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Prenatal Phthalate Exposures and Anogenital Distance in Swedish Boys

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BACKGROUND: Phthalates are used as plasticizers in soft polyvinyl chloride (PVC) and in a large number of consumer products. Because of reported health risks, diisobutyl phthalate (DiNP) has been introduced as a replacement for di(2-ethylhexyl) phthalate (DEHP) in soft PVC. This raises concerns because animal data suggest that DiNP may have antiandrogenic properties similar to those of DEHP. The anogenital distance (AGD)—the distance from the anus to the genitals—has been used to assess reproductive toxicity.

OBJECTIVE: The objective of this study was to examine the associations between prenatal phthalate exposure and AGD in Swedish infants.

METHODS: AGD was measured in 196 boys at 21 months of age, and first-trimester urine was analyzed for 10 phthalate metabolites of DEP (diethyl phthalate), DBP (dibutyl phthalate), DEHP, BBzP (benzylbutyl phthalate), as well as DiNP and creatinine. Data on covariates were collected by questionnaires.

RESULTS: The most significant associations were found between the shorter of two AGD measures (anoscrotal distance; AGDas) and DiNP metabolites and strongest for oh-MMeOP [mono-(4-methyl-7-hydroxyloctyl) phthalate] and oxo-MMeOP [mono-(2-ethyl-5-oxohexyl) phthalate]. However, the AGDas reduction was small (4%) in relation to more than an interquartile range increase in DiNP exposure.

CONCLUSIONS: These findings call into question the safety of substituting DiNP for DEHP in soft PVC, particularly because a shorter male AGD has been shown to relate to male genital birth defects in children and impaired reproductive function in adult males and the fact that human levels of DiNP are increasing globally.

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Introduction

Diesters of 1,2-benzenedicarboxylic acid (phthalic acid), commonly referred to as phthalates, belong to a group of chemicals with endocrine-disrupting properties, meaning that they can interfere with our normal hormonal balance, potentially resulting in adverse human health effects. A recent state-of-the-science report from the World Health Organization (WHO) provides new evidence for several human health risks from exposure to phthalates and other endocrine disruptors (EDCs) including reproductive health and sexual development (Bergman et al. 2013).

Phthalates are used as plasticizers in soft polyvinyl chloride (PVC) and found in a large number of commonly used consumer products including food, building materials, plastics, cosmetics, cleaning products, packages, and toys (Bornehag et al. 2005; Dodson et al. 2012; Rudel et al. 2011). Often, the fraction of phthalates in the plastics is as high as $\geq 30\%$. Because they are not covalently bound to the product matrix, phthalates can leach into the surrounding environment. As a result they are routinely found in indoor air (Bergh et al. 2011; Rudel and Perovich 2009) and dust (Abb et al. 2009; Bornehag et al. 2004, 2005; Langer

et al. 2010; Zhang et al. 2013), and food and water (Shi et al. 2012).

Phthalates are ubiquitous in the environment, and humans are exposed by multiple routes (oral, dermal, and inhalation), the pathway varying by phthalate. It has recently been shown, for example, that exposure to di(2-ethylhexyl) phthalate (DEHP) is primarily via food and for children also via mouthing of plastics, whereas exposure to diethyl phthalate (DEP) and benzylbutyl phthalate (BBzP) occurs primarily via dermal and inhalation pathways (Carlstedt et al. 2013; Koch et al. 2013; Wittassek et al. 2011). Robust biomonitoring data show phthalate metabolites in human fluids such as urine, both in children (Langer et al. 2014) and adults (Wittassek et al. 2011), blood (Frederiksen et al. 2010; Wan et al. 2013), and breast milk (Fromme et al. 2011). These compounds have also been found in amniotic fluid, suggesting that they cross the placental barrier and expose the fetus (Jensen et al. 2012).

One complication in studying phthalate exposure is that replacement of phthalates is continuously ongoing as risks are reported for older formulations. One example of this is the introduction of diisobutyl phthalate (DiNP) and more recently diisobutyl

cyclohexane-1,2-dicarboxylate (DINCH) to replace DEHP in soft PVC [European Chemicals Agency (ECHA) 2013]. Recent biomonitoring data show that urinary levels of DEHP metabolites are decreasing while DiNP metabolites are increasing in European (Göen et al. 2011) and American populations (Silva et al. 2013).

