

RESEARCH

Open Access



Maternal allergic disease history affects childhood allergy development through impairment of neonatal regulatory T-cells

Shan-shan Meng^{1,2}, Rong Gao¹, Bing-di Yan¹, Jin Ren¹, Fei Wu³, Peng Chen⁴, Jie Zhang¹, Li-fang Wang³, Yuan-ming Xiao³ and Jing Liu^{1*}

Abstract

Background: Maternal allergic disease history and impaired regulatory T-cells (Tregs) are critical risk factors for allergy development in children. However, the mechanisms that underlie these risk factors remain poorly defined. Therefore, the aim of this study was to assess whether maternal allergies affect the Tregs of offspring and lead to allergy development in childhood.

Methods: A total of 332 mothers of healthy newborns (234 from no allergic mothers, 98 from allergic mothers) were recruited to this study. Detailed questionnaires were administered yearly to determine the allergy status of the mothers and the newborns from birth to 3 years of age. Cord blood samples obtained at the time of birth were analysed for Treg counts, as well Treg activity, based on their response to Toll-like receptor (TLR) stimuli such as lipid A (LPA) and peptidoglycans (PPG). Surface markers, associated genes, suppressive capacity, and cytokine levels of Tregs were also measured. Possible correlations between Treg activity and maternal or neonate allergies were assessed. In addition, environmental microbial content and other known risk factors for allergies were measured.

Results: Cord blood mononuclear cells (CBMCs) from offspring with allergic mothers showed fewer CD4⁺CD25⁺FOXP3⁺ T cells, lower expression levels of associated genes, and reduced cytokine production of interleukin (IL)-10 and interferon- γ ($P < 0.05$), especially via the PPG-TLR2 pathway. Suppression of effector T cells by Tregs from children of mothers with allergies was impaired, especially IL-13 production by Type 2 T helper (Th2) cells ($P = 0.026$). Children who developed allergies in the first 3 years of life had lower numbers of CD4⁺CD25⁺FOXP3⁺ T cells and reduced FOXP3 expression and IL-10 production as newborns ($P < 0.05$). Maternal allergic background was identified as a risk factor for allergy development in the children (*Odds ratio* (OR) = 2.46, 95 % CI = 1.05–5.79); while declining Treg numbers, IL-10 production, and FOXP3 expression in neonates (PPG and LPA stimulated) were identified as independent risk factors for allergic diseases in offspring at 3 years of age after adjusting for maternal allergic history and environmental factors ($P < 0.05$).

Conclusion: Maternal allergy correlated with impaired Tregs in neonates, and this could enhance the susceptibility of offspring to allergic diseases in early childhood due to an imbalance of Th1 and Th2 cells.

Keywords: Allergy, Cord blood, Regulatory T-cell, Toll like receptor, Children

* Correspondence: liujing020901@163.com

¹Department of Respiratory Medicine, The Second Hospital of Jilin University, No.218, Ziqiang Street, Nanguan District, Changchun, China
Full list of author information is available at the end of the article

Background

The incidence of childhood allergic diseases has exhibited a marked increase in recent years, with the incidence in China being greater than 3.3 % [1]. A family history of allergic diseases, especially maternal allergies, is associated with a high likelihood of childhood allergy development [2, 3]. Some prospective cohort studies of Western countries have shown that susceptibility to allergic diseases in children may occur early in life [4–6], and during this time period regulatory T-cells (Tregs) could be involved. However, analogous follow-up studies in China have not been performed.

Because Tregs influence the early steps of allergy initiation [7], it is important to measure Treg cell abundance. New research has found that CD4⁺CD25⁺FOXP3⁺ T cells were found to be excellent markers for Tregs [8]. Stable expression of FOXP3 has been observed in mature Tregs, and this expression profile was found to be highly specific for Tregs [9]. Other Treg specific genes include glucocorticoid-induced TNFR-related protein (GITR) and lymphocyte activation gene 3 (LAG-3) [10, 11]. Furthermore, interleukin (IL)-10 is a relatively specific cytokine of Tregs, and secretion of this cytokine may act as an additional marker for Tregs [12]. The suppressive capacity of Tregs is required to maintain the balance of T helper (Th) cells, including Th1, Th2, and Th17 cells. Unchecked proliferation of these Th cell populations has been shown to play a key role in the pathogenesis of allergic diseases. Toll-like-receptor (TLR)-induced innate immune responses affect the early differentiation of T cells. In particular, TLR2 and TLR4 are induced by peptidoglycans (PPG) and lipid A (LPA), respectively, and have been shown to be important mediators of T cell differentiation. Immune system stimulation may be critical to early immune modulation, and several studies have reported that young children growing up in environments rich in microbial stimuli have a lower risk of allergy [13, 14]. At the molecular level, activation of the TLR2/4 pathways has been shown to upregulate Tregs and induce cytokine secretion in both children and animal models with allergy development [13–15].

We hypothesize that the extent of the T cell response to TLR2/4 stimulation may be influenced by a maternal background of atopy. Therefore, the aim of the present study was to examine whether TLR2/4 stimulation influences immunity in addition to the activity of T cell subsets such as Tregs. Secretion of cytokines IFN- γ (Th1 cells), IL-13 (Th2 cells), and IL-17 (Th17 cells) were also examined. Furthermore, follow-up studies were conducted to assess whether maternal allergic status affected the susceptibility of developing allergic diseases in the first three years of childhood.

