The influence of di-(2-ethylhexyl) phthalate on steroidogenesis by the ovarian granulosa cells of immature female rats

I Svechnikova, K Svechnikov and O Söder

Pediatric Endocrinology Unit, Department of Woman and Child Health, Karolinska Institute and University Hospital, Q2:08, 171 76 Stockholm, Sweden

(Correspondence should be addressed to I Svechnikova; Email: irina.svechnikova@ki.se)

Abstract

Phthalate esters are known to exert harmful effects on mammalian reproduction and fertility, but their potential adverse effects on the hormonal functions of the ovary have not yet been elucidated in detail. Here, we investigated the effects of di-(2-ethylhexyl) phthalate (DEHP) on the hypothalamic–pituitary–gonadal axis of young developing female rats, as well as on *ex vivo* steroidogenesis by granulosa cells (GCs) and secretion of LH by gonadotropes. Exposure of 20-day-old female rats to 500 mg DEHP by oral gavage once daily for 10 days reduced their serum levels of progesterone and estradiol, while tending to enhance levels of LH. Furthermore, primary cultures of GCs isolated from these rats exhibited an attenuated capacity to produce progesterone in response to stimulation by

LH and FSH, as well as a lower degree of transport of endogenous cholesterol into mitochondria. Moreover, the ability of primary cultures of pituitary cells isolated from DEHP-treated rats to produce and secrete LH in response to GnRH was significantly enhanced. In addition, 2-ethylhexanoic acid, a metabolite of DEHP, significantly potentiated GnRH-stimulated production of LH by cultures of pituitary cells isolated from untreated 20-day-old female rats. Together, these data indicate that DEHP exerts dual effects on the pituitary–gonadal axis, stimulating the hormonal function of the pituitary and, at the same time, by inhibiting steroidogenesis by GCs.

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Introduction

Phthalates, including the widely used plasticizer di-(2-ethylhexyl) phthalate (DEHP), are the most abundant pollutants of our general environment. These substances are components of food wraps and many medical devices (e.g. tubing, blood bags, and dialysis equipment), as well as of many cosmetic products. With time, they are leached out of these plastic products, and their volatility results in pronounced human exposure to phthalates.

Investigations on the influence of phthalates on the female reproductive system have focused primarily on fertility and teratogenicity in experimental animals and cattle (Collins *et al.* 1992). Little information concerning the effects of phthalates on pregnant women is presently available, although chronic occupational exposure of female factory workers to high levels of phthalates has been reported to be associated with reduced rates of pregnancy and an elevated frequency of miscarriage (Aldyreva *et al.* 1975). Exposure of adult rats to DEHP results in hypoestrogenic anovulatory cycles and polycystic ovaries (Davis *et al.* 1994a).

In the intestine and liver of humans and animals, DEHP is rapidly hydrolyzed by esterases to yield mono-(2-ethylhexyl) phthalate (MEHP) and 2-ethylhexanol (Rowland 1974,

Albro *et al.* 1983); and this hexanol is subsequently oxidized enzymatically to 2-ethylhexanoic acid (2-EHXA; Collins *et al.* 1992). MEHP, 2-ethylhexanol, and/or their metabolites are the immediate inducers of the majority of enzymes known to be affected by exposure to DEHP (Pollack *et al.* 1989).

In primary cultures of rat granulosa cells (GCs), stimulation of adenylate cyclase and progesterone synthesis by follicle-stimulating hormone (FSH) is inhibited by MEHP (Treinen *et al.* 1990). Moreover, this compound suppresses estradiol production by the same cells by a mechanism that is independent of the elevation in cAMP levels caused by FSH (Davis *et al.* 1994b). This latter phenomenon reflects direct inhibition by MEHP of transcription of the gene that encodes aromatase, the enzyme that converts androgen to estrogen (Lovekamp & Davis 2001). DEHP also suppresses the production of estradiol and ovulation in cycling rats (Davis *et al.* 1994a). However, in the case of immature female rats, the influence of *in vivo* administration of DEHP on the hypothalamic–pituitary–gonadal (HPG) axis and progesterone production by GCs isolated and cultured *ex vivo* has not yet been characterized. Here, we investigated the effects of DEHP on the HPG axis of immature female rats, as well as *ex vivo* steroidogenesis by GCs and secretion of luteinizing hormone (LH) by gonadotropes.

