human patients with allergic asthma following endobronchial allergen challenge, and genetic linkage studies showing an association between functional SNPs in the S1P receptor-1 and asthma. Thus, whether increased expression of ORMDL3 in hORMDL3<sup>3p3-Cre</sup> mice increases AHR in the absence of airway inflammation through reductions in sphingolipid levels is supported by some, but not all, studies of sphingolipids.

Interestingly, in Sptlc2<sup>−/−</sup> mice, the reduced synthesis of sphinganine and ceramide was associated with increases in AHR in the absence of inflammation, a phenotype we have observed in hORMDL3<sup>3p3-Cre</sup> mice. However, there are differences in the profile of sphingolipids reduced in hORMDL3<sup>3p3-Cre</sup> mice compared with Sptlc2<sup>−/−</sup> mice (sphinganine, sphingosine, S1P, ceramide in serum; sphinganine in lung) compared with Sptlc2<sup>−/−</sup> mice (sphinganine, ceramide in lung). In addition, hORMDL3<sup>3p3-Cre</sup> mice develop spontaneous airway remodeling and mucus expression not observed in Sptlc2<sup>−/−</sup> mice, suggesting that pathways to remodeling and mucus are independent of the ability of ORMDL3 to inhibit SPT and sphingolipid synthesis.

Studies have also examined the role of ceramide in mouse asthma models and demonstrated that allergen challenge increased lung levels of ceramide in WT mice, whereas pretreatment with pharmacologic inhibitors of ceramide reduced AHR, lung eosinophils, and Th2 cytokines. These studies suggest that the reduction in serum ceramide levels we have noted in hORMDL3<sup>3p3-Cre</sup> mice can contribute to an increased risk of allergy development including asthma and atopic dermatitis. In a recent study, an inducible lung epithelial gene regulating metalloproteases, chemokines, OAS, and ATP6. Proc Natl Acad Sci USA 2012;109:16648-53.

Prenatal phthalate exposure associates with low regulatory T-cell numbers and atopic dermatitis in early childhood: Results from the LINA mother-child study

To the Editor:

Phthalates serve as binders and plasticizers in everyday items, including cosmetics, household cleaners, food packaging, personal-care products, toys, and many other consumer products. Recent publications show that exposure to these chemicals may contribute to an increased risk of allergy development including asthma and atopic dermatitis. An immune modulatory capacity of these compounds is assumed but not fully elucidated by now. Epidemiological studies provide evidence that exposure to phthalates and their metabolites might be more critical in the prenatal period when the fetal immune system is developing. According to the fact that regulatory T (Treg) cells are key players in the modulation of immune responses, we asked whether phthalates may affect the number of these cells leading to the observed increased risk to develop atopic dermatitis. In a recent study, environmental factors influence the outcome of immune responses and the development of atopic dermatitis. The study aimed to assess the associations between prenatal phthalate exposure and the number of regulatory T cells in the maternal circulation, as measured by T<sub>reg</sub> cell counts.

In summary, in this study we have demonstrated that hORMDL3<sup>3p3-Cre</sup> mice had significantly reduced serum levels of the pathway of sphingolipids regulated by SPT (sphinganine, ceramide, sphingosine, and S1P), as well as reduced lung levels of sphinganine. In addition, we demonstrate that administration of S1P to naive hORMDL3<sup>3p3-Cre</sup> mice further increases their AHR associated with increased levels of peribronchial macrophages. These in vivo studies extend previous in vitro observations that ORMDL3 inhibits the generation of sphingolipids including ceramide and S1P. Previous studies have also demonstrated that increased ORMDL3 activates the ATF6α pathway of the endoplasmic reticulum unfolded protein response and that this regulates levels of SERC2b, which can contribute to AHR. In addition, ORMDL3 regulates levels of remodeling genes (TGF-β1, ADAM8) as well as CC and CXC chemokines implicated in asthma. Further studies are needed to determine which ORMDL3-regulated pathway (eg, inhibition of sphingolipid synthesis, inhibition of ATF6α and SERCA2b, inhibition of remodeling genes, or other as yet unidentified pathways) could contribute to increased AHR observed in hORMDL3<sup>3p3-Cre</sup> mice, and whether in patients with asthma SNPs associated with increased ORMDL3 expression result in reduced sphingolipid levels. Such insights are important in understanding why ORMDL3 on chromosome 17q21 is highly linked to human asthma.

