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Involvement of Estrogen Receptor β in the Induction of Polyovular Follicles in Mouse Ovaries Exposed Neonatally to Diethylstilbestrol

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Natural and synthetic estrogens, including diethylstilbestrol (DES), given during the critical period of newborn life induce abnormalities in ovaries of mice. Induction of polyovular follicles (PFs) containing two or more oocytes in a follicle is one example. In this study, the involvement of estrogen receptor subtypes ER α and ER β in induction of PFs by neonatal treatment with DES was analyzed by using ER α knockout (α ERKO) and ER β knockout (β ERKO) mice. Ovaries of mice injected with 3 μ g DES for 5 days from the day of birth were examined histologically from 10 to 60 days of age, and the expression of genes involved in folliculogenesis was analyzed by real-time quantitative PCR. The PF incidence (percent of PFs per 100 follicles greater than 50 μ m in diameter) in the ovary of α ERKO mice treated with DES was not different from that in the DES-treated wild-type mice. However, neonatal DES treatment did not increase the PF incidence in β ERKO mice, suggesting that PFs were induced by DES through ER β but not ER α . The expression of bone morphogenetic protein 15, growth differentiation factor 9, inhibin- α , Müllerian inhibiting substance, and other genes in the ovaries of DES-treated β ERKO mice was not different from that in the ovaries of DES-treated wild-type mice. These results indicate that ER β but not ER α is essential for DES to induce PFs in mice.

Key words: ovary, estrogen receptor, ovarian follicle, estrogen, mouse

INTRODUCTION

Newborn female mice have been used as a rodent model to understand the mechanisms of estrogen action responsible for the human diethylstilbestrol (DES) syndrome in which pregnant women during the period from the 1940s to 1970s were prescribed DES to prevent miscarriage. Daughters born from these DES-exposed mothers developed vaginal cancer and other reproductive abnormalities (Herbst and Bern, 1981). DES use in women was stopped in 1971. Neonatal DES treatment induces various abnormalities in female mouse reproductive organs, including persistent cornification of the vaginal epithelium (Takasugi, 1963, 1976) and tumorigenesis in the mammary gland (Lopez et al., 1988). Neonatal treatment of mice with DES induces several histological abnormalities in the ovary, including polyovular follicles (PFs) having two or more oocytes in a

follicle, ovarian cysts, hemorrhagic cysts, hypertrophy of the interstitial tissue, and the absence of corpora lutea. The lack of corpora lutea in the ovaries of neonatally DES-treated mice was caused by lack of a surge of luteinizing hormone (Hayashi and Aihara, 1989; Hayashi et al., 1991; Döhler and Jarzab, 1992).

Estrogen acts through binding to two cognate nuclear receptors, estrogen receptor (ER) α and ER β , in female reproductive organs. DES binds to both ER α and ER β with high affinity compared to 17 β -estradiol (E₂) (Kuiper et al., 1997). The uterus, mammary gland, placenta, liver, central nervous system, cardiovascular system, and bone have high ER α content. ER β is detected in the prostate, testis, ovary, pineal gland, thyroid gland, and skin, and is especially highly expressed in the prostatic epithelium, urogenital tract, ovarian follicles, lung, and muscle (Weihua et al., 2003). In mouse ovaries, ER α mRNA and protein are localized in interstitial and thecal cells, and ER β expression is detected in the nuclei of granulosa cells (Jefferson et al., 2002; Hishikawa et al., 2003). In studies with ER α knockout (α ERKO) and ER β knockout (β ERKO) mice, and the use of ER selective ligands, DES was shown to act through ER α

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but not ER β in the induction of squamous epithelial metaplasia in the uterus, proliferative lesions of the oviduct, persistent cornification of the vaginal epithelium, and hypertrophy of the ovarian interstitial tissue (Couse and Korach, 2004; Nakamura et al., 2008).

PFs are found in the ovary of intact mice, rats, hamsters, dogs, and other species of mammals, including humans (Iguchi, 1992; Iguchi and Sato, 2000). The PF incidence is different among strains in intact mice (Kent, 1960), and it is increased markedly by neonatal DES treatment (Forsberg, 1985; Iguchi, 1985; Iguchi et al., 1987). In C57BL/Tw mice, PFs are induced with high incidence by neonatal treatment with estrogens, progestins, and aromatizable androgen, testosterone (T), but not by the non-aromatizable androgens 5 α -dihydrotestosterone (5 α -DHT) or 5 β -DHT (Iguchi et al., 1986). In addition, simultaneous administration of aromatase inhibitor (4-hydroxy-4-androstene-3, 17-dione) and T does not induce PFs (Iguchi et al., 1988). The evidence suggests that T enhances PF induction as a result of its conversion to estrogen (Ryan et al., 1972). DES acts directly on the neonatal ovary to induce PFs both in vivo and in vitro (Iguchi et al., 1990); however, it has been unclear which ER subtype is involved in the PF induction by neonatal DES treatment.

Even without DES treatment, PFs are found in the ovaries of double mutant bone morphogenetic protein 15 (Bmp-15) $^{-/-}$ and growth differentiation factor 9 (Gdf-9) $^{+/+}$ mice (Yan et al., 2001). Mice overexpressing inhibin- α gene also show a high incidence of PFs (McMulle et al., 2001). Thus, the induction of PFs by DES may be accompanied by changes in BMP-15 and/or inhibin- α expression. In addition, studies of Müllerian inhibiting substance (MIS) and steroidogenic factor-1 (SF-1) deficient mice suggest that both are critical factors in regulating early ovarian differentiation (Durlinger et al., 1999).

In the present study, DES-treated α ERKO and β ERKO mice were used to examine whether DES acts through ER α and/or ER β in neonatal mouse ovaries to induce PFs and histological abnormalities. Using real-time quantitative PCR, changes in the expression of several genes reported to be involved in folliculogenesis and in PF occurrence were compared between DES-treated and oil-treated mouse ovaries.