Phthalates and reproductive health. The impact of endocrine-disrupting chemicals on male reproductive health has been a research focus for almost 20 years. During that time, multiple studies in laboratory animals (since 1995) (Sharpe et al. 1995) and in humans (since 2005) (Swan et al. 2005) have demonstrated the sensitivity of the developing male reproductive system to several phthalates. Phthalates may reduce the production of androgens by the testis and rodent studies have demonstrated that DEHP and dibutyl phthalate (DBP) disrupt androgen signaling when administered in the critical window for the development of the reproductive tract (MacLeod et al. 2010; van den Driesche et al. 2011). In humans, similar findings suggest that phthalates (e.g., DEHP and DBP) may be related to male reproductive and developmental abnormalities (ECHA 2013).

Anogenital distance and phthalate exposure. Anogenital distance (AGD)—the distance from the anus to the genitals—is a marker that has been used in animal studies to assess reproductive toxicity (U.S. Environmental Protection Agency 1996). AGD is a sexually dimorphic trait that develops *in utero* under androgen control and is 50–100% longer in males than females.

Numerous studies have shown that prenatal phthalate exposure (notably DEHP, DBP, BBzP) shortens male AGD in rodents (Foster 2005; van den Driesche et al. 2011). Only a few human studies have examined prenatal phthalate exposure and AGD. The first to examine this association in humans reported significant inverse relationships

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between male AGD and DEP, DBP, BBzP, and mono-isobutyl phthalate (MiBP) metabolites (Swan et al. 2005); and a later publication with a larger sample size and more powerful statistical methods found associations between male end points and metabolites of DEHP (Swan 2008). Recently a relationship between prenatal DEHP exposure and shorter AGD in male newborns was reported from Japan (Suzuki et al. 2012) and from Mexico (Bustamante-Montes et al. 2013). However, a study from Taiwan found a negative association between prenatal monobutyl phthalate (MBP) exposure and AGD in newborn girls but not in boys (Huang et al. 2009). To date no studies in humans have included metabolites of DiNP.

Aim of the study. Animal data suggest that DiNP may have antiandrogenic properties similar to those of DEHP (Boberg et al. 2011). This raises concerns because DiNP has been introduced to replace DEHP, and consequently DiNP exposure is rapidly increasing in populations globally. Although several studies have examined phthalates and AGD in humans, none has included DiNP. Therefore, in the present study we examined the relationship between first-trimester urinary metabolite concentrations of DEP, DBP, DEHP, and BBzP as well as DiNP in relation to anogenital distance in boys at 21 months of age.

Methods

Description of the SELMA study. The Swedish Environmental Longitudinal, Mother and child, Asthma and allergy (SELMA) study is a prospective birth cohort study in Sweden that includes > 2,000 mother-child pairs followed from early pregnancy (Bornehag et al. 2012). The study aim was to investigate the impacts of early-life exposure to endocrine-disrupting chemicals and other exposures, for multiple outcomes of growth, development, and chronic diseases.

The SELMA cohort was established by recruiting women in the 10th week of pregnancy in the county of Värmland, Sweden, between September 2007 and March 2010. Of 8,394 reported pregnant women, 6,658

were invited to participate in the study. Among the invited women, 2,582 agreed to participate, corresponding to a participating rate of 39%. Of the 4,076 nonparticipants, 2,091 women were invited to complete a nonrespondent questionnaire for us to examine possible selection bias. We found a self-selection bias in the established cohort when compared with the nonparticipant group—participating families smoked less (14% vs. 19%), had more frequent asthma and allergy symptoms in the family (58% vs. 38%), had higher education (university level) among the mothers (51% vs. 36%), and more often lived in single-family houses (67% vs. 60%). However, there was no obvious reason that this selection bias would have an impact on identification of environmental risk factors for health effects. Detailed recruitment procedures, selection criteria, and data relating to selection bias have been reported and discussed elsewhere (Bornehag et al. 2012). Biological samples (blood and urine) have been collected from the 2,582 pregnant women and their children. Information related to lifestyles, socioeconomic status, living conditions, diet, and medical history has been collected using annual questionnaires (Bornehag et al. 2012).

Measurements of AGD in baby boys. Boys born between 1 September 2009 and 20 November 2010 (i.e., those who were < 18 months of age in the SELMA study) ($n = 325$) and their parents were invited to participate in the study; of these, 228 participated (70%). Of these we were able to measure AGD in 225 boys, and of these, 196 children are included in the present analyses. Mean age of the 196 boys was 20.8 months. AGD was measured by two nurses (one pediatric staff nurse and one midwife) from the County Council of Värmland who were trained by research staff from The Infant Development and the Environment Study (TIDES) (Barret et al. 2014). The nurses had no knowledge of the mother's phthalate concentrations. All AGD measurements were made in the baby's home or at a pediatric clinic.