Methods

Study populations

A total of 332 mothers of healthy newborns in the city of Changchun, China were recruited to this study. Of these

mothers, 234 had no allergies, while 98 had been diagnosed with allergies. We defined maternal allergy history as a diagnosis of asthma [16], allergic rhinitis [17], and/or allergic eczema [18] with at least one positive specific IgE (sIgE). Enrollment occurred from January 2012 through December 2012 with the approval of the ethics committee of the Second Hospital of Jilin University (2012–003). Pregnant mothers were approached at least one month before delivery for consent and they completed a detailed questionnaire that assessed maternal and infant data. In addition, cord blood was collected from each neonate ($n = 332$). The study inclusion criteria consisted of healthy neonates and mothers with uncomplicated pregnancies. Exclusion criteria were preterm deliveries, parents with autoimmune diseases, perinatal infections, maternal antibiotic or systemic glucocorticoid use in the last trimester, rural newborns, unhealthy newborns, or paternal allergic diseases.

Follow-up studies were conducted from 2012 to 2015 by questionnaire when the children were 12, 24, and 36 months old. Follow-up questionnaires were only completed for 273 children. The population characteristics that were examined for the children included: vaccination status, pet exposure, air conditioner exposure, breast feeding, diet, frequency of viral respiratory infections, medications, household income, parental education, and cohabitation with a smoker. Allergic wheezing (as a proxy for asthma in children ≤ 5 years old [19]), allergic rhinitis, and allergic eczema were diagnosed by paediatricians based on clinical manifestations, the presence of at least one sIgE+, and parent-described patient symptoms. The cohort belonged to a low economic status group ($\leq 20,000$ Renminbi income/year) based on available data from the Statistics Department (<http://www.stats.gov.cn/>) on the Per Capita Disposable Income (PCDI) for the Jilin Province collected from 2012–2015.

Dust collection and analysis

To evaluate the presence of environmental allergens, dust samples were collected from the living rooms of the pregnant women one month before delivery. Dust samples were subsequently collected from the children's bedrooms at 12 ± 1 , 24 ± 1 , and 36 ± 1 months post delivery. All dust samples were collected on pre-weighed glass fibre filters. Collection was performed using vacuum cleaners with sampling nozzles (ALK, Horsholm, Denmark) according to a standardized protocol with photo and video instructions. Collected samples were stored at -20 °C until extraction. Dust-containing filters were weighed and extracted in a volume of 5–40 ml, as determined by the net dust weight (<0.5 g, 5 ml; 0.5 – 1.0 g, 10 ml; 1.0 – 2.0 g, 20 ml; > 2.0 g, 40 ml). Endotoxin levels were measured with a kinetic chromogenic *Limulus* amoebocyte lysate (LAL) test and glucan levels were detected in an inhibition enzyme immunoassay an inhibition EIA [20].

IgE detection

Total IgE and sIgE were measured in peripheral blood serum samples collected from both mothers and children using the RAST method. Levels of sIgE in serum against six foods (cow's milk, hen's egg, hazelnut, carrot, peanut, and wheat flour) and 13 inhalant allergens (*Dermatophagoides farinae*; *Dermatophagoides pteronyssius*; cat fur; dog fur; horse fur; *Alternaria* species; mugwort, plantain, birch, alder, hazel, and rye pollen; and grass pollen mix) were assayed. A positive sIgE was defined as one or more positive reactions ≥ 0.35 IU/mL to a panel of 19 common allergens.

Lymphocyte stimulation and cytokine production

Cord blood (20 ml) and peripheral blood (5–10 ml) samples were collected from healthy neonates within 24 h of birth. Cord blood mononuclear cells (CBMCs) were isolated from cord blood by density gradient centrifugation with Ficoll-Paque PLUS (GE Healthcare, Piscataway, NJ, USA) after dilution in PBS (Gibco, Carlsbad, CA, USA). Cells were washed in RPMI 1640 (Gibco) and diluted in 10 % human serum (Gibco) to a concentration of 5×10^6 cells/ml. CBMCs were cultured for three days in the presence of 10 μ g/ml PPG, 0.1 μ g/ml LPA (Sigma-Aldrich, St. Louis, MO, USA), or without a stimulation factor. Concentrations of IL-10, IL-13, IL-17, and interferon- γ (IFN- γ) in the CBMC supernatants were measured with a Human Cytokine Multiplex Assay kit (Bio-Rad, Munich, Germany) with Luminex detection. The lower limits of

detection for the assay (pg/mL) were: 0.9 (IL-10), 2.1 (IL-13), 0.2 (IL-17), and 1.3 (IFN- γ).

Flow cytometry

CBMCs were analysed by flow cytometry (BD Biosciences, San Jose, CA, USA) after three days of cultivation. For surface staining, 2 μ l of anti-human CD4-fluorescein isothiocyanate (FITC), 1 μ l of CD25-RPE-Cy5, 1 μ l of IgG1-FITC (DakoCytomation, Glostrup, Denmark), and 0.5 μ l of IgG2a RPE-Cy5 (BD Biosciences) were added. For CD4/CD25/FOXP3 co-staining, 8 μ l of anti-human CD4-FITC and 4 μ l of anti-human CD25-RPE-Cy5 antibodies were added to 1×10^6 cells in 100 ml PBS. Cells were then permeabilized and FOXP3-PE or the corresponding isotype control antibodies were added. Data were analysed with CellQuest software (BD Biosciences).