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Peter Rosenthal, BS
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Ruth Gordillo, PhD
David H. Broide, MB, ChB

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we demonstrated that low number of Treg cells at birth is associated with a higher risk to develop an atopic dermatitis within the first 3 years of life.\(^5\) In the present investigation, we aimed to evaluate whether a high maternal phthalate burden impacts the number of Treg cells in pregnancy as well as in early childhood. Associations to children’s atopic dermatitis were found with the primary metabolites of diethyl, di-n-butyl, butylbenzyl, or di-2-ethylhexyl phthalate.\(^2\) Therefore, in our study, we measured the concentration of these metabolites (monoethyl phthalate [MEP], monoisobutyl phthalate [MiBP], mono-n-butyl phthalate [MnBP], monobenzyl phthalate [MBzP], mono-(2-ethylhexyl) phthalate [MEHP]) in maternal urine and assessed the relationship to Treg-cell numbers and the development of atopic dermatitis as well as to hay fever and allergic sensitization to food (fx5) and inhalant (sx1) allergens.

Data were gained from our prospective birth cohort study LINA (Lifestyle and environmental factors and their Influence on Newborns Allergy risk). For this study, 629 mother-child pairs (622 mothers, 7 twin pairs) were recruited from March 2006 until December 2008 in Leipzig, Germany.\(^6\) Annually, around the birthday of the child, follow-up investigations were performed with questionnaire evaluations for allergic outcomes (atopic dermatitis, hay fever, asthma) and confounders as well as clinical visits including blood collection. Early morning urine samples were collected during the third trimester (gestational age 36 weeks, n = 610) and analyzed for the primary phthalate metabolites (MEP, MiBP, MnBP, MBzP, and MEHP) using a multianalyte procedure as described by Feltens et al.\(^7\) Absolute concentrations of phthalates were calculated on the basis of calibration curves and normalized to urinary creatinine concentrations. The number of Treg cells was determined by FOXP3 methylation-specific real-time PCR in the Treg-cell–specific demethylated region in blood samples from pregnancy (34th gestational week, n = 607), cord blood (n = 448), and from the second year of children’s life (n = 331) as described previously.\(^6\) This method enables the identification of cells with stable Treg-cell phenotype and function.\(^8\) Characteristics of the study population, detailed method description, and median values and interquartile ranges for all phthalate metabolites and Treg-cell numbers are given in this article’s Online Repository (Tables E2 and E3 at www.jacionline.org).

Associations between maternal urine phthalate metabolite concentrations and the number of Treg cells were calculated using a linear regression model adjusted for maternal atopic dermatitis, maternal smoking and/or ETS exposure at home, maternal education, cat ownership during pregnancy, and previous births. In addition, data from cord blood were adjusted for sex and those at the age of 2 years for sex and breast-feeding until 6 months. The number of Treg cells was determined by FOXP3 methylation-specific real-time PCR in the Treg-cell–specific demethylated region in blood samples from pregnancy (34th gestational week, n = 607), cord blood (n = 448), and from the second year of children’s life (n = 331) as described previously.\(^6\) This method enables the identification of cells with stable Treg-cell phenotype and function.\(^8\) Characteristics of the study population, detailed method description, and median values and interquartile ranges for all phthalate metabolites and Treg-cell numbers are given in this article’s Online Repository (Tables E2 and E3 at www.jacionline.org).

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TABLE I. Concentrations of phthalate metabolites in maternal urine (36th gestational week) in regard to the development of atopic dermatitis (physician diagnosed) in the first 3 years of children’s life

<table>
<thead>
<tr>
<th>Phthalate metabolites</th>
<th>Atopic dermatitis 0-3rd year, median (IQR) (ng/mg)*</th>
<th>adjusted mean ratios</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td>MEP</td>
<td>60 (28.3-143.9)§ 50.5 (23.9-99.4) .155</td>
<td></td>
<td></td>
</tr>
<tr>
<td>MiBP</td>
<td>73.3 (53.2-102.9)§ 62.4 (45.7-90.9) .036§</td>
<td></td>
<td></td>
</tr>
<tr>
<td>MnBP</td>
<td>112.7 (78.8-109.7) 96.9 (66.7-143.9) .057</td>
<td></td>
<td></td>
</tr>
<tr>
<td>MBzP</td>
<td>6.9 (3.7-11.7) 6.4 (3.7-10.6) .646</td>
<td></td>
<td></td>
</tr>
<tr>
<td>MEHP</td>
<td>7.7 (5.5-11.5) 7.2 (5-10.8) .438</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