We obtained two measurements of AGD using measurement methods that have been described elsewhere (Sathyannarayana et al. 2010). Briefly, the longer AGD measurement (AGDap) was measured from the center of the anus to the anterior base of the penis, and the shorter (AGDas) from the center of the anus to the posterior base of the scrotum. The infant was placed on his back on a flat surface with his hips relaxed outward and pulled back towards his shoulders by his mother or an assistant. The examiner stood in front of the infant and made three measurements of both long and short AGD with a dial caliper; the coefficient of variation (CV) was calculated for each AGD measure. Only one measurement was made for 12 boys regarding AGDas and for 24 boys regarding AGDap.

Phthalate metabolites in first-trimester urine. Of the 2,582 pregnant women participating in the SELMA study, a first morning urine sample was obtained from 2,356 women (91%) during weeks 9–11 of the pregnancy. The urine samples were stored frozen at -20°C . These samples were analyzed for the 10 phthalate metabolites listed in Table 1. Briefly, 0.2 mL of urine were added with 0.1 mL of ammonium acetate (pH 6.5) and 0.01 mL glucuronidase (*Escherichia coli*) and thereafter incubated at 37°C for 30 min. Then we added 0.05 mL of a 50:50 (vol:vol) water and acetonitrile solution of labeled (^3H or ^{13}C) internal standards of all analyzed compounds. A C18 column (2.1 mm i.d. \times 50 mm; Genesis Lightn; Grace, Deerfield, IL, USA) was used before the injector to reduce the interference of contaminants during the mobile phase. The phthalate metabolites in the samples were separated on a C18 column (1.5 μm , 2.0 mm i.d. \times 30 mm VisionHT; Grace). The mobile phases were water and acetonitrile with 0.08% formic acid. The samples were analyzed on a Shimadzu UFLC system (Shimadzu Corporation, Kyoto, Japan) coupled to a QTRAP5500 triple quadrupole linear ion trap mass spectrometer equipped with a TurboIon Spray source (AB Sciex, Foster City, CA, USA). The samples were analyzed in triplicate, and the

Table 1. Analyzed phthalate metabolites in the urine of 196 pregnant women, their parent compounds including CAS (Chemical Abstracts Service) number, and limit of detection (LOD).

Parent compound	Metabolite	Acronym	CAS no.	LOD (ng/mL)
Diester name (acronym)	CAS no.	Monoester name		
Diethyl phthalate (DEP)	84-66-2	Mono-ethyl phthalate	MEP	2306-33-4
Dibutyl phthalate (DBP)	84-74-2	Mono- <i>n</i> -butyl phthalate	MnBP	131-70-4
Butylbenzyl phthalate (BBzP)	85-68-7	Mono-benzyl phthalate	MBzP	2528-16-7
Di(2-ethylhexyl) phthalate (DEHP)	117-81-7	Mono-(2-ethylhexyl) phthalate	MEHP	4376-20-9
		Mono-(2-ethyl-5-hydroxyhexyl) phthalate	oh-MEHP	40321-99-1
		Mono-(2-ethyl-5-oxohexyl) phthalate	oxo-MEHP	40321-98-0
		Mono-(2-ethyl-5-carboxypentyl) phthalate	cx-MEPP	40809-41-4
Diisonyl phthalate (DiNP)	68515-48-0	Mono-(4-methyl-7-hydroxyoctyl) phthalate	oh-MMeOP	936021-98-6
		Mono-(4-methyl-7-oxo octyl) phthalate	oxo-MMeOP	936022-00-3
		Mono-(4-methyl-7-carboxyheptyl) phthalate	cx-MMeHP	936022-02-5

mean of the two closest were reported. All samples were analyzed in a randomized order. For quality control of the analyses, chemical blanks and two different in-house prepared quality control samples were analyzed in all sample batches. The limit of detection (LOD) was defined as the concentration corresponding to a peak area ratio of three times the standard deviation of the chemical blanks, and is shown in Table 1. The laboratory at Lund University is a reference laboratory for analyses of urinary phthalate metabolites in a European biomonitoring project [Consortium to Perform Human Biomonitoring on a European Scale (COPHES); <http://www.eu-hbm.info/cophes>]. The creatinine concentrations were analyzed according to an enzymatic method described by Mazzuchi et al. (2000). We calculated the sum of DEHP and DiNP metabolites by summing the metabolite concentrations on a molar basis.