Real-time quantitative RT-PCR

CBMCs were collected after three days of cultivation and total RNA was isolated immediately with the Tri-Pure Isolation Reagent (Roche, Mannheim, Germany). RNA concentration was determined with a Nano-Drop2000 (Thermo Scientific, Waltham, MA, USA), and samples were stored at -20 °C. Reverse transcriptase were performed (Invitrogen, Karlsruhe, Germany) for cDNA synthesis. Specific primer pairs were designed with Primer Express software (Vector NTI Advance10). Quantitative real-time PCR was performed with an iCyclerIQ-

Table 1 Characteristics of mothers and neonates

	Mothers without allergies (<i>n</i> = 234)	Mothers with allergies (<i>n</i> = 98)	<i>P</i> -value
Maternal age, y (IQR)	26 (23–32)	28 (22–35)	0.26
Body mass index (IQR)	23.15 (19.98–26.89)	22.04 (19.05–28.04)	0.23
Smoking status, n (%)			
Non smoking	225 (96.15)	97 (98.98)	0.29
Smoking during pregnancy	0 (0)	0 (0)	
Smoking until pregnancy	9 (3.85)	1 (1.02)	0.29
Maternal total IgE, UI/ml (IQR)	26.4 (9.42–57.9)	68.5 (42.5–222.4)	0.005
Maternal allergies [sIgE (+)], n (%)	0 (0)	98 (100)	
Asthma	0 (0)	23 (23.47)	
Allergic rhinitis	0 (0)	66 (67.35)	
Allergic eczema	0 (0)	30 (30.61)	
Endotoxins in maternal bedroom, EU/mg (IQR)	14.77 (4.20–55.18)	11.73 (2.64–50.14)	0.10
β (1,3)-glucans in maternal bedroom, μ g/mg (IQR)	3.42 (1.04–9.57)	2.33 (0.67–5.88)	0.16
Vaginal delivery, n (%)	210 (89.74)	82 (83.67)	0.14
Male neonate gender, n (%)	120 (51.28)	51 (52.04)	0.90
Gestational age, weeks (SEM)	39.91 (1.20)	39.93 (1.18)	0.94
Birth weight, g (SEM)	3403.3 (407.37)	3610.56 (427.19)	0.45
Birth length, cm (IQR)	51.75 (51.36–52.14)	53 (52.48–53.52)	0.19
Apgar 5 min score > 8, n (%)	234 (100)	98 (100)	

Data in boldface are significant with $P \leq 0.05$. Data are reported as either the mean (SEM) or median (IQR) depending on the data distribution

multicolor Real-Time PCR Detection System. SYBR Green (Applied Biosystems, Darmstadt, Germany) was used to detect double-stranded PCR products. The C_T means the number of PCR cycles which are required for the fluorescence signal to exceed the detection threshold value. The threshold cycle (C_T) of each target product is associated with the amplification plot of 18S. The level of mRNA from individual gene is described as gene expression. The fold difference in gene expression was calculated based on the difference in normalized C_T values between the stimulation and non-stimulation groups. The relative quantitative results showed changes of gene expressions in stimulated samples when compared with those in unstimulated ones.

Functional analysis of tregs

CD3⁻ T cells were isolated (CD3⁻ Cell Isolation kit, MiltenyiBiotec, Köln, Germany) and irradiated as antigen-presenting cells. CD3⁻ T cell purity reached 98 %. CD4⁺CD25⁺ T cells were isolated from CBMCs by tandem positive selection (MiltenyiBiotec). CD4⁺CD25⁻ T cells were collected from the flow through during CD25 selection. The purity of both the CD4⁺CD25⁻ and CD4⁺CD25⁺ T cell populations was > 95 %. CD4⁺CD25⁻ T cells (effector T cells) were labelled with 5 μ mol/L CFSE

(CellTrace™ CFSE Cell Proliferation Kit, Invitrogen), and were subsequently incubated for three days with irradiated CD3⁻ cells in co-culture with or without CD4⁺CD25⁺ T cells (Tregs). The co-cultures were stimulated with phytohaemagglutinin (0.8 μ g/ml). CFSE-stained CD4⁺CD25⁻ T cells were isolated by flow cytometry and ³H-thymidine incorporation was used to measure their proliferation. Concentrations of cytokines IL-13, IL-17, and IFN- γ were measured in the cell culture supernatants using a Human Cytokine Multiplex Assay Kit. The resulting values were compared between the effector T cells only group and the effector T cells plus Tregs group. The suppressive capacity of the Tregs on the effector T cells was further compared between the maternal allergy and non-allergy groups.

Statistical analysis

Data were analyzed using SigmaStat1.0 software. Differences in cell counts, gene expression, suppressive function, and cytokine production for the Tregs present in the neonate cord blood samples were compared between maternal allergy and non-allergy groups, and between children's allergy and non-allergy groups. Correlations between newborn Treg function and the development of childhood allergy were analysed. Metric data were

Table 2 Physical and environmental characteristics of children included in the follow-up study

	Children without allergies (n = 202)	Children with allergies (n = 71)	P-value
Male, n (%)	97 (48.02)	38 (53.52)	0.49
Body weight at 3 years of age, g (SEM)	15.1 (3.47)	16.7 (4.21)	0.14
Height at 3 years of age, cm (IQR)	97.6 (92.1–111.6)	97.4 (91.8–107.2)	0.58
Allergic diseases during 0–3 years of age, n (%)			
All allergies [sIgE (+)]	0 (0)	71 (100)	
Wheezing	0 (0)	19 (26.76)	
Allergic rhinitis	0 (0)	50 (70.42)	
Allergic eczema	0 (0)	47 (66.19)	
Maternal allergic history, n (%)	62 (30.69)	38 (53.52)	0.001
Average composition in dust from children's bedrooms			
Endotoxin, EU/mg (IQR)	13.65 (3.51–45.78)	11.65 (2.78–39.44)	0.05
β (1,3)-glucans, μ g/mg (IQR)	2.67 (0.54–7.45)	2.19 (0.51–5.56)	0.21
Pet exposure, n (%)	5 (2.47)	0 (0)	0.33
Air conditioner exposure, n (%)	22 (10.89)	9 (12.68)	0.54
Breast fed, n (%)	135 (66.83)	50 (70.42)	0.66
Respiratory virus infections, n (%)			
< 3 per year	139 (68.81)	47 (66.20)	0.44
3–5 per year	35 (17.33)	14 (19.72)	0.28
> 5 per year	28 (13.86)	10 (14.08)	0.54
Low household income, n (%)	92 (45.54)	26 (36.62)	0.21
Low parental education (middle school and lower), n (%)	35 (17.33)	12 (16.90)	1.0
Co-habitation with a smoker, n (%)	10 (4.95)	3 (4.23)	1.0

Data in boldface are significant with $P \leq 0.05$. Data were reported as either the mean (SEM) or median (IQR) depending on the data distribution

evaluated by either a *t*-test or the Mann–Whitney rank test with or without normal distribution. Data are reported as the mean ± standard error of the mean (SEM) or the median with a 25–75 % interquartile range (IQR), depending on their distribution. Attributes data were analyzed with a χ^2 test or Fisher’s exact test. Multiple logistic regression was performed for the correlation analyses. Odds ratio (OR) values were reported when applicable. Statistical significance was defined as a *P*-value < 0.05.