MATERIALS AND METHODS

Animals

As the developmental stage of the reproductive tract of mouse neonates corresponds to that of the 3- to 4-month-old human fetus, the neonatal mouse has been used as an animal model of human DES syndrome, in which DES was prescribed during 3–5 months of pregnancy to prevent abortion (Bern and Talamantes, 1981). Therefore, we used neonatal mice for this study.

Female C57BL/6J Jcl mice (CLEA Japan, Tokyo); α ERKO (ER α $^{-/-}$), wild-type (ER α $^{+/+}$), and heterozygous (ER α $^{+/-}$) females; and β ERKO (ER β $^{-/-}$), wild-type (ER β $^{+/+}$), and heterozygous (ER β $^{+/-}$) females were given daily subcutaneous (s.c.) injections of 3 μ g diethylstilbestrol (DES) (Sigma Chemical, St. Louis, MO, USA) dissolved in sesame oil or in the vehicle alone for 5 days, starting on the day of birth (day 0).

α ERKO and β ERKO mice were obtained by mating mice of a mixed C57BL/6/129sv background that were heterozygous for the ER gene disruption, as described previously (Dupont et al., 2000). Pup genotypes were determined by multiplex PCR.

All animals were fed a commercial diet (MF, Oriental Yeast,

Tokyo) and tap water ad libitum and kept at 24 \pm 1.0°C under 12 h light/12 h darkness (artificial illumination; lights on 0800–2000). Animals were maintained in accordance with the NIH Guide for the Care and Use of Laboratory Animals, and the procedures were approved by our institutional animal care committee.

Histological analysis

Ovaries of C57BL/6J mice at days 5, 10, 20, 30, 40, 50, and 60 (n=5), α ERKO and β ERKO mice at day 30 (n=7–10), and the respective wild-type and heterozygous mice (n=4–10) treated neonatally with DES or oil vehicle only were fixed in Bouin's solution overnight at room temperature, embedded in paraffin, and serially sectioned at 8 μ m. Sections were stained with haematoxylin and eosin. Every 13th section of ovaries was observed, and the incidence of PFs (%) was estimated by counting the number of PFs having more than two oocytes per follicle greater than 50 μ m in diameter per mouse, as described previously (Iguchi et al., 1986). The number of mice with PFs in C57BL/6J, α ERKO, or β ERKO mice treated neonatally with DES or the oil vehicle was expressed as the frequency of PFs.

Immunohistochemistry of ER β

Ovaries of C57BL/6J mice at days 0, 5, 10, and 20 (n=3) were fixed in 4% paraformaldehyde in PBS (pH 7.4) overnight at 4°C, embedded in paraffin, and sectioned at 4 μ m. Sections were deparaffinized, rehydrated, and microwaved for 8 min in 10 mM sodium citrate buffer (pH 6.0) for antigen retrieval. Sections were incubated with 0.3% H₂O₂ for 10 min to eliminate endogenous peroxidase. After washing with PBS, sections were treated with normal goat serum for 30 min, and then incubated overnight at 4°C with rabbit polyclonal antibody against mouse ER β (Zymed Laboratories, South San Francisco, CA, USA) at a dilution of 1:250, or with the rabbit IgG fraction (Dako Cytomation, Glostrup, Denmark) as a negative control. After incubation with secondary antibody streptavidin conjugated to horseradish peroxidase, the Elite Standard ABC Kit (Vector Laboratories, Burlingame, CA, USA) and 1 mg/ml diaminobenzidine (Sigma Chemical) in PBS containing 1% H₂O₂ were used for ER β detection.

Real-time quantitative PCR

Total RNA was isolated from ovaries of neonatally DES-treated or oil-treated C57BL/6J mice at days 5, 10, 20, 30, 40, 50, and 60, and neonatally DES-treated or oil-treated wild-type (ER β $^{+/+}$) and β ERKO (ER β $^{-/-}$) mice at days 10, 20, and 30, and reverse-transcribed with Super Script II reverse transcriptase (Invitrogen, Carlsbad, CA, USA) using 0.05 mM oligo dT primer (Invitrogen). Real-time quantitative PCR was carried out with an ABI PRISM 7000 Sequence Detection System using SYBR Green Master Mix (Applied Biosystems, Foster City, CA, USA). The relative mRNA expression of BMP-15, GDF-9, inhibin- α , MIS, kit (KIT), kit ligand (KL), insulin-like growth factor I (IGF-I), SF-1, steroidogenic acute regulatory protein (StAR), cholesterol side-chain cleavage cytochrome P450 (P450SCC), ER α , and ER β (Table 1) was determined by the standard curve method. Cyclophilin was chosen as an internal standard to control for variability in amplification due to differences in starting mRNA concentration. One to ten mice were used for each time point. Independent experiments were carried out at least three times for each gene.

Statistical analysis

Data are expressed as the mean \pm standard error. For multiple comparisons, treatment groups were compared using analysis of variance (ANOVA) followed by Dunnett's post-hoc test. Two-tailed Student's *t*-tests were used for single comparisons. Fisher's exact probability test was used to examine the significance of the association between the two kinds of classification. A statistically significant difference was defined as $p < 0.05$.

RESULTS

Polyovular follicles (PFs) in the ovary of DES-treated C57BL/6J mice

The histology of ovaries from 5- and 10-day-old mice showing ovarian follicles in a variety of stages was not altered by neonatal DES treatment. Corpora lutea were found in the ovaries of C57BL/6J control mice at day 40 or older, but not in the ovaries of any DES-treated mice (data not shown). This result corresponds with a previous report (Iguchi, 1992).