Statistical analyses. Urinary levels of phthalate metabolites were log-transformed to normalize distributions and geometric means (GMs) with 95% confidence interval (CIs). Associations between log-transformed phthalate metabolite concentrations and AGD were estimated using a general linear model, and we present the beta coefficients with 95% CI. We also categorized covariate-adjusted AGD (AGDas and AGDap) into “short” (< 25th percentile), “medium” (25th–75th percentile), and “long” (\geq 75th percentile) as a reference. We then used logistic regression models to estimate the odds ratio (OR) of having a boy with a short (adjusted) AGD compared with a long AGD as a function of log-transformed concentrations of phthalate metabolites in prenatal urine. Finally we examined the ORs for having a short AGD compared with a long AGD as a function of quartiles of phthalate metabolite concentration in the pregnant women’s urine, with the lowest quartile as a reference. Therefore, using these models we examined the relation between prenatal phthalate exposure and AGD in three ways: *a*) both AGD and phthalate metabolite concentrations modeled as continuous variables; *b*) categorical AGDs vs. continuous phthalate metabolite concentrations; and *c*) both AGD and phthalate metabolite concentrations modeled as categorical variables (Figure 1).

The literature indicates that both age and body size can influence measures of AGD. Thus, we controlled for body size by using weight-for-age percentiles by the use of WHO standards (WHO 2009). Weight-for-age percentiles are correlated with age, but much less correlated than weight and age (Swan 2008). Finally, in addition to weight-for-age, all models were adjusted for the boy’s age at examination (months), gestational week of urine sampling, and urinary

creatinine concentration. A *p*-value < 0.05 was considered statistically significant.

Ethical approval. The research ethics committee at Uppsala University, Sweden, approved the study, and informed consent was obtained from all participating adults.

Results

Study population. The study population for this analysis was quite similar to the entire SELMA study population. The 196 families included in the final analysis were somewhat better educated, smoked less, and were more likely to live in a single-family house and in a rural setting (Table 2).

AGD. AGD was measured on 225 boys (19–21 months of age) between June 2011 and August 2012, and prenatal phthalates metabolites were available for 199 of them. Of these, one child was excluded due to an error in weight measurements and two children were excluded due to a CV for the AGD measurements $> 10\%$, resulting in a study population of 196 children. These 196 boys had a mean age of 20.8 months

and a mean weight of 12.6 kg. In addition, AGDas was not measured on four of these boys, so for those analyses $n = 192$ (Table 3). The median AGDas was 40.7 mm [interquartile range (IQR) = 37.8–45.0 mm] while the median AGDap was 82.6 mm (IQR = 78.0–87.4 mm). The mean CV was 2.3% for AGDas and 1.6% for AGDap (Table 3).

Phthalate metabolites. Ten phthalate metabolites and creatinine were measured in urine collected during weeks 9–11. The GM and 95% CI for the concentration of the 10 metabolites as well as the sum of DEHP and DiNP metabolites are presented in Table 4. All phthalate metabolites were identified above LOD in all urine samples (Table 1). The distribution of the metabolite concentrations in the AGD study population was comparable with the distribution in the entire SELMA cohort ($n = 2,356$) (data not shown). Table 4 also provides unpublished data reflecting urinary levels of phthalate metabolites in pregnant women of boys and girls from the Study for Future Families (SFF), whereas published data reflect boys only (Swan 2008;