\*Concentrations were adjusted for creatinine (ng per mg of creatinine) to control for urine dilution.  
§P values from Mann-Whitney U test.  
\(|b|Cases with maternal phthalate measurements in maternal urine (36th gestational week).  
§§Significant values are marked in boldface (P < .05).
CI, 0.82-1, P = .046 and aMR, 0.91, 95% CI, 0.82-1, P = .047, respectively) and in children aged 2 years (aMR, 0.85, 95% CI, 0.76-0.96, P = .007 and aMR, 0.88, 95% CI, 0.78-0.99, P = .045, respectively) but not in mothers during pregnancy (aMR, 1.02, 95% CI, 0.94-1.12, P = .529 and aMR, 1.07, 95% CI, 0.98-1.17, P = .095, respectively) (Fig 1). We repeated the analysis on the basis of a subgroup of 208 children with available data for maternal urine phthalate concentrations and Treg-cell numbers during pregnancy, as well as Treg-cell measurements in cord blood and at the age of 2 years and gained similar results. In addition here, at the age of 2 years, a reduced number of Treg cells was observed in association with high concentrations of MnBP during pregnancy (see Fig E1 in this article’s Online Repository at www.jacionline.org). Although cord blood Treg-cell numbers are only affected by maternal exposure during pregnancy, we cannot exclude that phthalate exposure after birth also contributed to the observed low Treg-cell numbers at the age of 2 years.

Furthermore, the impact of high phthalate metabolite levels in pregnancy on the development of atopic dermatitis during the first 3 years in children’s life was assessed by calculating odds ratios adjusted for sex, maternal atopic dermatitis, maternal smoking, and/or ETS exposure at home, siblings, maternal education, cat ownership, and breast-feeding until 6 months.

<table>
<thead>
<tr>
<th>Phthalate metabolites</th>
<th>Atopic dermatitis 0-3rd year; % (n/N), 19.5% (435)*</th>
</tr>
</thead>
<tbody>
<tr>
<td>MEP</td>
<td>Crude OR (95% CI) P value Adjusted OR (95% CI) P value</td>
</tr>
<tr>
<td>MiBP</td>
<td>2.15 (1.11-4.14) 0.022 2.21 (1.1-4.45) 0.026</td>
</tr>
<tr>
<td>MnBP</td>
<td>1.79 (0.95-3.37) 0.072 1.79 (0.91-3.52) 0.090</td>
</tr>
<tr>
<td>MBzP</td>
<td>1.29 (0.68-2.47) 0.427 1.28 (0.65-2.52) 0.470</td>
</tr>
<tr>
<td>MEHP</td>
<td>1.47 (0.77-2.79) 0.242 1.5 (0.76-2.98) 0.238</td>
</tr>
</tbody>
</table>

*Cases with maternal phthalate measurements in maternal urine (36th gestational week).
†Analysis was performed using a logistic regression model with phthalate values categorized into quartiles; odds ratios (ORs) were adjusted for sex, maternal atopic dermatitis, maternal smoking, and/or ETS exposure at home, siblings, maternal education, cat ownership, and breast-feeding until 6 months.
‡Significant values are marked in boldface (P < .05).

In addition to the above-described link between MiBP and atopic dermatitis, we could show that exposure to high concentrations of MEP and MnBP during the fetal period was associated with an increased risk to sensitization to food allergens (fx5) at the age of 2 years and in trend also at the age of 1 year (see Table E4 in this article’s Online Repository at www.jacionline.org). We furthermore could demonstrate that a high number of Treg cells at birth is protective against sensitization to food allergens at the age of 1 year and a high number of Treg cells at the age of 2 years was associated with a lower risk for sensitization to inhalant allergens at this age (see Table E5 in this article’s Online Repository at www.jacionline.org). As Treg cells are keeping immune responses in balance, lower numbers of these cells may facilitate the development of allergic diseases, as we could show earlier for atopic dermatitis.2,3 Interestingly, maternal Treg-cell numbers in pregnancy were not associated with phthalate metabolite concentrations, pointing out that the immature immune system of the fetus and toddler is more vulnerable to these chemicals. Although we are not able to provide a mechanistic explanation for the impact of MEP and MiBP on Treg-cell numbers (which might implicate the differentiation and/or proliferation of these cells), our epidemiological findings may encourage for the performance of in vitro assays in this direction in future studies. Taken together, our data suggest that MiBP being associated with reduced numbers of Treg cells may facilitate the development of an atopic dermatitis in early childhood. Because DiBP, the parent compound from which MiBP originates, is widely used in food packages, adhesives, and cosmetics, the usage of these products especially during pregnancy should be avoided.
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1These authors contributed equally to this work.

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Dysregulation of lipidomic profile and antiviral immunity in response to hyaluronan in patients with severe asthma

To the Editor:

Features of patients with severe asthma include a greater frequency and severity of hospitalizations caused by pneumonia, severe influenza, and sinopulmonary infections. Viral infections are frequent triggers of asthma exacerbations. Impaired antiviral responses in asthmatic patients have been noted. However, the mechanisms of this phenomenon are not well understood.