PFs greater than 50 μ m in diameter were found in the ovaries of all DES-treated C57BL mice at day 10 or older. In control mice, 20–100% of mice had PFs in the ovary from days 10 to 60; therefore, DES treatment did not significantly increase the PF frequency (percent of mice with PFs), except in 60-day-old mice (Table 2). The number of follicles greater than 50 μ m in diameter was significantly higher in the ovaries of 20- to 60-day-old mice than in those of 10-day-old mice in both the oil- and DES-treated groups. In 30- and 50-day-old DES-treated mice, the total number of follicles was larger compared to the age-matched oil controls (Fig. 1A). The PF incidence (percent of PFs per 100 follicles greater than 50 μ m in diameter) was significantly higher in DES-treated mice from days 20 to 60 than in the age-matched controls. The PF incidence was highest ($15.7 \pm 2.02\%$) in neonatally DES-treated mice at day 20 and decreased gradually with age (Fig. 1B). The total number of PFs per ovary in DES-treated mice increased to day 30 and then decreased with age. PFs with two or three oocytes were significantly higher at days 20 to 40 (Fig. 1C).

Expression of genes involved in folliculogenesis in C57BL/6J mice

To analyze the mRNA expression of folliculogenesis-related genes, including BMP-15, GDF-9, inhibin- α , MIS, Kit, KL, and IGF-I, real-time quantitative PCR was performed. In

Table 1. Sequences of the oligonucleotide primers used for RT-PCR and real-time quantitative PCR.

Gene	Forward sequence (5' \rightarrow 3')	Reverse sequence (5' \rightarrow 3')
BMP-15	TCCTTGCTGACGACCCTACATT	GGTCAGCCGAACGATGGTATAA
GDF-9	TGGTGGACCTGCTGTTTAACCT	CCAGAAGACATGGCCTCCTTTA
Inhibin- α	CTGCTCTCAATATCTCCTTCCAAGAG	CCATGGCAGTAGTGGAAGATGATGAA
MIS	CCTACATCTGGCTGAAGTGATATGG	GAGGCTCTTGGAACCTCAGCAA
Kit	TACACGTGCAGCAACAGCAA	GAAGGCCAACCAGGAAAAGTT
KL	GGAAAATAGTGGATGACCACGTGT	TGGCCTCTTCGGAGATTCTTT
IGF-I	CAGGCATTGTGGATGAGTGTTG	CACAGTACATCTCCAGTCTCCTCAGA
P450SCC	GTGAATGACCTGGTGCTTCGT	TCGACCCATGGCAAAGCTA
SF-1	TTGCCTCCTGAAAGCCACTCT	CGCGAAAGCAGGAAAGACA
StAR	GCTGCCGAAGACAATCATCA	GATGGACAGACTTGCAGGCTT
ER α	CCTAGCTCAGTCTCTTCTCATTCTT	GGCACAACGTTCTTGCATTTC
ER β	GGTGATTTCAAGAGTGGAATCTC	CCATGCCCTTGTTACTGATGTG
cyclophilin	AGGTCCTGGCATCTTGTCAT	CCATCCAGCCATTGAGTCTTG

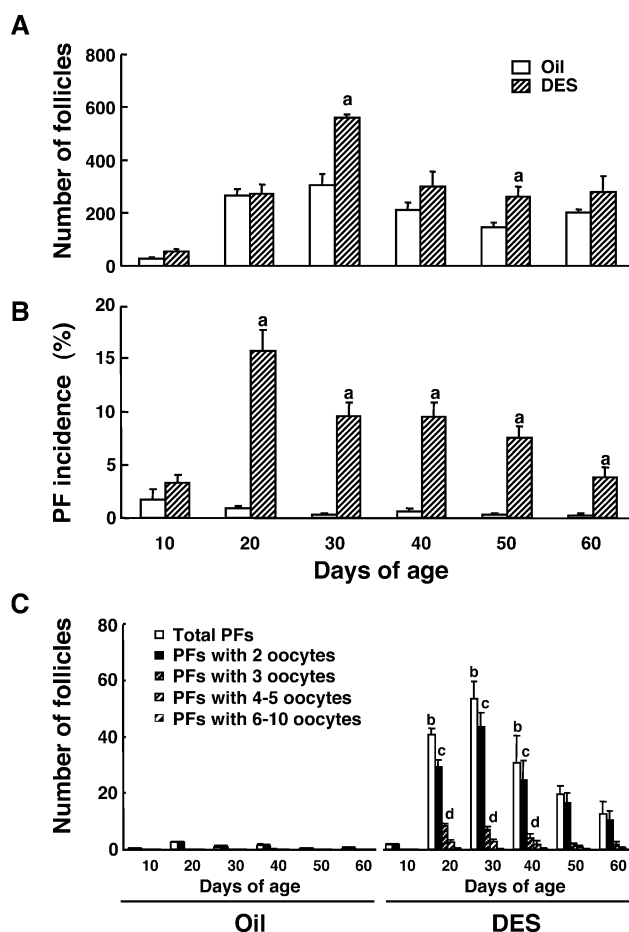


Fig. 1. (A) Number of follicles and (B) incidence of polyovular follicles (PFs) in ovaries of C57BL/6J mice treated neonatally with oil or with 3 μ g DES. (C) Number of total PFs and PFs with 2, 3, 4 to 5, or 6 to 10 oocytes per follicle in ovaries of C57BL/6J mice treated neonatally with oil or with 3 μ g DES. Data are expressed as the mean \pm standard error. a, significantly different (*p* < 0.05) from age-matched oil-treated controls (Student's *t*-test); b, c, d, significantly different (*p* < 0.05) from 10-day-old mice treated neonatally with DES (Dunnett's test). Five mice were used for each time point.

Table 2. Number and frequency of mice with PFs among C57BL/6J mice treated neonatally with the oil vehicle or with 3 μ g DES.