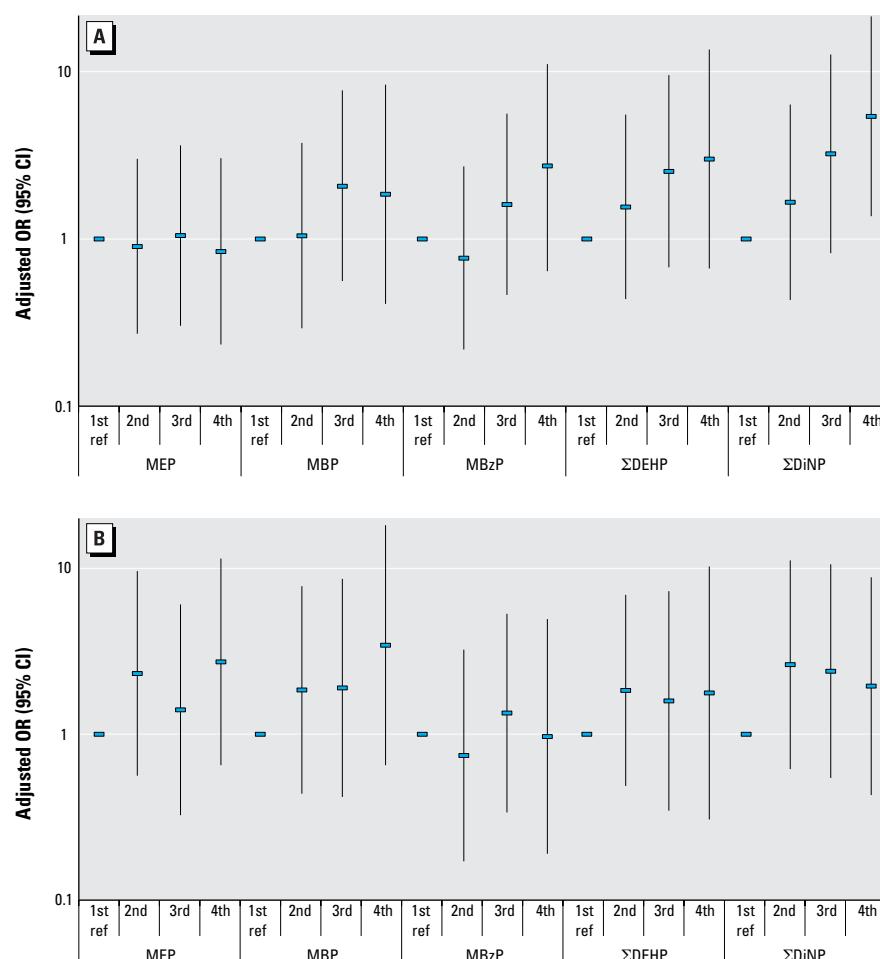


Figure 1. Adjusted odds ratio (OR) for having a short AGD (in the 1st quartile) compared with a long AGD (4th quartile) as a function of quartile of phthalate metabolite concentration in the pregnant women’s urine: (A) AGDas; (B) AGDap. ref, reference (1st quartile). Adjusted for age (months), gestational week of urine sampling, weight-for-age percentile, and creatinine.

Swan et al. 2005), as well as from the U.S. National Health and Nutrition Examination Survey (NHANES) study where data are from women of childbearing age [Centers for Disease Control and Prevention (CDC) 2014].

AGD in relation to phthalate exposure. Most of the phthalate metabolites were negatively associated with AGD both before and after adjustment for covariates in multiple

linear regression models; however, most of the associations did not reach significance. Strongest and most significant inverse associations were found between AGDas and DiNP metabolites and most strongly for oh-MMeOP [mono-(4-methyl-7-hydroxyoctyl) phthalate] and oxo-MMeOP [mono-(2-ethyl-5-oxohexyl) phthalate] and the sum of DiNP metabolites (Table 5).

Table 2. Characteristics of the study population of 196 boys and the families in the entire SELMA study (%).

Characteristic	Present study (n = 196)	SELMA ^a (n = 1,974)
Surrounding for the home		
Urban	16.8	19.3
Suburban	47.1	47.9
Rural	36.1	32.9
Type of home		
Single-family house	64.2	60.8
Row house	7.1	7.9
Apartment in multifamily house	26.4	29.5
Other	2.4	1.8
Education for the mother (highest)		
Elementary school	2.8	3.9
High school (Gymnasium)	33.5	36.0
University	58.5	54.0
Other	5.2	6.0
Smoking in family		
Any smoker in the family	16.0	18.4

^aData from a questionnaire at week 10 of pregnancy when urine samples were collected from the pregnant women (n = 1,974).

Table 3. Measurements of the 196 boys including AGD and other characteristics.

Characteristic	Mean \pm SD	Percentile		
		25th	50th	75th
Age of child (months)	20.8 \pm 1.6	19.7	20.7	21.8
Weight of child (kg)	12.6 \pm 1.3	11.7	12.6	13.3
Gestational week for urine sampling	10.0 \pm 2.3	9.0	10.0	11.0
AGD (mm)				
AGDas	41.4 \pm 5.98	37.8	40.7	45.0
AGDap (n = 192)	82.2 \pm 6.94	78.0	82.6	87.4
CV (%) ^a				
AGDas (n = 184)	2.28	1.03	1.59	2.91
AGDap (n = 168)	1.58	0.66	1.13	1.84

^aFor 12 boys only one measurement was made for AGDas, and for 24 boys only one AGDap measurement was made.

Table 4. Distribution of urinary levels of phthalate metabolites (ng/mL) in 196 women in the present study and in two American studies, SFF and NHANES.