The asthmatic airway wall undergoes many alterations, including increased and changed deposition of extracellular matrix. Hyaluronan (HA), a major component of extracellular matrix, accumulates in the lung and serum of asthmatic patients and correlates with disease severity. Low-molecular-weight (LMW) forms of HA generated during tissue injury or inflammation have been linked to asthma, but the mechanisms of that link are not well understood.

Recently, we described the mechanism by which LMW HA can activate cytosolic phospholipase A2 (cPLA2) and arachidonic acid (AA) production. Previously, we reported increased expression of cPLA2 in PBMCs of patients with severe asthma. cPLA2 is a rate-limiting enzyme in eicosanoid production, which is responsible for liberation of AA from cellular membranes. AA is the precursor of leukotrienes, prostaglandins (PGs), hydroxyeicosatetraenoic acids (HETEs), thromboxanes, lipoxins, and epoxides, many of which are involved in asthma pathogenesis, and they are altered in patients with viral infections.

A thorough analysis of LMW HA’s effects on the lipidomic profile or global gene expression in asthmatic patients has never been performed, and its possible influence on the disease progression has not been noted. Therefore we performed a systemic analysis of LMW HA signaling in PBMCs of patients with mild-to-moderate and severe asthma by using liquid chromatography and mass spectrometry combined with microarray, real-time PCR, and multiplex protein analyses.

Details of methodology are provided in the Methods section in this article’s Online Repository at www.jacionline.org.

Thirteen asthmatic patients and six control subjects were enrolled under a National Heart, Lung, and Blood Institute review board–approved protocol (99-H-0076). Severe asthma was defined according to European Respiratory Society/American Thoracic Society guidelines. Participants’ demographic and phenotypic characteristics are presented in Tables E1 and E2 in this article’s Online Repository at www.jacionline.org.

Of the 151 lipid species profiled, we detected 68 before or after LMW HA stimulation. At baseline, fewer metabolites were generated through the COX pathway compared with the combined lipoxigenase, CYP450, and nonenzymatic pathways in each group (Fig 1A, and see Fig E1A, in this article’s Online Repository at www.jacionline.org). The relative percentage of all COX metabolites at baseline was significantly lower in patients with severe asthma than in control subjects (Fig 1A, and see Fig E1A). However, after LMW HA stimulation, there was a significant increase in the relative percentage of COX-generated mediators and a corresponding decrease of metabolites derived from the other pathways in all groups (Fig 1A, and see Fig E1A).

Analysis of individual lipids revealed that 22 were significantly upregulated by using LMW HA (Fig E1B). Although at baseline several lipid species tended to have lower concentrations in patients with severe asthma, the increase in 9 metabolites after LMW HA treatment was significantly more pronounced in those patients compared with control subjects (Fig 1B). These included the COX metabolites thromboxane B2 (TXB2), prostaglandin E2, D2, and B2 (PGE2, PGD2, and PGB2) and metabolites from other pathways, including 15-hydroxyeicosatetraenoic acid (15-HETE) (through lipoxigenase), 11,12-eicosadienoic acid (11,12-EET), and 14,15-EET (through CYP450), and non-AA metabolites, such as 13-hydroxyoctadecatrienoic acid (13-HODE) and (16)-epoxydocosapentaenoic acid 16(17)-EpDPE. Only in patients with severe asthma did the levels of most of these metabolites reach the range of activation of their cognate receptors, although 15-HETE, 14,15-EET, and 16(17)-EpDPE were significantly upregulated also in patients with mild-to-moderate asthma. Treatment with a specific cPLA2α inhibitor significantly decreased LMW HA-induced release of AA, as well as 15-HETE and 14,15-EET, from PBMCs of asthmatic patients.

http://dx.doi.org/10.1016/j.jaci.2016.09.034
METHODS

Study design

Six-hundred twenty-nine mother-child pairs were recruited within the prospective birth cohort study LINA (Lifestyle and environmental factors and their Influence on Newborns Allergy risk) from March 2006 until December 2008 in Leipzig, Germany. Blood samples were obtained during pregnancy (mother, 34th week of gestation), at birth (venous umbilical cord blood), and annually around the child’s birthday. Early morning urine samples were obtained from mothers in the 36th week of gestation. Data on confounding variables, prenatal exposure, lifestyle factors, and children’s disease outcomes were gained from questionnaires filled in by the parents 4 weeks before birth and annually thereafter. The present investigation comprises of mothers having urine sample analyses and measurement of Treg cells in blood samples as well as children with Treg-cell measurements in cord blood and at the age of 2 years.

Participation was voluntary and informed consent was given by the parents. This study was approved by the Ethics Committees of the University of Leipzig (file references 046-2006, 160-2008).