Materials and Methods

Materials

Dulbecco's modified Eagle's medium (DMEM)–Ham's nutrient mixture F-12 (supplemented with L-glutamine and HEPES), BSA (7.5% solution), cholesterol lipids (250×), and penicillin–streptomycin (Gibco/BRL, Life Technologies); ovine FSH, ovine LH, 22R-hydroxycholesterol (22R-OHC), DEHP, EHXA, and aminoglutethimide (AMG; an inhibitor of cytochrome P450_{sc}; Sigma); MEHP (TCI Europe, Brussels, Belgium); trilostane (an inhibitor of 3 β -hydroxysteroid dehydrogenase; Stegram Pharmaceuticals, Billingham, Sussex, UK); and the cell proliferation WST-1 kit (Roche Diagnostic GmbH) were purchased from the sources indicated. The rat LH used for iodination (NIDDK-rLH-I-10, Lot # AFP-11536B), antiserum against rat LH (NIDDK-anti-rLH-S11, Lot # AFPC697071P), and LH utilized as a reference standard (NIDDK-rLH-RP-3, Lot # AFP-7187B) were obtained from Dr Parlow of the National Hormone and Peptide Program, NIDDK, USA. The secondary goat antiserum against rabbit IgG was purchased from the Sigma.

Animals

Immature (20-day-old) Sprague–Dawley female rats (B&K Laboratories, Sollentuna, Sweden) were randomly divided into two groups of ten animals each, one of which was administered DEHP (500 mg/kg dissolved in corn oil) and the other a corresponding volume of corn oil alone (controls) by gavage once daily for 10 days. This dose of DEHP has been shown to effectively elicit response, but not toxicity in male rats of approximately the same age (Supornsilchai *et al.* 2007). Twenty-four hours after the last gavage, the animals were anesthetized with pentobarbital, blood samples collected by intracardiac puncture, and plasma prepared by centrifugation at 2500 g for 15 min, and thereafter maintained at –80 °C for subsequent analysis of progesterone, estradiol, and LH. The ovaries and pituitary glands were removed from the same rats for isolation of granulosa and pituitary cells.

The animal experiments performed here were pre-approved by the Northern Stockholm Committee for Ethical Animal Experimentation.

Isolation and primary culture of GCs

GCs were isolated from immature rat ovaries, as described previously (Treinen *et al.* 1990). In brief, after trimming away the surrounding tissue, the ovaries were placed in 3 ml DMEM:F-12 (1:1) medium supplemented with penicillin (100 U/ml), streptomycin (100 µg/ml), and 0.5% BSA and then punctured with a 25-gauge needle. The resulting cell suspension was first filtered through a 70 µm cell strainer, following which the cells were collected by centrifugation at 200 g for 7 min, washed once with the same medium, and

thereafter resuspended in a corresponding volume of fresh DMEM:F-12 (1:1) medium containing cholesterol lipid concentrate (1:250; added to mimic the cholesterol content of the follicular fluid *in vivo*). Subsequently, 100 000 GCs in 100 µl were placed into each well of 96-well plates (Falcon, Franklin Lakes, NJ, USA), cultured for 2 h at 37 °C under 5% CO₂ and then incubated in the absence or presence of FSH (50 mU/ml), LH (100 mU/ml), and/or 22R-OHC (10 µM) for an additional 24 h in a total volume of 250 µl.

Isolation and primary culture of pituitary cells

Cells were isolated from the anterior pituitary glands of both control and DEHP-treated rats as follows: the glands were first minced utilizing a scalpel, and the pieces thus obtained were incubated with collagenase (2 mg/ml) for 30 min at 37 °C. The resulting cell suspension was then filtered through nylon gauze (70 µm; Becton, Franklin Lakes, NJ, USA), following which the cells were collected by centrifugation at 300 g for 7 min, washed twice with DMEM:F-12 (1:1) medium containing 0.5% (w/v) BSA, and finally resuspended in the same medium supplemented with 15 mM HEPES (pH 7.4), 0.5% BSA, glutamine (365 mg/l), penicillin (100 IU/ml), and streptomycin (100 µg/ml). The viability of these cells, as assessed by Trypan blue exclusion, was routinely >90%.