Results

Cohort characteristics

Table 1 reports the available data for the 332 mother/neonate pairs of the present cohort (including 234 pairs in the maternal non-allergy group and 98 pairs in the maternal allergy group). Total maternal IgE levels were significantly higher in the maternal allergy group compared with the maternal non-allergy group (*P* = 0.005). Otherwise, there were no distinguishing characteristics between the mothers of the two groups, except there were four allergic asthmatic mothers who took low doses of inhaled glucocorticoids at gestational weeks 20–30.

A follow-up study was conducted for the 273 children for whom a completed follow-up questionnaire was completed at 0–3 years of age (Table 2). There were 59 children that were lost to follow-up due to a change in address or telephone number. While approximately half (53.52 %) of the allergic children had allergic mothers, only 30.69 % of the non-allergic children had allergic mothers. It was observed that the children with maternal allergy histories were more prone to allergic diseases (*P* = 0.001). The average concentration of endotoxin dust in the samples collected from the bedrooms of the allergic children was less than that in the bedroom samples for the non-allergic children (*P* = 0.05). However, there was no difference in β (1,3)-glucan exposure, diet, frequency of viral respiratory infections, household income, parental education, or cohabitation with a smoker between the children with and without allergies.

Treg abundance and activity

Treg abundance in the CBMCs (characterized as CD4⁺CD25⁺FOXP3⁺ T cells, Fig. 1) and cytokine production are listed in Table 3. Tregs were less abundant in the

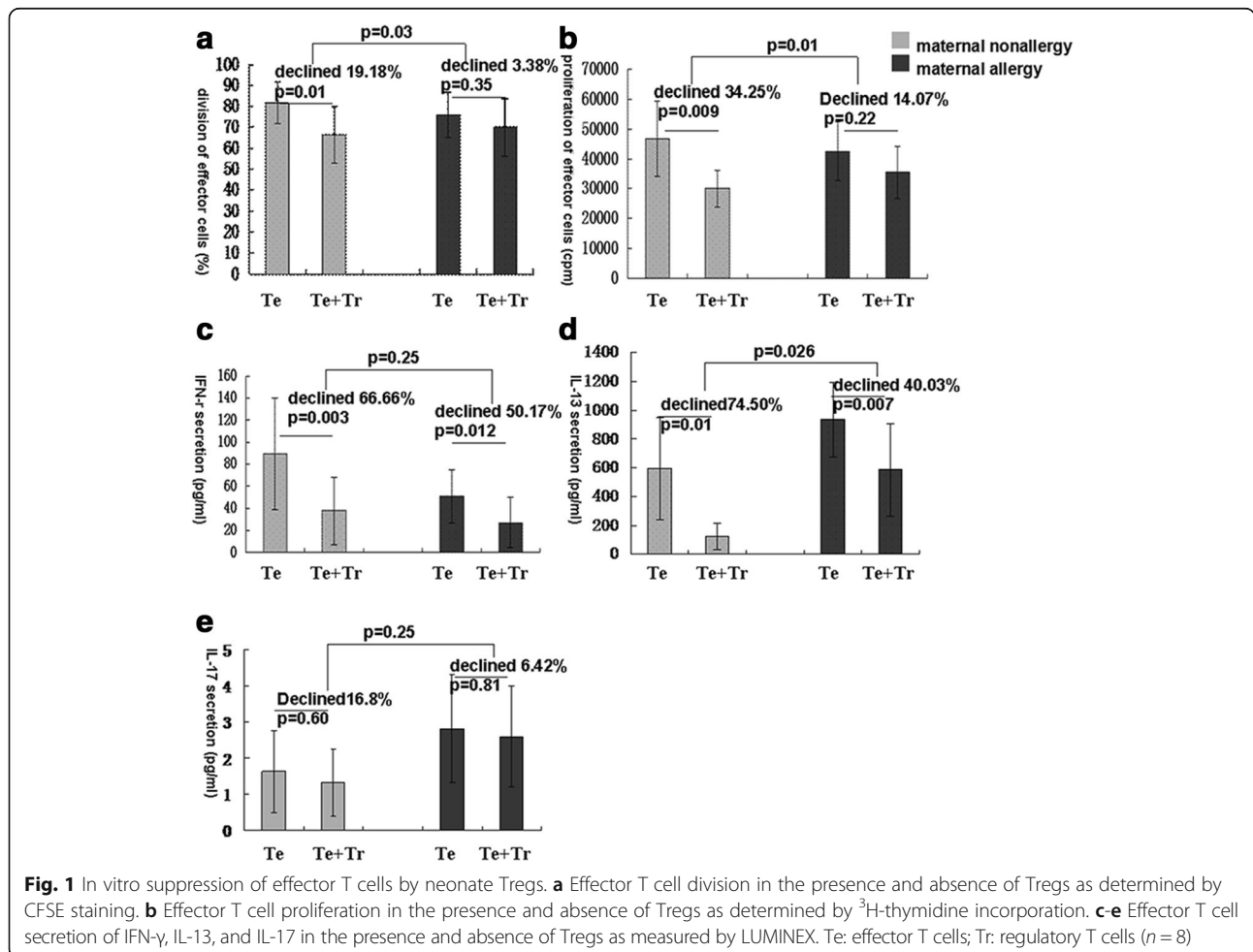


Table 3 Tregs and cytokines in CBMCs of offspring with and without maternal allergy