Quantification of Treg cells

Isolation of genomic DNA. Genomic DNA (gDNA) was isolated from whole blood using the DNA Blood Mini Kit according to manufacturer’s instructions (Qiagen, Hilden, Germany). Briefly, 200 μL whole blood was treated with 20 μL Proteinase K (activity, 6000 μAU/mL) and incubated with Buffer AL in a ratio of 1:1 (10 minutes, 56°C, 350 rpm). Cell lysis was stopped by adding 200 μL ethanol. The complete sample volume was transferred onto a QIAamp Mini spin column and centrifuged (12,000 rpm). DNA bound to the QIAamp membrane was washed twice using Buffers AW1 and AW2 (12,000 rpm, 1 minute; 13,000 rpm, 5 minutes). DNA was eluted by adding Buffer AE and stored at 4°C until subsequent analysis, applying the mean of the 2 reference genes for normalizing.

Bisulfite conversion of gDNA and quantification of Treg cells by FOXP3 methylation-specific real-time PCR. Bisulfite treatment of gDNA was conducted using EpiTect 96 Bisulfite Kit (Qiagen, Hilden, Germany) as described by the manufacturer. Briefly, 1 μg of gDNA, 85 μL of the Bisulfite Mix, and 35 μL of the DNA Protect Buffer were mixed in a 96-well EpiTect Conversion Plate, and bisulphite conversion was performed in a thermal cycler according to manufacturer’s instruction. Converted gDNA was stored at −20°C until further analysis.

Quantification of demethylation in Treg-cell–specific demethylated region was performed by real-time PCR by Epiontis (Berlin, Germany; for details, see data in this article’s Online Repository at www.jacionline.org). As described previously, the number of Treg cells in blood is presented as percentage corresponding to the measured amount of Treg-cell–specific demethylated region demethylation in the FOXP3 gene.

Phthalate metabolite measurement in urinary samples

Phthalate quantification was carried out for 610 early morning maternal urine samples. In the present study, data for 542 mothers, the subset with Treg-cell measurements in their blood, are presented. Urine samples were collected at 36th weeks’ gestation and stored in polypropylene tubes at −80°C until further analysis. Primary phthalate metabolite (MEP, MEBP, MnBP, MBzP, MEHP) quantification was carried out for all samples using a multianalyte procedure as described by Feltens et al.13 Absolute concentrations of phthalate metabolites were calculated on the basis of calibration curves and normalized to urinary creatinine concentrations as previously described.13 Concentrations are given in ng per mg creatinine.

RESULTS

Allergic outcomes

Information on allergic outcomes was collected using questionnaires self-administered by the parents. Atopic dermatitis and hay fever were recorded as physician diagnoses (“Has a doctor diagnosed your child with atopic dermatitis in the last 12 months?”); “Has a doctor diagnosed your child with hay fever in the last 12 months?”). The lifetime prevalence within the first 3 years of life was determined by adding the information of having doctor’s-diagnosed atopic dermatitis or hay fever from each year. The control group was defined including children having never developed symptoms or being diagnosed for this disease until the age of 3 years.

Allergic sensitization was assessed by the measurement of specific IgE against food allergens (fx5) and inhalant allergens (sx1). Samples were categorized as positive by the cutoff value of more than 0.35 kU/L.

IgE measurement

The concentrations of specific IgE against food (fx5) and inhalant allergens (sx1) in sera of 1-year-old children were determined by the Phadia CAP System (Phadia GmbH, Freiburg, Germany). The allergen multipanel fx5 consists of hen’s egg, cow’s milk, wheat, fish, peanut, and soy. sx1 includes timothy, rye, mugwort, birch, house dust mite (Dermatophagoides pteronyssinus), cat, and dog. Samples with a specific IgE concentration of more than 0.35 kU/L were regarded as positive.

Confounding variables

Information on maternal history of atopy and prenatal exposure was assessed by detailed questionnaires answered by parents during the third trimester of pregnancy and annually after delivery. To address maternal atopy history, we included asthma, hay fever, and atopic dermatitis (“Did you ever suffer from asthma, allergic rhinitis, atopic dermatitis?”). Smoking or exposure to environmental tobacco smoke at home during pregnancy was recorded with the question “Did you or anybody else smoke inside your dwelling during the last twelve months?” If yes; occasionally, once per week, or daily. Maternal education was assessed by asking “Which is the highest education level you have?” “low, 9 yr of schooling or less,” “Hauptschulabschluss”; “intermediate, 10 yrs of schooling” “Mittlere Reife”; “high, 12 yrs of schooling or more” “(Fach-)hochschulreife.” Furthermore, the number of present siblings, the presence of pets (including cats) during pregnancy, and the duration of breast-feeding were recorded.