Treatments		Days of age					
		10	20	30	40	50	60
Oil	No. of mice with PFs / No. of mice examined	2 / 5	5 / 5	2 / 5	4 / 6	2 / 5	1 / 5
	PFs frequency (%)	40	100	40	67	40	20
DES	No. of mice with PFs / No. of mice examined	5 / 5	5 / 5	5 / 5	5 / 5	5 / 5	5 / 5*
	PFs frequency (%)	100	100	100	100	100	100

*, significantly different (*p* < 0.05) from oil-treated 60-day-old mice (Fisher's exact probability test).

the ovaries of control mice, the expression of BMP-15, GDF-9, MIS, and IGF-I showed a peak at day 10. The expression of Kit and KL decreased with age. Inhibin- α expression gradually increased to day 20 and remained at a plateau until day 50. Neonatal DES treatment did not affect the expression of any of the genes studied from days 5 to 60, except for inhibin- α , which showed a significantly higher level in DES-treated ovaries than in controls at day 60 (Fig. 2).

Expression of ER α and ER β mRNA and ER β protein in ovaries of C57BL/6J mice treated neonatally with DES

In oil controls, ER α expression showed no alteration from day 0 to day 60. In contrast, ER β mRNA was significantly higher at days 20 and 30 compared to day 0 (Fig. 3). Neither the ER α nor the ER β mRNA level was altered by neonatal DES treatment (Fig. 3), even though control mice from days 40 to 60 had corpora lutea.

No distinct difference in ER β immunoreactivity was discernible between PFs and normal follicles (Fig. 4A). ER β was detected distinctly in the nuclei of granulosa cells. No

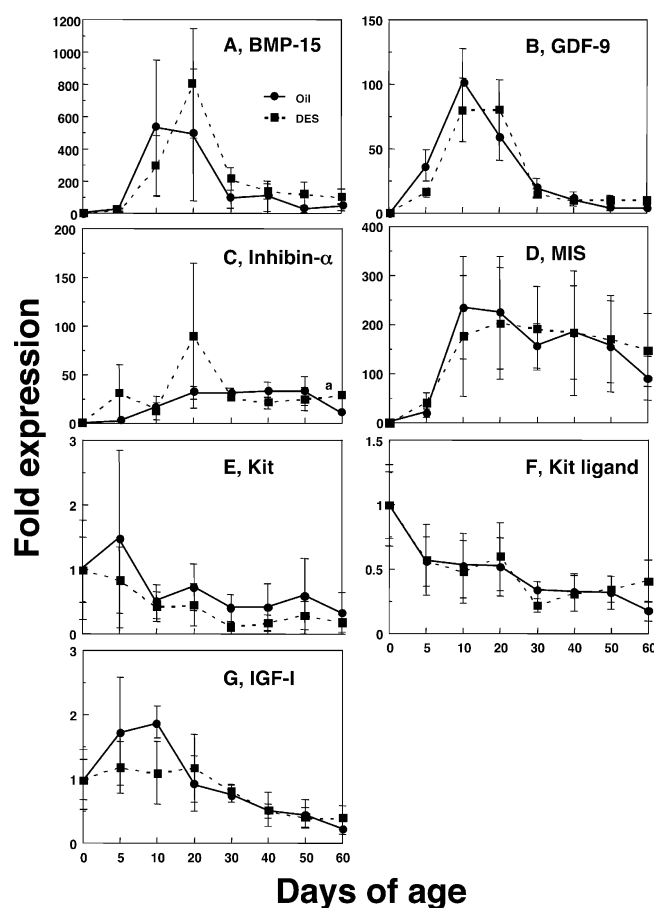


Fig. 2. Ontogenic expression of genes reported to be involved in folliculogenesis in neonatally oil- or DES-treated C57BL/6J mice, determined by real-time quantitative PCR. All measurements are normalized to the cyclophilin mRNA level and are shown relative to the mRNA expression level at day 0. Data are expressed as the mean \pm standard error. a, significantly different ($p < 0.05$) from age-matched oil-treated mice (Student's t -test). One to ten mice were used for each time point, and independent experiments were carried out at least three times for each gene.

ER β staining was observed in the ovaries of mice at day 0; however, granulosa cells began to express ER β protein from day 5 and showed positive immunohistochemical staining through day 20 (Fig. 4C, E). ER β showed no difference in immunoreactivity between the ovaries from DES-treated mice and those from oil controls (Fig. 4C–F).

Histological analysis of ovaries from α ERKO and β ERKO mice at day 30

Ovaries of oil-injected wild-type mice (ER α +/+ and ER β +/+) at day 30 showed various stages of follicles (Fig. 5A). PFs were encountered in the age-matched wild-type mice treated neonatally with DES (Fig. 5B). Hemorrhagic cysts and follicular cysts were found in the ovaries of both oil-treated and DES-treated α ERKO (ER α -/-) mice (Fig. 5C, D). In DES-treated α ERKO mice, PFs were induced as in the DES-treated wild-type mice (Fig. 5D). Unlike α ERKO mice, the ovaries of β ERKO (ER β -/-) mice treated with oil or DES showed normal follicles at all stages of development, without hemorrhagic cysts or follicular cysts (Fig. 5E, F). Neonatal DES treatment in β ERKO mice (Fig. 6B) did not increase the PF incidence. The ovaries of heterozygotes (ER α +/- and ER β +/- mice) were not histologically different from those of wild-type mice, and the effects of DES on the ovary of these heterozygous mice were similar to those in wild-type mice having PFs (data not shown).