	Maternal non-allergy (n = 222)	Maternal allergy (n = 90)	P-value
CD4 ⁺ CD25 ⁺ FOXP3 ⁺ T cells, % in CBMCs (SEM)			
U	1.78 (0.71)	1.57 (0.76)	0.12
LPA	2.05 (0.75)	1.81 (0.76)	0.08
PPG	2.33 (0.69)	1.99 (0.68)	0.01
FOXP3 gene expression, fold difference (IQR)			
LPA	3.14 (1.79–5.24)	2.50 (0.83–4.88)	0.021
PPG	3.36 (1.32–4.92)	2.69 (1.22–5.86)	0.024
GITR gene expression, fold difference (IQR)			
LPA	2.83 (1.53–6.24)	1.50 (0.95–3.03)	0.16
PPG	3.93 (1.87–8.46)	2.46 (0.95–5.89)	0.15
LAG3 gene expression, fold difference (IQR)			
LPA	2.83 (1.39–6.45)	1.53 (0.90–3.03)	0.12
PPG	2.93 (1.88–5.66)	2.00 (0.85–2.95)	0.028
IL-10 production, pg/ml (IQR)			
U	0.19 (0.01–0.27)	0.035 (0.01–0.23)	0.81
LPA	173.71 (109.24–198.26)	135.16 (91.37–241.89)	0.43
PPG	1104.47 (806.43–1519.56)	868.44 (603.76–1153.45)	0.006
IL-13 production, pg/ml (IQR)			
U	0.16 (0.01–0.33)	0.13 (0.01–0.36)	0.52
LPA	3.81 (1.45–7.80)	5.87 (1.48–13.21)	0.06
PPG	25.44 (12.41–53.98)	33.38 (13.14–61.53)	0.021
IFN-γ production, pg/ml (IQR)			
U	1.02 (0.01–1.99)	1.02 (0.45–1.99)	0.49
LPA	5.52 (3.47–17.64)	3.95 (1.64–9.66)	0.04
PPG	14.28 (6.57–33.42)	8.66 (5.02–18.96)	0.047
IL-17 production, pg/ml (IQR)			
U	0.09 (0.01–0.47)	0.12 (0.01–0.67)	0.38
LPA	2.39 (0.24–4.39)	2.26 (0.81–4.10)	0.86
PPG	2.7 (1.65–10.99)	4.03 (2.74–19.25)	0.049

U unstimulated, LPA stimulation with 0.1 μg/ml LPA, PPG stimulation with 10 μg/ml PPG
 Data in boldface are significant with $P \leq 0.05$. Data are reported as the mean (SEM) or median (IQR) depending on the data distribution

CBMCs collected from the neonates with maternal allergies than in those without allergies, especially following PPG stimulation ($P = 0.01$, Table 3). Both PPG/LPA-stimulated *FOXP3*, and PPG-stimulated *LAG3*, gene expression levels were lower in the neonates with maternal allergies than in the neonates from mothers without allergies ($P < 0.05$). Maternal allergic background also affected cytokine responses. IL-10 levels in the PPG-TLR2 innate immune pathway were lower in the offspring from the mothers with allergies than those from the mothers without allergies ($P = 0.006$), in agreement with the observations in CD4⁺CD25⁺FOXP3⁺T cells and the

FOXP3/LGA3 expression profiles for these groups. Production of the Th1 cytokine, IFN-γ, was also reduced upon LPA and PPG stimulation in the newborns with allergic mothers compared to the newborns with non-allergic mothers ($P = 0.04$ and $P = 0.047$, respectively), while increased production of PPG-stimulated IL-13 and IL-17 was observed as well ($P = 0.021$ and $P = 0.049$, respectively).

In Table 4, Treg abundance, related gene expression, and cytokine production data are compared for neonate cord blood samples based on their childhood allergy status, independent of maternal allergy status. Children with allergies had a lower number of PPG-stimulated CD4

Table 4 Tregs and cytokines in CBMCs of offspring with and without childhood allergies

	Children without allergies (n = 202)	Children with allergies (n = 71)	P-value
CD4 ⁺ CD25 ⁺ FOXP3 ⁺ T cells, % in CBMCs (SEM)			
U	1.91 (0.67)	1.17 (0.61)	0.12
LPA	2.15 (0.73)	1.45 (0.54)	0.08
PPG	2.44 (0.63)	1.57 (0.49)	0.01
FOXP3 gene expression, fold difference (IQR)			
LPA	3.14 (1.60–5.25)	1.41 (0.70–2.64)	0.01
PPG	4.03 (1.32–6.02)	1.98 (1.25–4.14)	0.02
GITR gene expression, fold difference (IQR)			
LPA	2.14 (1.0–5.14)	2.79 (0.96–10.44)	0.12
PPG	2.42 (0.99–6.96)	3.67 (0.95–8.61)	0.44
LAG3 gene expression, fold difference (IQR)			
LPA	2.14 (0.61–4.65)	1.19 (0.68–3.36)	0.51
PPG	2.0 (0.54–3.73)	2.22 (0.63–4.74)	0.61
IL-10 production, pg/ml (IQR)			
U	0.10 (0.01–0.26)	0.15 (0.01–0.21)	0.35
LPA	177.93 (110.74–229.50)	135.28 (88.14–192.87)	0.09
PPG	1078.74 (730.84–1518.49)	830.74 (623.0–1038.23)	0.016
IL-13 production, pg/ml (IQR)			
U	0.16 (0.01–0.37)	0.21 (0.01–0.35)	0.63
LPA	4.5 (1.65–11.83)	5.07 (1.70–13.26)	0.80
PPG	31.70 (12.88–50.53)	30.59 (11.90–61.45)	0.53
IFN-γ production, pg/ml (IQR)			
U	1.02 (0.01–1.99)	1.66 (0.45–2.51)	0.09
LPA	5.08 (2.81–16.86)	4.91 (2.82–7.72)	0.55
PPG	13.15 (6.12–30.33)	10.22 (4.96–26.50)	0.53
IL-17 production, pg/ml (IQR)			
U	0.01 (0.01–0.47)	0.01 (0.01–0.43)	0.96
LPA	1.71 (0.19–4.39)	0.84 (0.01–2.80)	0.12
PPG	2.94 (1.28–11.41)	2.34 (0.74–6.48)	0.18