Ten thousand of these cells were placed into each well of 96-well plates. They were pre-incubated for 1 h at 37 °C under 5% CO₂ and subsequently cultured in the absence or presence of gonadotropin-releasing hormone (GnRH; 0.01, 0.1, and 1.0 ng/ml) for an additional 3 h. In similar experiments, cells isolated from 20-day-old female rats, treated neither with DEHP nor with vehicle, were cultured with or without DEHP, MEHP, or EHXA (1 µM) for 24 h at 37 °C, at which time fresh culture medium containing the same compound together with GnRH (1 ng/ml) was added and incubation continued for another 3 h. Following these procedures, the culture media were collected and stored at –80 °C until subsequent assay for LH.

Determination of hormone concentrations

Plasma samples and media from the GC cultures were used for determination of the concentrations of progesterone and estradiol present employing the Coat-a-Count RIA and Double Antibody Estradiol kits respectively (Diagnostic Products Corp., Los Angeles, CA, USA) in accordance with the manufacturer's instructions. In addition, the LH levels in the same samples of serum media from the pituitary cell cultures were measured employing rat LH labeled with ¹²⁵I by the routine Chloramine-T procedure.

Monitoring cholesterol transport into mitochondria

For this purpose, 100 000 GCs isolated from control or DEHP-treated rats and suspended in 100 µl medium were placed into each well of 96-well Falcon plates (Falcon), where

they were first incubated at 37 °C under an atmosphere containing 5% CO₂ for 24 h. Thereafter, fresh medium containing AMG (0.5 mM), an inhibitor of cytochrome P450_{sc}, and trilostane (5 µM), an inhibitor of 3β-hydroxysteroid dehydrogenase, was added and incubation continued for an additional 30 min, following which the medium was supplemented with LH (100 mU/ml) and incubation performed for 3 h more. Inhibition of mitochondrial cholesterol metabolism by AMG and trilostane results in accumulation of this steroid in this organelle in response to hormonal stimulation (Potts *et al.* 1978, Brueggemeier *et al.* 2005).

Subsequently, the cells were washed twice with the basic medium and then cultured in the presence of trilostane (5 µM) in AMG-free medium at 37 °C under 5% CO₂ for 2 h, after which the levels of pregnenolone and its metabolite dehydroepiandrosterone (DHEA; formed by cytochrome P450_{c17} via the Δ⁵ pathway) in the medium were measured. The total combined amount of these steroids produced by the cells reflects the amount of cholesterol available to the cytochrome P450_{sc} system (Potts *et al.* 1978). After lysing the cells with the M-PER Mammalian Protein Extraction Reagent (Pierce, Rockford, IL, USA), protein was determined using the Micro BCA Protein Assay from the same company and the total amount of these hormones produced expressed per milligram protein.

Statistical analysis

Differences in the values obtained for animals or cells subjected to different treatments were evaluated for statistical significance with Student's *t*-test and one-way ANOVA, followed by the Student–Newman–Keul's or Dunn's test, in all cases employing the SigmaStat (v 3.00) software package (SPSS Inc., Chicago, IL, USA).

Results

Oral administration of DEHP to immature female rats decreases their serum levels of progesterone and estradiol

Administration of DEHP by oral gavage to 20-day-old female rats for 10 days resulted in two- and fivefold reductions in the concentrations of progesterone and estradiol respectively in their plasma, together with a tendency toward an elevated level of plasma LH (Fig. 1A–C). This exposure did not exert any obvious toxic effects on the animals, as reflected in their unaltered body and ovary weights (Table 1).