PFs in DES-treated α ERKO and β ERKO mice

The number of follicles in ovaries of α ERKO (ER α -/-)

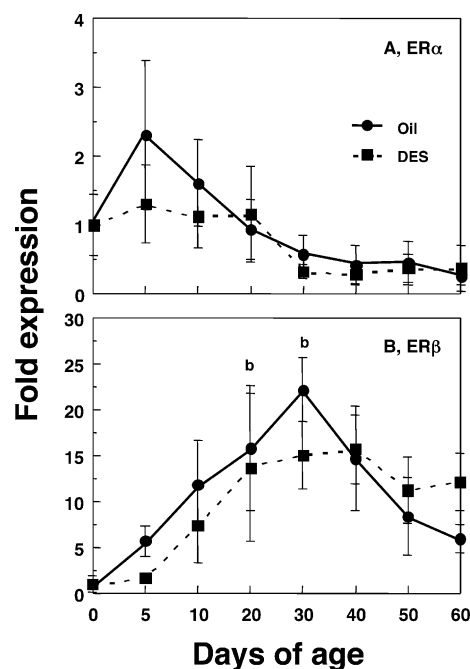


Fig. 3. Ontogenic expression of ER α and ER β mRNAs in the ovaries of oil- or DES-treated C57BL/6J mice, determined by real-time quantitative PCR. Cyclophilin was used as an internal standard to control for variability in amplification due to differences in starting mRNA concentration. Data are expressed as the mean \pm standard error. One to ten mice were used for each time point, and independent experiments were carried out at least three times for each gene. b, significantly different ($p < 0.05$) from 0-day-old control mice.

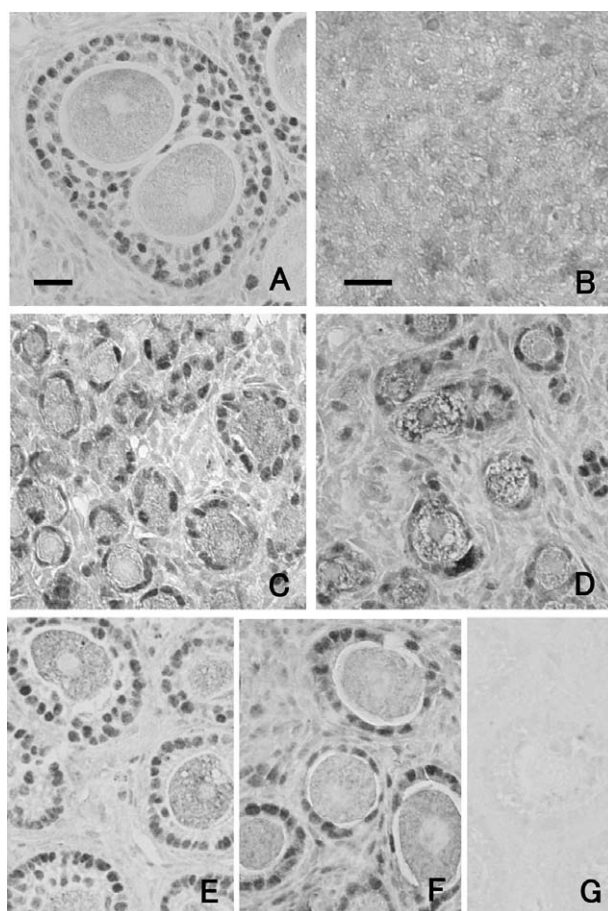


Fig. 4. Immunohistochemical staining of ER β protein in ovaries of C57BL/6J mice. (A) ER β positive staining in a PF at day 20. (B–F) ER β expression in the ovary of a day 0 mouse (B), day 5 mice treated neonatally with oil (C) or with DES (D), and day 10 mice treated with oil (E) or with DES (F). (G) Ovary of a day-10 mouse incubated with the rabbit IgG fraction as a negative control. The cytoplasm of oocytes was occasionally stained, but this was also observed in the negative control. Three mice were used for each time point. Scale bars, 25 μ m.

and β ERKO (ER β –/–) mice was not different from respective wild-type mice (ER α +/, ER β +/) or heterozygous (ER α +/, ER β +/-) mice. In ER β wild-type mice (ER β +/), DES treatment significantly increased the number of follicles (Fig. 6A).

Neonatal DES treatment significantly increased the PF incidence (percent of PFs per 100 follicles greater than 50 μ m in diameter) in ovaries of ER α wild-type, heterozygous, and α ERKO mice (Fig. 6B). The PF incidence was not different among DES-treated α ERKO mice, ER α heterozygous mice, and wild-type mice (9.2–10.5%). DES-treated ER β wild-type and heterozygous mice showed a high incidence of PFs; however, DES treatment did not increase the PF incidence in β ERKO mice compared to that in oil-treated β ERKO mice (Fig. 6B). In the oil controls, 20–60% of mice had PFs in the ovary at day 30, regardless of genotype, and DES treatment did not alter the PF frequency (percent of mice with PFs), except in ER β heterozygous mice (Table 3).

The total number of PFs and the number of PFs with 2, 3, 4 to 5, or 6 to 10 oocytes were not different among gen-

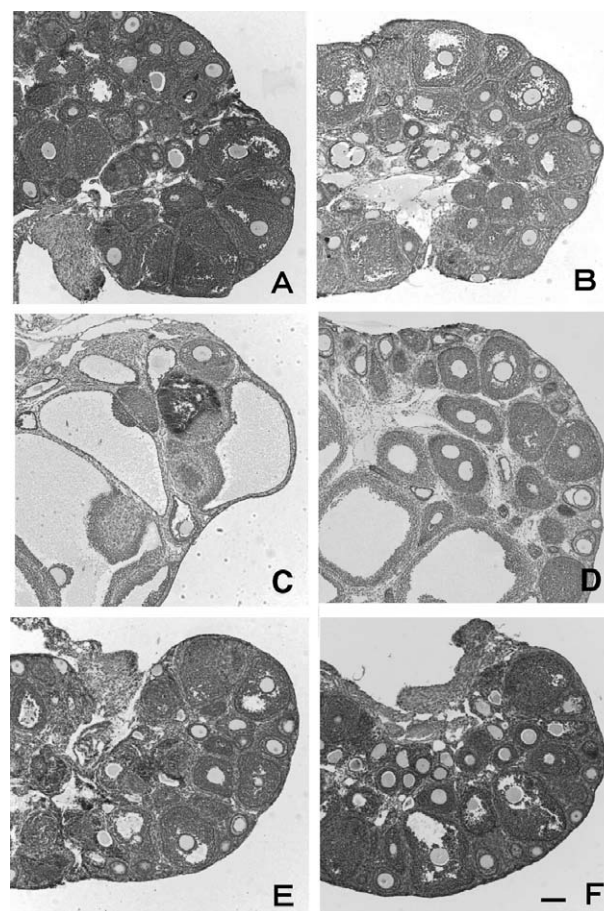


Fig. 5. Ovarian histology of day-30 wild-type ER α +/+ (A) and ER β +/+ (B) mice, α ERKO (ER α –/–) mice (C, D), and β ERKO (ER β –/–) mice (E, F) treated neonatally with oil (A, C, E) or with 3 μ g DES (B, D, F). Four to ten mice were used. Scale bar, 100 μ m.