Phthalate	Metabolite	Present study, 2008–2009 (1st trimester) (n = 196)			SFF, ^a 2000–2003 (2nd trimester) (n = 380)	NHANES, ^b 2005–2010 (women 20–40 years) (n = 1,069)	
		GM (95% CI)	25th	50th	75th		
DEP	MEP	63.64 (54.35, 74.52)	30.67	60.56	134.12	149.00 (124.66, 178.10)	100.99 (89.28, 114.24)
DBP	MBP	67.62 (60.02, 76.17)	43.13	66.03	111.78	15.04 (13.49, 16.77)	18.98 (17.34, 20.77)
BBzP	MBzP	15.99 (13.50, 18.95)	7.85	15.08	35.58	9.97 (8.64, 11.50)	7.37 (6.63, 8.19)
DEHP	MEHP	3.27 (2.87, 3.73)	1.91	3.30	5.86	3.23 (2.82, 3.70)	2.61 (2.31, 2.95)
	oh-MEHP	14.42 (12.82, 16.22)	8.69	15.28	22.85	11.74 (10.28, 13.40)	18.30 (16.14, 20.73)
	oxo-MEHP	9.68 (8.59, 10.91)	5.67	9.99	15.60	10.51 (9.26, 11.93)	11.67 (10.36, 13.14)
	cx-MEPP	14.21 (12.63, 15.98)	8.00	14.53	22.50	19.97 (16.45, 24.24)	28.78 (25.71, 32.21)
	Σ DEHP ^c	142.61 (126.99, 160.16)	84.56	148.13	220.71	118.33 (104.37, 134.17)	—
DiNP	oh-MMeOP	6.81 (5.60, 8.28)	2.83	6.27	14.23	—	—
	oxo-MMeOP	3.05 (2.56, 3.63)	1.33	2.75	6.22	—	—
	cx-MMeHP	10.81 (9.16, 12.75)	4.95	8.26	16.43	—	—
	Σ DiNP ^c	67.74 (57.00, 80.52)	28.28	55.91	124.92	—	—
Creatinine (mmol/L)	—	9.55 (8.94, 10.20)	7.00	9.60	13.10	—	—

^aUnpublished data from mothers of boys and girls; published data reflect boys only (Swan 2008; Swan et al. 2005). ^bBased on pooled 2005–2010 NHANES cycles, and women 20–40 years of age (CDC 2014). ^cThe unit is nmol/L.

These associations were also seen when AGD was stratified and phthalate concentrations were compared by adjusted AGD quartile (Table 6). The ORs for having an AGDas in the 2nd–3rd quartile when compared with the 4th quartile as a function of (log-transformed) DiNP metabolite concentrations were in the range of 1.1–1.2 but nonsignificant, whereas the ORs for having an AGDas in the 1st quartile when compared with the 4th quartile was in the range of 2.6–3.1 and significant. We also saw weaker inverse associations between AGDap and DiNP metabolites, which did not reach statistical significance. Associations between prenatal DEHP metabolite exposure and shorter AGDas were also seen, though these associations did not reach statistical significance.

Finally, the odds of having a short AGDas (in the 1st quartile) when compared with a long AGDas (4th quartile) as a function of quartile of phthalate metabolite concentration showed a linear dose–response relationship for prenatal DiNP metabolite exposure (Figure 1A). Similar but somewhat weaker associations were seen between AGDas and DEHP metabolites, as well as MBzP metabolites, whereas no associations were seen with MEP and MBP metabolites. No consistent patterns were seen between AGDap and any of the phthalate metabolites (Figure 1B).

Discussion

Our finding that prenatal phthalate exposure is associated with shorter male AGD is consistent with several earlier studies. However, Swan (2008) found associations between a shorter male AGD and prenatal exposure, particularly for DEHP metabolites, in mothers recruited in 2000–2003, whereas we saw a stronger association with DiNP metabolites. Mothers of the 196 boys included in the current analysis were pregnant in 2009–2010. Phthalates in commerce changed

considerably between 2000 and 2010. Since then, DiNP has largely replaced DEHP in soft PVC applications such as flooring materials, and was measurable in 100% of the current mothers. Additionally, in the present study we observed associations primarily with the shorter AGD measure (AGDas) whereas in the SFF study associations were stronger for AGDap (Swan 2008)—a difference that may be attributable partly to subtle differences in measurement methods between the two studies, but is largely unexplained.

The inverse association between prenatal DiNP exposure and AGDas was significant, as expressed in Table 5, but the AGDas reduction was small in relation to the quite large increase in exposure. For example, more than an IQR increase in DiNP exposure (Table 4) was related to a 4% (1.69 mm) reduction in AGDas in the model (Table 5). About the same effect size has been found for DEHP exposure and AGD reduction in the SFF study (Swan 2008). However, such small relative differences have been associated with male genital birth defects and impaired reproductive function in adult males, as described below.