U unstimulated, LPA stimulation with 0.1 μg/ml LPA, PPG stimulation with 10 μg/ml PPG
 Data in boldface are significant with $P \leq 0.05$. Data are reported as the mean (SEM) or median (IQR) depending on the data distribution

$CD25^+FOXP3^+$ T cells ($P = 0.01$), reduced *FOXP3* expression in response to PPG and LPA ($P = 0.02$ and $P = 0.01$, respectively), and lower IL-10 production ($P = 0.016$), compared to the children without allergies.

The suppressive capacity of tregs

Tregs from the children with allergic mothers suppressed the division and proliferation of effector T cells less efficiently than the Tregs from the children with non-allergic mothers ($P = 0.03$ and $P = 0.01$, respectively; Fig. 2). The Tregs of the former group also exhibited a significant decline in the production of IL-13 compared to the latter group ($P = 0.026$, Fig. 2). In contrast, production of IL-17 and IFN- γ were unaffected ($P > 0.05$, Fig. 2).

Risk factors for allergic diseases

Maternal allergy was a significant risk factor for developing allergic diseases ($OR = 2.46$, $95\% CI = 1.05-5.79$; Table 5). After adjusting for maternal allergy, environmental exposure, diet, frequency of viral respiratory infections, household income, parental education, and cohabitation with a smoker as factors, the percent of $CD4^+CD25^+FOXP3^+$ T cells in the PPG-stimulated cord blood

samples was found to negatively correlate with allergy status, allergic rhinitis, and allergic eczema in children, yet not with wheezing (Table 6). Conversely, a decrease in the percent of $CD4^+CD25^+FOXP3^+$ T cells in the unstimulated cord blood samples was identified as a risk factor for wheezing (*adjusted OR* = 0.017, $95\% CI = 0.001-0.46$; Table 6). Decreased PPG-stimulated IL-10 production, as well as decreased LPA-stimulated and PPG-stimulated *FOXP3* expression, in cord blood were further identified as risk factors for a positive allergic status in children and allergic rhinitis (Tables 7 and 8, respectively).

Discussion

In this birth cohort study, Tregs in the cord blood samples obtained from neonates with allergic mothers were fewer in number. The Tregs also exhibited compromised suppressive function, lower levels of expression for associated genes, and decreased IL-10 production. Moreover, the PPG-TLR2 innate immune response pathway was particularly affected. As a result, an increased susceptibility to allergies in early childhood (0–3 years old) was observed. Maternal allergic disease history and neonatal impairment of Tregs were both

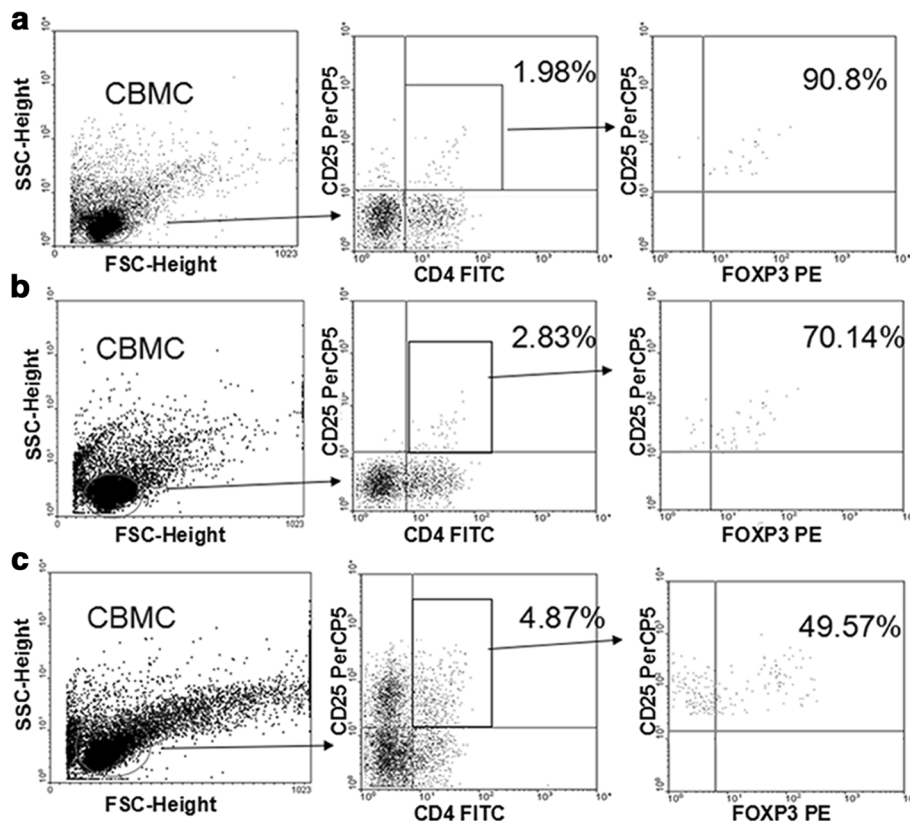


Fig. 2 Percent of $CD4^+CD25^+FOXP3^+$ T cells in untreated and stimulated CBMCs. $CD4^+CD25^+FOXP3^+$ T cell populations were determined by fluorescent antibody staining and FACS. **a** Unstimulated $CD4^+CD25^+FOXP3^+$ T cells represented 1.80 % of the CBMC population (=90.8 % * 1.98 %). **b** $CD4^+CD25^+FOXP3^+$ T cells represented 1.98 % of the LPA-stimulated CBMC population (=70.14 % * 2.83 %). **c** $CD4^+CD25^+FOXP3^+$ T cells represented 2.41 % of the PPG-stimulated CBMC population (=49.57 % * 4.87 %). See Methods for details on CBMC culturing, stimulation, and fluorescence staining

