial glands. Tissues were fixed in 10% neutral buffered formalin that was supplemented with zinc chloride (Anatech, Battle Creek, MI). Following appropriate fixation time, tissues were trimmed, dehydrated through ascending grades of ethyl alcohol, cleared in xylene, and infiltrated with paraffin wax. Processing was performed on a Miles Tissue-Tek VIP 3000 automated tissue processor. Tissues were embedded in paraffin and 4-μm sections were prepared using a standard rotary microtome (Leitz 1512). The sections were heat-dried and, following decration, stained with hematoxylin and eosin (Richard-Allan, Richland, MI) using the Code-On automatic stainer. The sections were dehydrated through ascending grades of alcohol, rinsed in xylene, and mounted using a synthetic mounting medium.

The hematoxylin staining of mammary gland whole mounts was performed as described previously [Medina 1973]. Briefly, both ingunal glands were dissected from euthanized mice and fixed in 10% formalin for a minimum of 24 hr. Following fixation, lipid was removed in two changes of acetone for 24 hr. After a 1-hr wash in each of 100% and 95% ethanol, the glands were stained in hematoxylin for 24 hr. The glands were dehydrated for at least 1 hr in tap water and subsequently dehydrated by washing in 70% and 95% ethanol, followed by absolute ethanol and xylene. Finally, processed mammary glands were stored permanently in methylsalicylate.

Decidual response

The decidual response was measured in wild-type and homozygote female mice. Two weeks following ovariec-tomy, mice (6 weeks old) were administered a hormone regimen as described previously [Ledford et al. 1976]. Briefly, mice were primed with three daily subcutaneous injections (days 1, 2, and 3) of 100 ng of 17 β-estradiol in 0.1 ml of sesame oil. Mice did not receive injections on days 4 and 5. On days 6, 7, and 8, the animals were sensitized with daily injections of 1 μg of progesterone and 6.7 ng of 17 β-estradiol in 0.1 ml of sesame oil. Six hours after the third Pe injection, the left horn of the uterus was dramatically stimulated by insertion of a burred needle into the horn just proximal to the cervix and longitudinally scratching the entire length of the uterine horn along the antimesometrial side. The right horn was not stimulated. Daily injections of Pe were administered until day 14 (6 days poststimulation) when the animals were sacrificed. The uteri were extracted and matched horns were cut to equal lengths and weighed. Six mice from each genetic group were examined. Following uterine weight measurements, tissue samples were processed further for histological analysis.

Immunofluorescence and confocal laser scanning microscopy

Following dissection, the inguinal mammary gland was treated with 0.5% Triton X-100 in PBS for 5 min before fixation in 70% ethanol/PBS for 3 hr. Approximately 2×2-mm sections of the gland were cut and rehydrated in PBS for 30 min. Thin slices from each section were cut using a scalpel and teased gently to expose duct and end-bud structure. Each tissue slice was incubated overnight with a rabbit anti-keratin-14 polyclonal antibody [a gift from Dr. Dennis R. Roop, Baylor College of Medicine, Houston, TX] at a dilution of 1:500 in 1% BSA in PBS. Following primary antibody incubation, tissue slices were washed in PBS before a 1 hr incubation with a fluorescein isothiocyanate (FITC)-tagged anti-rabbit secondary antibody [Pierce Chemicals, Rockford, IL]. Finally, tissue slices were washed in PBS and mounted on glass slides for confocal microscopy.

Optical sectioning of mammary gland tissue sections was performed using the multiprobe 2001 confocal laser scanning microscope [Molecular Dynamics]. Optical sections were obtained serially and three dimensional images were reconstructed using Image Space software [Molecular Dynamics].

Behavioral testing

Sexual receptivity experiments were performed on wild-type and homozygote female mice. Two weeks following ovariecto-
my, mice (60 days old) were administered 1 μg of estradiol benzoate subcutaneously in 50 μl of sesame oil followed 48 hr later by 100 μg of progesterone in the same volume of vehicle. Six mice of each genetic group were tested separately but simultaneously on a weekly basis for a total period of 5 weeks for sexual receptivity after the administration of estradiol and six hr after the administration of progesterone. The evaluation of sexual behavior commenced with the introduction of each female into the home cage of a sexually active (stimulus) male. Each female was tested for 10 mounts by the male or for a maximum of 30 min. Only mounts in which the male showed pelvic thrusting were scored. During the testing period, if a male did not mount, the female was placed with a different male. The female's response to a mount was categorized as either a lordosis response, consisting of a full arching of the back or no lordosis, consisting of no response to mounts with pelvic thrusting by the male. This receptive behavior was quantified and expressed as the LQ, which is defined as the number of lordosis responses of the female divided by the number of mounts by the male and multiplied by 100 (Mani et al. 1994b). Finally, all female mice were coded so that observers were blind with respect to the genotype of the animal under study.

Animals

For oocyte and embryo collection experiments, 6-week-old virgin mice were superovulated by an intraperitoneal injection of 5 IU of PMSG (Diosynth, Chicago, IL) and induced to ovulate 48 hr later with 5 IU of hCG (Pregnyl, Organon, Inc., West Orange, NJ). Immediately following hCG treatment, females were housed overnight with B6C3F1 fertile males (Harlan Sprague-Dawley, Indianapolis, IN). The presence of a copulation plug the following morning indicated day 1 of pregnancy. In these experiments, oocytes and embryos were collected from oviducts 24 hr following hCG administration. For ovarian histology, ovaries were routinely excised either 13, 24, or 48 hr following hCG treatment, the ovarian histology described herein represents the 24 hr time point.

To evaluate the response of the uterus and mammary gland to estrogen and progesterone stimulation, ovariectomized virgin mice (6 weeks old) were injected subcutaneously with 1 μg of 17β estradiol and 1 mg of progesterone (Sigma Chemical Co., St Louis, MO) dissolved in 50 μl of sesame oil, daily for 3 weeks.

Progestosterone receptor assays

Uterine tissue was excised immediately after euthanization of ovariectomized mice that were administered 10 μg of estradiol benzoate 72 hr previously. Tissue was homogenized in four volumes of 10 mM Tris-HCl, 1.5 mM Na2 EDTA, 10% glycerol, and 12 mM monothioglycerol (pH 7.4) using a Polytron homogenizer (Brinkman Instruments, Westbury, NY). Tissue homogenates were centrifuged at 100,000 rpm for 10 min in an ultracentrifuge using a TLA 100.3 rotor. In decreasing amounts (1–30 nm) of [3H]labeled R5020 (SA 86.90 Ci/m mole, New England Nuclear Corp., Boston, MA) were incubated with 100-μl aliquots of the cytosol supernatant either in the presence or absence of 100 nm unlabeled R5020. After an overnight incubation at 4°C, bound and free [3H]-labeled R5020 were separated by dextran-coated charcoal treatment. Samples were counted subsequently in a Beckman LS8000 liquid scintillation counter. The concentrations of cytosol protein were assayed by the method of Bradford (1976). All sample manipulations were carried out at 4°C.

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