Our finding of an association between AGD and prenatal exposure to DiNP, and

to a lesser extent to DEHP (nonsignificant), is not unexpected. DEHP is a known antiandrogen (Foster 2005; Gray et al. 2000), and DiNP may be as well. Prenatal DiNP reduces testosterone production in Harlan rat pups (Hannas et al. 2011), although it is reported to be a less potent antiandrogen than DEHP. Furthermore, DiNP exposure in rats during gestation and perinatally increased the incidence of reproductive malformations in male offspring and caused alterations in fetal testicular testosterone production (Borch et al. 2004; Gray et al. 2000). In addition, DiNP is an analog of DEHP—the reason for replacing DEHP with DiNP in soft PVC is that the two molecules have common chemical properties, which may also include antiandrogenic action. Our finding is also in line with some animal data. Prenatal exposure to DiNP has been found to reduce anogenital distance in male offspring rats in three studies; in Wistar rats (Boberg et al. 2011), in Wistar Imamichi rats (Lee et al. 2006), and in Sprague-Dawley rats (Ostby et al. 2001). In another study by Clewell et al. (2013b), AGD was reported to be reduced, but only on postnatal day 14 and at the highest exposure level. Boberg et al. (2011) found similar dose-related

effects of DiNP as those previously shown for DEHP and DBP. They stated that their finding supports the conclusion that DiNP has antiandrogenic properties and is therefore a reproductive toxicant. However, they also stated that DiNP appears to be less potent than DEHP and DBP. On the other hand, in three other studies no association was found between prenatal DiNP exposure and AGD in offspring rats (Clewell et al. 2013a; Gray et al. 2000; Masutomi et al. 2004).

AGD measurements in SELMA boys were longer than those in SFF (Swan 2008; Swan et al. 2005). Median AGDap was 82.6 mm in SELMA and 70.2 mm in SFF; median AGDas was 40.7 mm in SELMA and 36.8 mm in SFF. These differences are at least partly attributable to difference in age at examination; boys in SELMA averaged 20.8 months of age compared with 12.8 months in SFF. The other three studies reporting associations between prenatal phthalate exposure and AGD in offspring children are not comparable with ours because AGD in those studies was measured at birth (Bustamante-Montes et al. 2013; Sathyarayana et al. 2010; Suzuki et al. 2012).

Prenatal urinary levels of phthalate metabolites in the 196 mothers were comparable with those in the entire SELMA cohort, suggesting that there was no substantial selection bias (data not shown). The urinary levels of most phthalate metabolites are comparable with those in SFF and NHANES, but there are differences. MBP metabolite levels are higher in Swedish pregnant women (SELMA), whereas MEP is higher in pregnant women in the United States (SFF, NHANES). MBzP (monobenzyl phthalate) metabolites levels are slightly higher in Sweden whereas the picture for DEHP metabolites is less clear, with both higher and lower levels in Swedish data (Table 4). There are no data for DiNP metabolites in SFF. However, based on more recent data for the DiNP metabolite cx-MMeHP

Table 5. Association between AGD in boys and log-transformed concentrations of phthalate metabolites in prenatal urine from an adjusted^a linear regression model.

Phthalate	Metabolite	AGDas		AGDap	
		β (95% CI)	p-Value	β (95% CI)	p-Value
DBP	MBP	-1.41 (-4.39, 1.57)	0.351	-2.06 (-5.29, 1.18)	0.211
DEP	MEP	0.63 (-1.29, 2.54)	0.518	-1.30 (-3.40, 0.81)	0.225
BBzP	MBzP	-1.66 (-3.56, 0.25)	0.088	-0.65 (-2.74, 1.44)	0.542
DEHP	MEHP	-1.28 (-3.74, 1.17)	0.304	-1.74 (-4.43, 0.95)	0.203
	oh-MEHP	-1.24 (-3.99, 1.51)	0.374	-1.50 (-4.50, 1.49)	0.324
	oxo-MEHP	-0.77 (-3.48, 1.94)	0.576	-1.25 (-4.19, 1.70)	0.406
	cx-MEPP	-0.89 (-3.69, 1.92)	0.534	-0.64 (-3.69, 2.40)	0.677
	Σ DEHP	-1.16 (-4.01, 1.68)	0.420	-1.39 (-4.49, 1.70)	0.375
DiNP	oh-MMeOP	-1.61 (-3.06, -0.16)	0.029	-1.23 (-2.83, 0.37)	0.131
	oxo-MMeOP	-1.82 (-3.47, -0.17)	0.031	-1.67 (-3.49, 0.15)	0.072
	cx-MMeHP	-1.51 (-3.26, 0.24)	0.091	-1.39 (-3.32, 0.53)	0.156
	Σ DiNP	-1.69 (-3.35, -0.02)	0.047	-1.46 (-3.29, 0.38)	0.119

^aAdjusted for age (months), gestational week of urine sampling, weight-for-age percentile, and creatinine.

Table 6. Association between AGD (short, medium, and long) and log-transformed concentrations of phthalate metabolites in prenatal urine calculated in an adjusted^a logistic regression model [OR (95% CI)].