Statistical analysis

Statistical analyses were performed using Statistica for Windows Version 10.0 (StatSoft Inc [Europe], Hamburg, Germany). The chi-square test for cross-relationship was performed to ensure the equal distribution of parameters in the analyzed subgroups and the entire LINA cohort. All P values of less than .05 were considered to be significant. Because measured phthalate metabolite concentrations and Treg-cell numbers were not normally distributed, a logarithmic transformation was performed. Furthermore, the data were categorized into quartiles and used in the regression models. To analyze the relationship between maternal urine phthalate metabolite concentrations from pregnancy and the number of Treg cells in blood samples from pregnancy, cord blood, and at children’s age of 2 years, linear regression models were used. The linear regression models were adjusted for the possible confounding factors maternal atopic dermatitis, maternal smoking and/or ETS exposure at home, maternal education, cat ownership, previous births (presence of older siblings), and also (only for children) for sex and breast-feeding until 6 months. Data are presented as mean ratios, which are the back-transformed effects from the regression model of the logarithmically transformed outcome. Odds ratios were calculated to show the relationship between prenatal phthalate concentrations in maternal urine and the development of atopic dermatitis (physician diagnosed) within the first 3 years of life. The logistic regression models were adjusted for sex, maternal atopic dermatitis, maternal smoking and/or ETS exposure at home, presence of older siblings, maternal education, cat ownership, and breast-feeding until 6 months.

Study population characteristics

Characteristics of the study population are listed in Table E1. There were no differences in the distribution of considered
parameters in the analyzed subgroups compared with the entire LINA cohort. Out of the 622 mothers participating in the study, 542 had urinary samples with phthalate metabolite measurements as well as Treg-cell measurements in blood. Out of the 470 children with cord blood samples, in 448 the number of Treg cells was assessed whereas at the age of 2 years, Treg-cell numbers were measured in 331 out of 339 blood samples. In the analyzed subgroup of children with prenatal maternal urine phthalate measurements, 85 (19.5%) developed an atopic dermatitis until the age of 3 years compared with 92 (18.5%) children from the entire LINA cohort.

REFERENCES


FIG E1. Relationship between maternal urine phthalate metabolite concentrations and the number of Treg cells. Results based on the subgroup of children with available data for maternal urine phthalate concentrations in pregnancy as well as measurements of Treg cells in pregnancy, in children's cord blood, and at the age of 2 years (n = 208). Shown are the mean ratios with 95% CI. Mean ratios were adjusted for maternal atopic dermatitis, maternal smoking and/or ETS exposure at home during pregnancy, maternal education, cat ownership during pregnancy, and previous births. In addition, data from cord blood were adjusted for sex and those at the age of 2 years for sex and breast-feeding until 6 months.
<table>
<thead>
<tr>
<th>Parameters</th>
<th>Entire LINA cohort, n (%), N = 629</th>
<th>Maternal urine samples in pregnancy, n (%), N = 542</th>
<th>Cord blood samples at birth, n (%), N = 448</th>
<th>Blood samples at children’s age of 2 y, n (%), N = 331§</th>
<th>P value*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sex of the child</td>
<td>Male 327 (52)</td>
<td>282 (52)</td>
<td>211 (47.1)</td>
<td>165 (49.8)</td>
<td>.885</td>
</tr>
<tr>
<td></td>
<td>Female 302 (48.0)</td>
<td>260 (48)</td>
<td>237 (52.9)</td>
<td>166 (50.2)</td>
<td>.954</td>
</tr>
<tr>
<td>Maternal history of atopy</td>
<td>No 330 (53.1)</td>
<td>280 (51.7)</td>
<td>242 (54)</td>
<td>166 (50.2)</td>
<td>.947</td>
</tr>
<tr>
<td></td>
<td>Yes 292 (46.9)</td>
<td>262 (48.3)</td>
<td>206 (46)</td>
<td>165 (49.8)</td>
<td>.947</td>
</tr>
<tr>
<td>Maternal atopic dermatitis</td>
<td>No 517 (82.2)</td>
<td>443 (81.7)</td>
<td>373 (83.3)</td>
<td>265 (80.1)</td>
<td>.947</td>
</tr>
<tr>
<td></td>
<td>Yes 111 (17.6)</td>
<td>99 (18.3)</td>
<td>75 (16.7)</td>
<td>66 (19.9)</td>
<td>.947</td>
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<tr>
<td>Maternal education</td>
<td>Low 24 (3.8)</td>
<td>20 (3.7)</td>
<td>13 (2.9)</td>
<td>6 (1.8)</td>
<td>.98</td>
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<td>Intermediate 194 (30.8)</td>
<td>165 (30.4)</td>
<td>143 (31.9)</td>
<td>113 (34.1)</td>
<td>.947</td>
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<tr>
<td></td>
<td>High 411 (65.3)</td>
<td>357 (65.9)</td>
<td>292 (65.2)</td>
<td>212 (64)</td>
<td>.947</td>
</tr>
<tr>
<td>Breast-feeding until 6 mo</td>
<td>No 139 (23.92)</td>
<td>112 (20.7)</td>
<td>102 (22.8)</td>
<td>77 (23.3)</td>
<td>.987</td>
</tr>
<tr>
<td></td>
<td>Yes 442 (76.07)</td>
<td>386 (71.2)</td>
<td>311 (69.4)</td>
<td>242 (73.1)</td>
<td>.987</td>
</tr>
<tr>
<td>Cat ownership during pregnancy</td>
<td>No 516 (82)</td>
<td>443 (81.7)</td>
<td>364 (81.2)</td>
<td>262 (79.2)</td>
<td>.957</td>
</tr>
<tr>
<td></td>
<td>Yes 113 (18)</td>
<td>99 (18.2)</td>
<td>84 (18.8)</td>
<td>69 (20.8)</td>
<td>.957</td>
</tr>
<tr>
<td>Presence of older siblings</td>
<td>No 420 (66.8)</td>
<td>359 (66.2)</td>
<td>298 (66.5)</td>
<td>218 (65.9)</td>
<td>.999</td>
</tr>
<tr>
<td></td>
<td>Yes 209 (33.2)</td>
<td>183 (33.8)</td>
<td>150 (33.5)</td>
<td>113 (34.1)</td>
<td>.999</td>
</tr>
<tr>
<td>Smoking during pregnancy</td>
<td>Never 527 (84.7)</td>
<td>465 (85.8)</td>
<td>383 (85.5)</td>
<td>287 (86.7)</td>
<td>1</td>
</tr>
<tr>
<td></td>
<td>Occasionally 43 (6.9)</td>
<td>37 (6.8)</td>
<td>23 (5.8)</td>
<td>20 (6)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Once per week 4 (0.7)</td>
<td>4 (0.7)</td>
<td>3 (0.7)</td>
<td>1 (0.3)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Daily 48 (7.7)</td>
<td>36 (6.6)</td>
<td>36 (8)</td>
<td>23 (6.9)</td>
<td></td>
</tr>
</tbody>
</table>