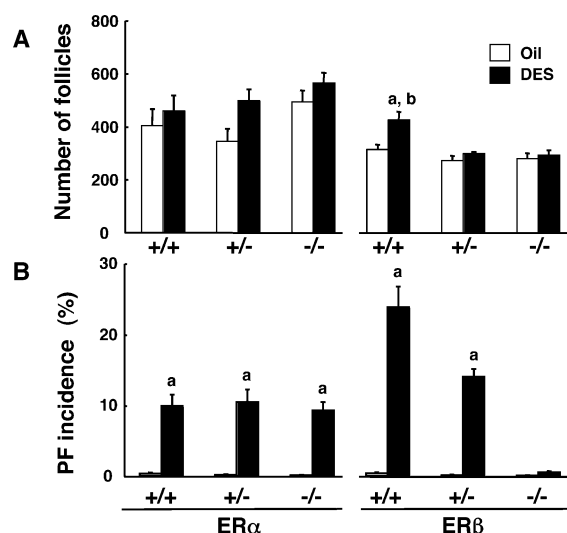
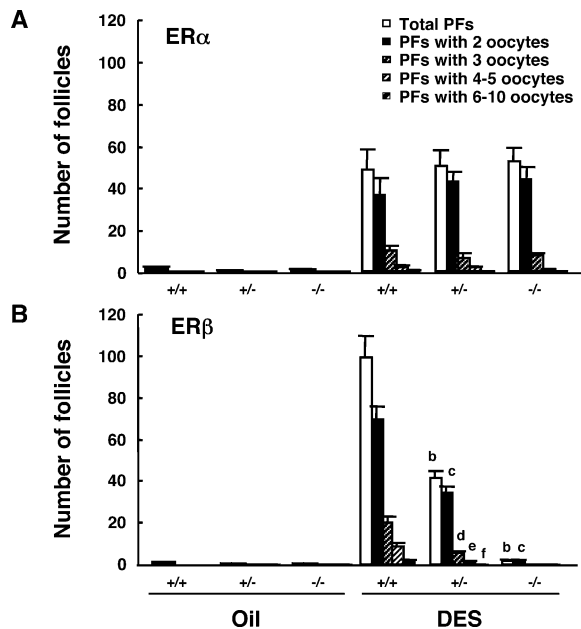


Fig. 6. (A) Number of follicles and (B) incidence of PFs in ovaries of 30-day-old wild-type (ER α +/, ER β +/), heterozygous (ER α +/-, ER β +/-), α ERKO (ER α –/–), and β ERKO (ER β –/–) mice treated neonatally with oil or with 3 μ g DES. Data are expressed as the mean \pm standard error. a, significantly different (p <0.05) from genotype-matched oil-treated mice; b, significantly different (p <0.05) from DES-treated heterozygotes and β ERKO mice (Student's t -test).

Table 3. Number and frequency of mice with PFs among different phenotypes of 30-day-old mice treated neonatally with DES or with the oil vehicle alone.

Treatments	Genotypes	No. of mice with PFs	Frequency of mice with PFs (%)
Oil	ER α +/+	3 / 5	60
	ER α +/-	2 / 5	40
	ER α -/-	3 / 7	43
	ER β +/+	3 / 5	60
	ER β +/-	1 / 4	25
	ER β -/-	2 / 10	20
DES	ER α +/+	5 / 5	100
	ER α +/-	5 / 5	100
	ER α -/-	7 / 7	100
	ER β +/+	5 / 5	100
	ER β +/-	10 / 10*	100
	ER β -/-	5 / 8	63

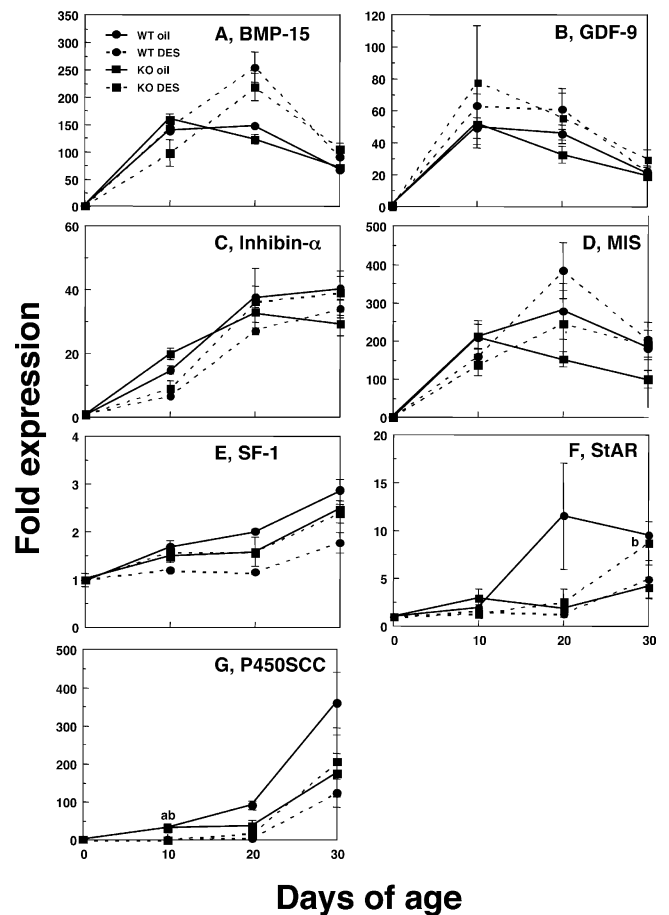
*, significantly different ($p < 0.05$) from oil-treated ER β +/- mice (Fisher's exact probability test).