Table 5 Association of allergy in children with maternal allergic history and endotoxin or β (1, 3)-glucan exposure

	OR (95 % CI) P-value		
	Maternal allergic history	Endotoxin (EU/mg)	β (1,3)-Glucans (μ g/mg)
All allergies	2.46 (1.05–5.79) 0.03	0.90 (0.75–1.02) 0.15	1.05 (0.95–1.16) 0.27
Allergic wheezing	3.02 (0.68–13.35) 0.14	0.88 (0.75–1.03) 0.11	1.06 (0.92–1.20) 0.42
Allergic rhinitis	2.13 (0.815–5.56) 0.12	0.93 (0.85–1.01) 0.17	1.01 (0.95–1.07) 0.68
Allergic eczema	3.21 (1.18–8.69) 0.02	0.94 (0.84–1.06) 0.35	1.03 (0.93–1.15) 0.54

Data in boldface are significant with $P \leq 0.05$

identified as risk factors for susceptibility to allergies in the children examined, independent of social and environmental factors for the children and their families. These results are consistent with those of previous studies of allergies in European populations [4, 5].

Many studies have reinforced the importance of in utero exposure for fetal immune development and the programming of susceptibility to asthma and allergic diseases [21]. In the present study, these factors were adequately considered to avoid potential interference. In addition, based on the results of a study by Schaub et al. where a higher percentage of allergic neonates with allergic mothers were more likely to have allergic fathers [5], paternal allergies were established as an exclusion criterion for the present study in order to focus on the effect of maternal allergic disease history on offspring. The study by Schaub et al. also reported increased birth weights and lengths for the newborns with allergic mothers [5]. However, no differences in birth weight or length between the groups of the present study were observed. This may be due to the similarity in the diets and medication use of the mothers included in the present cohort. Over the course of child development, many social and environmental factors should be considered when assessing influences on allergy development. Therefore, factors which have been shown to influence allergic disease development in children were considered in the present study, including: environmental exposure, diet, frequency of viral respiratory infections, household income, parental education, and co-habitation with a smoker [22]. It was observed that the endotoxin content in the living environments of the children with allergies was lower than in the living environments of the children without allergies. These results are consistent with

those from Karvonen et al. where endotoxin exposure in early life was found to be closely related to the development of allergic diseases in children [23]. Consequently, microbial content was adjusted for in the analyses of risk factors for children's allergic status.

As shown in Fig. 1a, b and c, FOXP3⁺ T cells represented > 90 % of the unstimulated CD4⁺CD25⁺ T cells. Therefore, CD4⁺CD25⁺ T cells were isolated from the unstimulated CBMCs to serve as a Treg population that would have its suppressive function examined in vitro. There were fewer CD4⁺CD25⁺FOXP3⁺ T cells present in both the neonates with allergic mothers and the neonates who developed allergies in early childhood. While the expression levels of GITR, LAG3, and FOXP3 were previously shown to positively correlate with each other [5], the specificity and sensitivity of their expression slightly differed in the TLR-mediated innate immune responses that were induced. It has been shown that Tregs secreting IL-10 play an important role in allergic diseases, independent of CD25⁺ Tregs [12]. Therefore, IL-10 production was assayed as a readout of function for this Treg subgroup. However, IL-10 is also secreted from Th2 cells. In the present data, PPG-stimulation reduced IL-10 secretion in the newborns with allergic mothers, while secretion of the Th2-specific cytokine, IL-13, increased (Table 3). These results suggest that IL-10 production is characteristic of this Treg subgroup in the TLR-mediated innate immune response. Overall, the risk factors for allergies in children of the present cohort included a reduced percentage of CD4⁺CD25⁺FOXP3⁺ T cells, reduced FOXP3 expression, and PPG-stimulated IL-10 secretion (Tables 6, 7 and 8). However, the other cytokines assayed, IL-13, INF- γ , and IL-17, were not identified as risk factors (data not shown). To date,

Table 6 Association of allergy in children with % CD4⁺CD25⁺ FOXP3⁺ T in neonate CBMCs

	Adjusted OR (95 % CI) P-value		
	Unstimulated	LPA-stimulated	PPG-stimulated
All allergies	0.51 (0.18–1.51) 0.228	0.65 (0.23–1.85) 0.416	0.12 (0.04–0.43) 0.01
Allergic wheezing	0.017 (0.001–0.46) 0.016	2.15 (0.35–13.39) 0.41	0.36 (0.046–2.77) 0.32
Allergic rhinitis	0.49 (0.148–1.60) 0.23	0.39 (0.12–1.28) 0.12	0.26 (0.074–0.91) 0.035
Allergic eczema	1.08 (0.36–3.29) 0.88	0.57 (0.18–1.80) 0.34	0.16 (0.044–0.56) 0.04

Data in boldface are significant with $P \leq 0.05$. Data were adjusted for maternal allergy, environmental and social factors

Table 7 Association of allergy in children with Treg-specific cytokine IL-10 levels