Treatment of immature female rats with DEHP inhibits *ex vivo* steroidogenesis by their GCs

In an attempt to identify the primary target through which DEHP impairs the biosynthesis of sex hormones, the capacities of GCs isolated from treated and control rats to produce progesterone under *ex vivo* conditions were

Figure 1 Plasma levels of progesterone (A), estradiol (B), and LH (C) in 20-day-old female rats exposed to DEHP (500 mg/kg per day) by oral gavage for 10 days. The control animals received a corresponding volume of vehicle (corn oil) alone by the same route. Each bar represents the mean ± S.E.M. (*n* = 9–10). ***P* < 0.01; ****P* < 0.001 when compared with the corresponding control value.

compared. Following stimulation with FSH or LH, progesterone production by GC isolated from DEHP-treated animals was 30% lower (*P* < 0.05) than by control cells (Fig. 2). Moreover, these cells isolated from young developing female rats, whether treated or not, were poorly responsive to gonadotropins, suggesting that the signaling machinery involved was not yet fully developed. More detailed information concerning the site in steroidogenesis inhibited by DEHP was provided using 22R-OHC, a derivative of cholesterol that diffuses readily across membranes. DEHP exerted no effect on progesterone synthesis from 22R-OHC,

Table 1 The weights of the whole body and both ovaries of 20-day-old female rats prior to and following treatment with di-(2-ethylhexyl) phthalate (DEHP) or vehicle alone for 10 days

	Age (days)	Body weight (g)	Weight of ovaries (mg)
Group			
Prior to treatment	20		
Control		42.6 ± 0.9	ND
DEHP		45.7 ± 1.6	ND
Following treatment	30		
Control		69.5 ± 1.7	12.3 ± 0.6
DEHP		77.0 ± 3.7	12.7 ± 0.7

The rats received DEHP (500 mg/kg) or the corresponding volume of vehicle (corn oil) alone each day for 10 days (n=10). ND, not determined.

suggesting that this phthalate may be disturbing the translocation of cholesterol across mitochondrial membranes.

Treatment of immature female rats with DEHP reduces ex vivo transport of cholesterol into the mitochondria of their GCs

Since the suppression by DEHP of steroidogenesis in GCs was abolished by 22R-OHC, we examined whether treatment with this phthalate affects the transport of cholesterol into the mitochondria of GCs. Indeed, the level of DHEA produced (which reflects the amount of cholesterol available to the mitochondrial cytochrome P450scc system) was significantly reduced (33%) by such treatment (Fig. 3). The level of pregnenolone detected was negligible, indicating that when 3β-HSD (3β-hydroxysteroid dehydrogenase) was blocked, this entire compound was converted further into DHEA by CYP17α.

Figure 2 Progesterone production by granulosa cells isolated from the ovaries of DEHP-treated (gray bars) and control (black bars) 20-day-old female rats and cultured *ex vivo*. Following incubation of these cells with FSH, LH, and 22R-OHC or vehicle only (0) for 24 h, the concentrations of progesterone in the culture media were determined by RIA. These results depict the means ± S.E.M. of the values obtained with three independent preparations of granulosa cells. *P<0.05 when compared with the corresponding control value.

Figure 3 *Ex vivo* transport of cholesterol into the mitochondria of granulosa cells isolated from DEHP-treated (gray bars) and control (black bars) immature female rats. Following pretreatment of the cells with aminoglutethimide (AMG; 0.5 mM) and trilostane (5 μM) for 30 min and subsequent incubation in the presence (LH) or absence (basal) of LH (100 mU/ml) for 3 h, the DHEA present in the culture medium was measured by RIA procedure. The results presented are the means ± S.E.M. for three independent experiments. **P<0.01; ***P<0.001 when compared with the corresponding control values.

Treatment of immature female rats with DEHP enhances ex vivo secretion of LH by their pituitary cells

Our findings that the serum levels of steroid hormones in DEHP-treated immature female rats were reduced, together with a tendency toward an increase in the serum level of LH, suggest that the *ex vivo* observations presented above might reflect activation of the pituitary by the attenuated levels of steroid hormones in the treated animals. Consequently, release of LH by pituitary cells isolated from control and DEHP-treated rats was examined and proved to be significantly (twofold) elevated, both with and without stimulation by GnRH, following exposure (Fig. 4).