Because of missing data, case number may vary for some variables.
*P value from chi-square test for cross-relationship.
†Children with Treg-cell measurements in pregnancy.
‡Children with Treg-cell measurements in cord blood.
§Children with Treg-cell measurements at the age of 2 years.
||History of atopy is defined as occurrence of asthma or atopic dermatitis or hay fever.
¶Low, 9 years of schooling or less “Hauptschulabschluss”; intermediate, 10 years of schooling “Mittlere Reife”; high, 12 years of schooling or more “(Fach-)hochschulreife.”
#Maternal smoking and/or ETS exposure during pregnancy at home.
### TABLE E2. Maternal urine phthalate metabolite concentrations (36th gestational week, n = 542, samples with Treg-cell measurements)

<table>
<thead>
<tr>
<th>Phthalate metabolite</th>
<th>Median (IQR) (ng/mg)*</th>
</tr>
</thead>
<tbody>
<tr>
<td>MEP</td>
<td>50 (24.8-102.3)</td>
</tr>
<tr>
<td>MiBP</td>
<td>66.6 (48.9-99.8)</td>
</tr>
<tr>
<td>MnBP</td>
<td>104.9 (69.9-146.8)</td>
</tr>
<tr>
<td>MBzP</td>
<td>6.7 (3.8-11.6)</td>
</tr>
<tr>
<td>MEHP</td>
<td>7.3 (5.1-11.4)</td>
</tr>
</tbody>
</table>

*IQR, Interquartile range.*

*Concentrations were adjusted for creatinine (in ng per mg of creatinine) to control for urine dilution.*
TABLE E3. Treg-cell numbers in samples with phthalate measurements as detected by means of demethylation in the FOXP3 gene

<table>
<thead>
<tr>
<th>Treg cell</th>
<th>Median (IQR) (%)*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Maternal blood (34th gestational week)</td>
<td>0.91 (0.56-1.34)</td>
</tr>
<tr>
<td>Birth (cord blood)</td>
<td>1.18 (0.75-1.71)</td>
</tr>
<tr>
<td>Children aged 2 y</td>
<td>3.31 (2.49-4.39)</td>
</tr>
</tbody>
</table>

IQR, Interquartile range; TSDR, Treg-cell–specific demethylated region.