**Fig. 7.** (A) Number of total PFs and (B) PFs with 2, 3, 4 to 5, or 6 to 10 oocytes per follicle in ovaries of 30-day-old wild-type (ER α +/+, ER β +/+), heterozygous (ER α +/-, ER β +/-), α ERKO (ER α -/-), and β ERKO (ER β -/-) mice treated neonatally with oil or with 3 μ g DES. b, c, d, e, f, significantly different ($p < 0.05$) from total PFs, with 2 oocytes, 3 oocytes, 4–5 oocytes, or 6–10 oocytes in wild-type (ER β +/+) mice treated with DES, respectively. Four to ten mice were used for each time point.

otypes in the DES-treated α ERKO mouse group (Fig. 7A). DES-treated ER β wild-type (ER β +/+) mice showed PFs with 10 oocytes. DES-treated β ERKO mice had PFs with two oocytes, but the number of PFs was significantly lower than in wild-type and heterozygous (ER β +/-) mice treated neonatally with DES (Fig. 7B).

Expression of genes involved in folliculogenesis and steroidogenesis in β ERKO mice

The expression of BMP-15 showed a peak at day 10 in

**Fig. 8.** Ontogenic expression of genes involved in folliculogenesis or steroidogenesis in oil- or DES-treated wild-type (ER β +/+) or β ERKO (ER β -/-) mice, determined by real-time quantitative PCR. All measurements are normalized to the cyclophilin mRNA level and are shown relative to the mRNA expression level of 0-day-old mice. Data are expressed as the mean \pm standard error. a, significantly different ($p < 0.05$) from age-matched oil-treated wild-type mice (Student's t -test); b, significantly different ($p < 0.05$) from age-matched oil-treated β ERKO mice (Student's t -test). One to ten mice were used for each time point, and independent experiments were carried out at least three times for each gene.

oil-injected wild-type mice and β ERKO mice. Neonatal DES treatment shifted the peak to day 20 in both wild-type and β ERKO mice (Fig. 8A). The expression of GDF-9 showed a peak at day 10 in wild-type and β ERKO mice, irrespective of DES treatment. Expression of inhibin- α , SF-1, and P450SCC increased with age. Neonatal DES treatment did not affect the expression of GDF-9, inhibin- α , MIS, or SF-1 in either wild-type or β ERKO mice (Fig. 8B–E). The expression of StAR peaked at day 20 in wild-type mice but remained low in β ERKO mice; however, neonatal DES treatment significantly increased the expression in β ERKO mice at day 30 (Fig. 8F). In both wild-type and β ERKO mice, neonatal DES treatment significantly reduced the expression of P450SCC at day 10 (Fig. 8G).

DISCUSSION

In mouse ovaries, ER α protein and mRNA are detected

in the nuclei of interstitial and thecal cells, and ER β protein and mRNA are found in the nuclei of granulosa cells, but not in cells in atretic follicles or corpora lutea (Hishikawa et al., 2003). The localization of ERs suggests that ER β may play a principal role in follicular development, while ER α may be involved in LH action and androgen synthesis. Indeed, α ERKO mouse ovaries exhibit not only hemorrhagic and cystic follicles but also growing follicles in the pre-antral to small antral stages. The phenotypes of β ERKO mouse ovaries suggest a reduction in completed folliculogenesis (Couse and Korach, 1999). Thus, ER α and ER β are quite distinct in function in the rodent ovary.

In the present study, neonatal DES treatment increased neither the PF frequency nor the PF incidence in β ERKO mice, whereas DES treatment significantly increased the PF incidence in α ERKO mice. PFs were also found in C57BL/6J control mice (20–100%), but the PF incidence was low. All DES-treated C57BL/6J mice showed PFs, and the PF incidence increased significantly at day 20 and older. Nakamura et al. (2008) showed that neonatal treatment with a specific ligand of ER α (propyl pyrazole triol) induces PFs in C57BL mice, but they also showed in a transactivation assay that the ligand could activate ER β activity at a higher concentration. Therefore, the possibility cannot be ruled out that the ER α -specific ligand acts through ER β in the mouse ovary. Jefferson et al. (2002) reported the induction of PFs by neonatal treatment with 100 μ g/pup of genistein, a potent phytoestrogen, at day 19, and they found PFs in only one of 12 β ERKO mice. While the binding affinity of genistein to ER β is 20 times higher than that to ER α , DES binds to ER α and ER β with equal relative affinity (Kuiper et al., 1998). Although the affinities of ER α and ER β to DES were equal, DES treatment did not significantly increase the PF incidence in β ERKO mice, but did significantly increase the PF incidence in α ERKO mice, suggesting that neonatal DES treatment induces PFs through ER β but not ER α . In the present study, ER β protein was not detected at day 0, but it was detected by day 5. In addition, neonatal treatment with an ER β -specific agonist, 2,3-bis(4-hydroxyphenyl)-propionitrile (25 μ g/pup for 5 days from the day of birth, $n=5$), also induced PFs in all wild-type mice at day 30. The PF incidence in oil controls and the ER β agonist-treated mice was 0.6 and 4.3%, respectively (unpublished data).