Effects of in vitro exposure to DEHP or its metabolites on LH secretion by pituitary cells isolated from immature female rats treated neither with DEHP nor with vehicle

Subsequent experiments were designed to test the hypothesis that DEHP and/or metabolites of this phthalate can directly stimulate the pituitary of developing rats to produce LH. In order to assess contribution of all these compounds into the activating effect of DEHP on these cells observed *ex vivo*, the effects of DEHP and its predominant metabolites, MEHP and 2-EHXA, on hormonal function of cultured pituitary cells were studied. Indeed, 2-EHXA, but neither DEHP nor MEHP, significantly enhanced (by 30%, P<0.05) GnRH-stimulated production of LH by cultures of pituitary cells isolated from 20-day-old female rats with no pretreatment whatsoever (Fig. 5). However, this compound had no effect on basal production of LH by the same cells (the control and 2-EHXA-exposed values were 40.8 ± 1.0 and 46.7 ± 2.5 ng/10⁴ cells/24 h respectively).

Figure 4 *Ex vivo* production of LH by pituitary cells isolated from DEHP-treated (open circles) and control (filled circles) immature female rats, both with and without stimulation by GnRH. The pituitary cells were incubated without or with various concentrations of GnRH for 3 h, after which the concentration of LH in the culture medium was determined employing RIA procedure. The results represent means $\pm$ S.E.M. of three independent pituitary cell preparations. * $P < 0.05$; ** $P < 0.01$; *** $P < 0.001$ when compared with the control value (i.e. cells from control animals without exposure to DEHP).

Figure 5 *In vitro* effects of DEHP and its metabolites, MEHP and 2-ethylhexanoic acid (EHXA), on GnRH-stimulated production of LH by pituitary cells isolated from 20-day-old female rats treated neither with DEHP nor with vehicle. The cells were cultured with or without DEHP, MEHP, or EHXA (1 μ M) for 24 h at 37 °C. At this time point, fresh culture medium containing the same compound together with GnRH (1 ng/ml) was added and incubation continued for 3 more h, after which the concentration of LH in the culture medium was determined by RIA procedure. These results are means $\pm$ S.E.M. for three independent pituitary cell preparations. * $P < 0.05$ when compared with the basal level (0); † $P < 0.05$ when compared with GnRH alone.

Discussion

The present investigation demonstrates that the treatment of pre-pubertal 20-day-old female rats with DEHP lowers their serum levels of progesterone and estradiol and, at the same time, suppresses progesterone production by and transport of cholesterol into the mitochondria of GCs isolated from their ovaries. In addition, we found that primary cultures of pituitary cells prepared from DEHP-treated rats exhibit an enhanced capacity to secrete LH, which may explain the observed tendency for serum gonadotropin levels to increase in these treated animals. Moreover, we demonstrate here that 2-EHXA, a metabolite of DEHP, significantly enhances GnRH-stimulated production of LH by pituitary cells isolated from 20-day-old female rats that have not been treated in any way. To the best of our knowledge, this is the first example of direct modulation of the hormonal function of the pituitary by a metabolite of DEHP. Together, these results indicate that DEHP exerts a dual effect on the pituitary–ovary axis, both stimulating production of LH by the pituitary and inhibiting steroidogenesis in GCs.

Our observation that administration of DEHP to immature female rats by oral gavage reduces their serum levels of progesterone and estradiol (Fig. 1) is consistent with previous findings that serum levels of cholesterol and triglycerides are lowered by dietary administration of phthalates and/or their metabolites to rats (Moody & Reddy 1982). Two metabolites of DEHP, 2-ethylhexanol and EHXA, specifically reduce the level of cholesterol, the precursor of all steroid hormones, which may attenuate production of, for example, progesterone and estradiol. This reasoning is also consistent with the previous report that MEHP, one of the major and active metabolites of DEHP, inhibits transcription of the aromatase gene in GCs (Lovekamp & Davis 2001).