*The number of Treg cells is presented as percentage of Treg cells in whole blood corresponding to the measured amount of TSDR demethylation in the FOXP3 gene. The detection limit of the FOXP3 TSDR demethylation assay is 0.03%; all values were above this limit.
TABLE E4. Relationship between maternal urine phthalate metabolite concentrations (36th gestational week) and allergic outcomes in the first 3 years of children’s life

<table>
<thead>
<tr>
<th>Phthalate metabolites</th>
<th>Hay fever</th>
<th>Sensitization to food allergens (fx5)</th>
<th>Sensitization to inhalant allergens (sx1)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>0-3rd year, 3.6% (17 of 475)</td>
<td>First year, 15.3% (79 of 515)</td>
<td>Second year, 12.1% (41 of 340)</td>
</tr>
<tr>
<td>MEP</td>
<td>1.05 (0.62-1.76)</td>
<td>1.23 (0.97-1.57)</td>
<td><strong>1.49 (1.05-2.13)</strong></td>
</tr>
<tr>
<td></td>
<td>( P = .857 )</td>
<td>( P = .091 )</td>
<td><strong>( P = .024 )</strong></td>
</tr>
<tr>
<td>MiBP</td>
<td>1.09 (0.66-1.80)</td>
<td>0.97 (0.77-1.23)</td>
<td>1.08 (0.77-1.52)</td>
</tr>
<tr>
<td></td>
<td>( P = .734 )</td>
<td>( P = .835 )</td>
<td>( P = .630 )</td>
</tr>
<tr>
<td>MnBP</td>
<td>1.01 (0.61-1.66)</td>
<td>1.24 (0.98-1.58)</td>
<td><strong>1.47 (1.05-2.07)</strong></td>
</tr>
<tr>
<td></td>
<td>( P = .972 )</td>
<td>( P = .065 )</td>
<td><strong>( P = .025 )</strong></td>
</tr>
<tr>
<td>MBzP</td>
<td>0.98 (0.58-1.67)</td>
<td>0.83 (0.65-1.05)</td>
<td>0.95 (0.68-1.33)</td>
</tr>
<tr>
<td></td>
<td>( P = .943 )</td>
<td>( P = .127 )</td>
<td>( P = .780 )</td>
</tr>
<tr>
<td>MEHP</td>
<td>1.34 (0.79-2.26)</td>
<td>0.83 (0.65-1.05)</td>
<td>1.23 (0.88-1.72)</td>
</tr>
<tr>
<td></td>
<td>( P = .267 )</td>
<td>( P = .604 )</td>
<td>( P = .212 )</td>
</tr>
</tbody>
</table>

Note the reduced case numbers for allergic sensitization because blood samples were not available for all children participating in the follow-up investigation.

*Analysis was performed using a logistic regression model with phthalate values categorized into quartiles; odds ratios (ORs) were adjusted for sex, parental history of atopy, maternal smoking and/or ETS exposure at home, siblings, maternal education, cat ownership, and breast-feeding until 6 months.

†According to the Pharmacia CAP System, concentrations of >0.35 kU/L were regarded as positive.

‡Significant values are marked in boldface (\( P < .05 \)).
<table>
<thead>
<tr>
<th>Treg cells</th>
<th>Hay fever</th>
<th>Sensitization to food allergens (fx5)</th>
<th>Sensitization to inhalant allergens (sx1)</th>
</tr>
</thead>
<tbody>
<tr>
<td>0-3rd year</td>
<td>3.6% (17 of 475)</td>
<td>15.3% (79 of 515)</td>
<td>12.1% (41 of 340)</td>
</tr>
<tr>
<td>First year</td>
<td>15.3% (79 of 515)</td>
<td>9.7% (0.67-1.41)</td>
<td>0.83 (0.55-1.21)</td>
</tr>
<tr>
<td>Second year</td>
<td>12.1% (41 of 340)</td>
<td>0.89 (0.51-1.57)</td>
<td>1.02 (0.75-1.39)</td>
</tr>
<tr>
<td>Third year</td>
<td>13.8% (40 of 289)</td>
<td>0.89 (0.51-1.57)</td>
<td>0.84 (0.61-1.16)</td>
</tr>
</tbody>
</table>

*Analysis was performed using a logistic regression model with Treg-cell values categorized into quartiles; odds ratios (ORs) were adjusted for sex, parental history of atopy, maternal smoking and/or ETS exposure at home, siblings, maternal education, cat ownership, and breast-feeding until 6 months.

†According to the Pharmacia CAP System, concentrations of >0.35 kU/L were regarded as positive.

‡Significant values are marked in boldface (P < .05).