The estrogen-related receptors (ERRs), an orphan family of nuclear receptors identified initially by similarity to ER α (Giguère et al., 1988), are not activated by natural estrogens, but DES can bind and inhibit the transcriptional activity of ERR α , β , and γ (Tremblay et al., 2001). While ERR α has a potential role in the regulation of metabolic homeostasis, loss of ERR β or ERR γ results in mid-gestational or neonatal death due to placental defects or failure of the perinatal transition in metabolic state (Luo et al., 1997; 2003; Alaynick et al., 2007; Dufour et al., 2007). However, the expression of ERRs in the ovaries of neonatal wild-type and β ERKO mice has not been examined; therefore, the role of ERRs in the induction of PFs by DES needs to be clarified in the near future.

An increase and decrease of ER β mRNA with age may be correlated with follicle number and follicular development, because ER β is high in preantral and small antral follicles, but not in the Graafian follicles or corpora lutea in rats

(Fitzpatrick et al., 1999). In contrast, ER α is found in the interstitial cells of the ovary from the day of birth, and mRNA expression is not altered until 26 days of age in mice (Jefferson et al., 2000). The ontogeny and distribution of ER α and ER β in the rat ovary are similar to that in the mouse ovary (Sar and Welsch, 1999). In addition, no ER β staining was observed in the ovaries of mice at day 0; however, positive immunohistochemical staining of ER β was evident in granulosa cells from days 5 through 20, coinciding with the changes in ER β mRNA expression.

In adult human females, the incidence of PFs ranges from 24% to 85%, depending on the study (Gougeon, 1981; Dandekar et al., 1988); both reports agree that the proportion of PFs among total follicles in the human ovary is less than 1%. No report is available concerning PFs in DES-exposed humans; however, chronic implants of DES increase PFs in adult squirrel monkeys (Graham and Bradley, 1971). In the human ovary, ER β is localized in the granulosa cells of follicles at all stages of development, whereas ER α is absent from granulosa cells but present in thecal cells (Pelletier and El-Alfy, 2000). Thus, it appears that each receptor subtype can exert a different function according to its specific cellular localization in the human ovary.

In the ovary, ER β is highly expressed in granulosa cells, as discussed above. Our results show clearly that ER β participates in the induction of PFs by neonatal DES exposure. ER β expression is observed in the granulosa cells, but not in the oocytes, of rat and human ovaries (Sar and Welsch, 1999; Jakimiuk et al., 2002; Scobie et al., 2002), while ER β mRNA is detected in the cumulus-oocyte complexes and denuded oocytes of mice after PMSG and hCG injections (Hiroi et al., 1999). Jefferson et al. (2006) demonstrated that the suppression of oocyte cell death by neonatal genistein treatment results in PFs. Our previous report suggests that a delay in or inhibition of oocyte death and an increase in inhibin- α by DES through ER β may be a trigger for PFs formation (Kim et al., in press). Therefore, DES may indirectly affect oocyte cell death and induce PFs.

The PF incidence increases only when neonatal DES treatment starts within 3 days after birth (Iguchi et al., 1986). Exposure to DES from days 15 to 18 of gestation in utero also induces a high incidence of PFs in offspring (Iguchi and Takasugi, 1986). Folliculogenesis proceeds in ovaries of the fetus and neonate. Syncytia, clusters formed by germ cells, are invaded by somatic cells and undergo programmed breakdown before the formation of primordial follicles (Epifano and Dean, 2002). Thus, DES may induce PFs by disrupting the interaction between oocytes and somatic cells or granulosa cells, and/or by disrupting syncytial breakdown in folliculogenesis.

Folliculogenesis is under the control of molecular interactions between germ cells and somatic cells, and is thought to involve several genes, e.g. Fig α , Wnt4, Kit, KL, Cx43, MIS, GDF9, and FecX (Epifano and Dean, 2002). Double mutant Gdf9 $^{+/-}$ and Bmp15 $^{-/-}$ mice show PFs in their ovaries without DES treatment (Yan et al., 2001). The expression of GDF-9 and BMP-15 is localized in oocytes (McGrath et al., 1995; Dube et al., 1998), and that of inhibin- α and MIS in granulosa cells (Meunier et al., 1988; Ikeda et al., 2002). In the present study, neonatal DES exposure did not alter the expression of these genes in C57BL/6J, ER β wild-type,

or β ERKO mice; therefore, further studies are needed to clarify the genes related to PF induction by neonatal DES exposure.

Kipp et al. (2007) reported that neonatal DES or estrogen treatment decreased activin β -subunit mRNA and protein levels, with a reduction in the number of small antral follicles and the induction of PFs on postnatal day 19. Moreover, mice overexpressing inhibin- α show PFs in their ovaries without DES treatment (McMulle et al., 2001). In our study, the expression of inhibin- α subunit mRNA in DES-treated C57BL/6J mice was significantly higher than in oil controls only at day 60. At day 5, inhibin- α subunit mRNA showed no significant change. Thus, the up-regulation of inhibin- α at day 60 may not be directly related to PF induction by DES. The balance of inhibin and activin expression in the ovary could be important in the induction of PFs, and this needs to be clarified in the near future.

MIS and SF-1 are detected in rat granulosa cells, and MIS is a target gene of SF-1 (Ikeda et al., 1994, 2002; Hanley et al., 2000). Both MIS and SF-1 are critical factors regulating early ovarian differentiation (Durlinger et al., 1999). Therefore, we studied the effects of DES on expression of these genes; however, we detected no change in the expression of these genes.

In both wild-type and β ERKO mice, DES treatment significantly reduced the expression of P450SCC at day 10. The expression of P450SCC is localized in interstitial cells (Ikeda et al., 2001), and these cells contain ER α protein. Thus, DES possibly suppresses P450SCC mRNA expression via ER α . Since hypertrophy of the interstitium is not observed in neonatally DES-treated α ERKO mice (Couse et al., 2001), ER α may mediate the effects of DES on interstitial cells.

In conclusion, neonatal DES treatment induces PFs through ER β but not ER α , while DES acts on ovarian interstitial cells through ER α . Further studies are essential to clarify the molecular mechanisms involved in the induction of PFs.

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