In addition, a decrease in serum levels of estradiol following exposure of cycling rats to DEHP had been observed earlier, along with lowered serum levels of progesterone at certain stages of the estrous cycle (Davis *et al.* 1994a). Similar inhibition of serum and total ovarian progesterone levels by dibutyl phthalate was shown in female rats during pregnancy (Gray *et al.* 2006). However, the present study represents for the first time that the *ex vivo* steroidogenic capacity of GCs isolated from rats receiving DEHP by oral administration has been examined, and we show that the responsiveness of these cells to gonadotropins is attenuated. This may result in decreased production and secretion of progesterone into the serum, which could also contribute to our present findings.

The dose of DEHP (500 mg/kg) used in this investigation was shown to effectively increase the levels of corticosterone and ACTH in plasma of 20- and 40-day-old male rats (Supornsilchai *et al.* 2007). As in our study, no overt toxicity was observed in these animals. The dose of DEHP used in this investigation (500 mg/kg) was lower when compared with doses commonly used by others. In similar studies on female rats, doses of DEHP as high as 1500–2000 mg/kg have been

used (Laskey & Berman 1993, Davis *et al.* 1994a). In a study on human exposure, the calculated intake of DEHP by women reached a maximum of 15 µg/kg per day (McKee *et al.* 2004). However, for certain groups of neonatal patients at the neonatal intensive care unit, exposure to DEHP leaching from medical devices could reach 1780 µg/kg per day (Koch *et al.* 2006). High levels of phthalate metabolites have also been found in human urine (Hauser *et al.* 2004). These observations show that there is a great deal of uncertainty about the individual levels of exposure to phthalates. Further, to correlate toxic effects observed in animals with acceptable levels of exposure for humans, tenfold increases in the levels of exposure are commonly used to compensate for the inter- and intraspecies variability (Xu *et al.* 2005).

Transport of cholesterol into the mitochondria of GCs is also impaired by exposure of rats to DEHP. This phenomenon was reflected by the fact that under conditions where further conversion of pregnenolone to progesterone is blocked by trilostane, accumulation of DHEA, the final product, by GC from DEHP-treated rats is significantly less than in the case of control rats. Similar impairment of cholesterol transport in the fetal testis by di(*n*-butyl) phthalate associated with attenuated expression of scavenger receptor B1 and the steroidogenic acute regulatory protein has been reported (Thompson *et al.* 2004). The suggestion that inhibition of cholesterol transport is at least one of the deleterious effects of phthalates on steroidogenesis is strongly supported by our finding that in the presence of a cholesterol derivative that readily penetrates membranes, 22R-OHC, GCs from control and DEHP-treated rats produce equal amounts of progesterone.

Our finding that the capacity of pituitary cells isolated from DEHP-treated immature female rats to produce LH is enhanced might probably reflect less pronounced negative feedback by progesterone and estradiol on the hypothalamus and/or the pituitary of these developing animals. However, our data also indicate that the DEHP metabolite, 2-EHXA, stimulates LH production by the pituitary directly; of course, an indirect action via attenuated negative feedback on the pituitary may also be occurring. The existence of such an indirect action is supported by the observation that exposure of pre-pubertal male rats to DEHP for 28 days stimulated their pituitary–gonadal axis, thereby resulting in elevated serum levels of LH (Akingbemi *et al.* 2001), as well as by the findings that chronic administration of DEHP to male rats results in hypertrophy of gonadotropes in the anterior pituitary and promotes development of pituitary tumors (Kluwe *et al.* 1982). Recently, we have also shown that oral exposure of 20- and 40-day-old male rats to DEHP elevates their serum level of ACTH (Supornsilchai *et al.* 2007).

Although the question as to whether DEHP and/or its metabolites can penetrate the blood–brain barrier remains to be resolved, it seems possible that 2-EHXA, which is a relatively small molecule, can do so. This idea is supported by the fact that the (*R*)-enantiomer of sodium 2-ethylhexanoate is capable of penetrating the placental barrier and exerting

severe teratogenic effects in mice (Collins *et al.* 1992), as well as by our own data revealing that 2-EHXA directly enhances the sensitivity of the gonadotropes to GnRH.

In conclusion, oral administration of DEHP to immature female rats reduces their plasma levels of progesterone and estradiol and enhances the responsiveness of their pituitary cells to GnRH. Such alterations in the HPG axis, caused by exposure to phthalates from various sources, may have deleterious effects on the health of the developing organism.

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