Phthalate	Metabolite	AGDas			AGDap		
		Short 1st quartile	Medium 2nd–3rd quartile	Long 4th quartile (ref)	Short 1st quartile	Medium 2nd–3rd quartile	Long 4th quartile (ref)
DBP	MBP	1.84 (0.43, 7.91)	0.73 (0.21, 2.60)	1.0	1.83 (0.37, 9.02)	0.84 (0.23, 3.01)	1.0
DEP	MEP	1.06 (0.47, 2.40)	0.84 (0.32, 2.16)	1.0	2.23 (0.80, 6.24)	1.11 (0.47, 2.62)	1.0
BBzP	MBzP	1.92 (0.74, 5.00)	0.75 (0.33, 1.79)	1.0	1.49 (0.53, 4.17)	1.21 (0.52, 2.81)	1.0
DEHP	MEHP	1.71 (0.52, 5.64)	1.04 (0.37, 2.93)	1.0	1.98 (0.51, 7.66)	4.01 (1.32, 12.20)	1.0
	oh-MEHP	1.85 (0.50, 6.95)	0.89 (0.27, 2.92)	1.0	2.62 (0.58, 11.78)	3.05 (0.91, 10.16)	1.0
	oxo-MEHP	1.47 (0.40, 5.37)	0.72 (0.22, 2.32)	1.0	2.23 (0.50, 9.87)	3.05 (0.93, 10.07)	1.0
	cx-MEPP	1.76 (0.46, 6.70)	0.98 (0.30, 3.28)	1.0	1.76 (0.38, 8.27)	2.80 (0.82, 9.57)	1.0
	Σ DEHP	1.82 (0.47, 7.06)	0.88 (0.26, 3.00)	1.0	2.44 (0.51, 11.72)	3.36 (0.96, 11.82)	1.0
DiNP	oh-MMeOP	2.61 (1.24, 5.48)	1.08 (0.55, 2.09)	1.0	1.47 (0.67, 3.22)	1.17 (0.61, 2.24)	1.0
	oxo-MMeOP	2.99 (1.28, 7.00)	1.17 (0.54, 2.53)	1.0	1.73 (0.70, 4.23)	1.28 (0.61, 2.70)	1.0
	cx-MMeHP	3.11 (1.27, 7.66)	1.21 (0.52, 2.78)	1.0	1.77 (0.67, 4.65)	1.41 (0.62, 3.21)	1.0
	Σ DiNP	3.02 (1.28, 7.09)	1.16 (0.52, 2.56)	1.0	1.69 (0.68, 4.20)	1.32 (0.61, 2.84)	1.0

ref, reference.

^aAdjusted for age (months), gestational week for urine sampling, weight for age percentile, and creatinine.

[mono-(4-methyl-7-carboxyheptyl) phthalate], concentrations in pregnant women in Sweden and the United States were similar in 2008 (Silva et al. 2013).

The associations we observed between prenatal DiNP metabolites and a shorter AGD in baby boys raises concern for at least two reasons. First, anogenital distance has been shown to be associated with adverse health effects in humans. Three studies reported that male infants and boys with hypospadias or undescended testis had reduced AGD (Hsieh et al. 2012; Jain and Singal 2013; Thankamony et al. 2014). Moreover, a shorter AGD in adult men has been related to decreased fertility (Eisenberg et al. 2011), impaired semen quality (Mendiola et al. 2011), and lower serum testosterone levels (Eisenberg et al. 2012). Shortened AGD has also been suggested as a biomarker of testicular dysgenesis syndrome (Sharpe 2005). Second, the use of DiNP is increasing as it is used to replace DEHP in soft PVC, and consequently the human urinary levels of DiNP metabolites are rapidly increasing globally. The total global market for phthalates has been estimated by the ECHA to be about 6 million tons, with 1.4 million tons in Europe, the Middle East, and Africa, 1.1 million tons in the Americas, and 3.5 million tons in Asia, and phthalates represent 84% of the global plasticizer market (ECHA 2013). Of the global market of plasticizers, DiNP and DiDP (diisodecyl phthalate) were reported to represent about 32% (ECHA 2013). Data from the American Chemistry Council (2012) indicate that the annual world production of DiNP was estimated to be about 1.5 million tons in 2013, assuming a production growth of 2.5% during the last years. Biomonitoring data from Europe and the United States also show that human urinary levels of DEHP metabolites are decreasing while DiNP metabolite concentrations are increasing (Göen et al. 2011; Silva et al. 2013).

Conclusions

The use of DiNP in soft PVC is increasing, and biomonitoring data show that human DiNP metabolite levels are rapidly increasing globally. Our data suggest that this substitute phthalate may not be safer than the chemical it is replacing. We find that DiNP is associated with a shorter AGD in boys at the age of 21 months, which is of concern because AGD has been related to male genital birth defects and impaired reproductive function in adult males.

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