	Adjusted OR (95 % CI) P-value		
	Unstimulated	LPA-stimulated	PPG-stimulated
All allergies	0.17 (0.01–2.34) 0.19	0.65 (0.25–1.67) 0.37	0.33 (0.13–0.86) 0.03
Allergic wheezing	1.16 (0.07–18.98) 0.91	0.33 (0.07–1.51) 0.15	0.69 (0.15–3.26) 0.64
Allergic rhinitis	0.21 (0.07–5.94) 0.35	1.27 (0.41–3.98) 0.68	0.29 (0.09–0.89) 0.04
Allergic eczema	0.14 (0.04–4.97) 0.28	0.81 (0.27–2.43) 0.70	0.50 (0.17–1.46) 0.21

Data in boldface are significant with $P \leq 0.05$. Data were adjusted for maternal allergy, environmental, and social factors

the traditional Th1/Th2 imbalance theory has not been sufficient to explain the complex pathogenesis for allergic diseases. In addition, Tregs, Th17 cells, Th9 cells, and other subgroups of T cells have all been implicated in allergic disease pathogenesis. Based on the importance of Tregs maintaining their suppressive function, we hypothesize that impairment of Tregs in neonates with allergic mothers leads to dysregulation of the differentiation of Th0 cells into Th1, Th2, and Th17 cells, and this is the driving force for allergy development in children.

When the children of this study were divided into those with or without allergies, regardless of maternal allergy status, down-regulation of Tregs according to the TLR2/4 pathway at the newborn stage partly predicted the children's allergic status. However, it was not clear whether the down-regulation of Tregs derived exclusively from maternal allergy history. By using multiple logistic regression, exclusion of maternal allergic disease history and environmental factors led to the identification of a reduced number of CD4⁺CD25⁺FOXP3⁺ T cells, reduced *FOXP3* gene expression, and reduced IL-10 secretion in neonates as independent risk factors for the allergic status of children (Tables 6, 7, and 8).

It was previously demonstrated that impairment of Tregs in neonates with allergic mothers was more profound in the TLR2/4-mediated innate immune response pathways, due in part to genetic variations in the TLRs present in cord blood [24]. Genetic-immunological interactions of the TLR pathway influence Tregs early in life and are modulated by maternal allergic disease history. Moreover, it is likely these interactions are relevant for immune maturation in the development of allergic

diseases in childhood. In addition to a genetic component, the time, frequency, and intensity of activation of TLR-mediated immune pathways are important for the maturation and differentiation of Tregs and other Th cells. For instance, increased microorganism exposure in early life in children and in mouse models has been shown to regulate the immune system by enhancing the Th1 pathway and weakening the Th2 pathway through TLR2/4 immune responses [25–27].

The results of the present study have revealed that maternal allergy status has a significant impact on Treg differentiation in the TLR2/4 immune pathway, yet not in the unstimulated pathway. The present results also confirm that the occurrence of allergic disease is the result of interactions between genetic and environmental factors. In particular, the neonatal period appears to be a key immune maturation period, and environmental endotoxin levels during this period can influence the extent of differentiation that takes place to generate different T cell subgroups in children via the TLR2/4 pathway. The number of Tregs present in fetal umbilical cord blood was observed to increase following PPG and LPA stimulation, thereby facilitating the differentiation of naive T cells into new Tregs. Maternal allergy history can influence modulation of differentiation and inhibit the differentiation rate of Tregs through the interactions of other T cell subgroups.

Thorburn et al. [28] showed that high fibre or acetate diets during pregnancy, which influence gut microbiota, can lead to marked suppression of allergic airway diseases in offspring by enhancing Treg numbers and function. These results are consistent with the observation that the intake of probiotics during pregnancy and early life have the potential to prevent childhood eczema [29] and IgE-associated allergies to some extent [30]. However, avoiding activation of the TLR-mediated immune pathway appears safer than a desensitization approach for preventing allergic disease development in the offspring of allergic pregnant women since the latter can up-regulate CD4⁺CD25⁺ T cells and IL-10 production in offspring [31, 32]. In the risk factor analysis performed in the present study, a decrease in PPG-stimulated Tregs was identified as a risk factor for children's allergic status, allergic rhinitis, and allergic eczema. Meanwhile, a

Table 8 Association of allergy in children with FOXP3 expression

	Adjusted OR (95 % CI) P-value	
	LPA-stimulated	PPG-stimulated
All allergies	0.78 (0.63–0.97) 0.04	0.80 (0.65–0.97) 0.05
Allergic wheezing	0.78 (0.52–1.16) 0.22	0.76 (0.52–1.13) 0.18
Allergic rhinitis	0.77 (0.59–0.99) 0.05	0.85 (0.69–1.04) 0.12
Allergic eczema	0.83 (0.66–1.04) 0.11	0.81 (0.64–1.03) 0.08

Data in boldface are significant with $P \leq 0.05$. Data were adjusted for maternal allergy, environmental, and social factors

decrease in unstimulated Tregs was only a risk factor for allergic wheezing. Thus, micro-stimulation of the TLR2 and TLR4 pathways with adjustment of diet and environmental exposure may be an effective early intervention for preventing allergic disease in neonates that have lower numbers of Tregs or that have a maternal allergic disease history by establishing a more beneficial Th1/Treg balance.

Conclusions

In summary, maternal allergic disease history and deficiency in Tregs in newborns are risk factors for the development of allergic diseases in children. The impairment of Tregs number and function in neonates could lead to susceptibility to allergic status and diseases through the imbalance of Th1/2 and Th17 cells. Because deficiency in Tregs is more obvious in the TLR2-mediated innate immune response pathway, we could seek possible micro-stimulation to up-regulate Tregs in the TLR2 pathway and provide early prevention for allergic diseases in children with maternal allergic histories.

Abbreviations

CBMC: Cord blood mononuclear cell; C_T: Threshold cycle; FITC: Fluorescein isothiocyanate; FOXP3: Forkhead/winged helix transcription factor; GITR: Glucocorticoid-induced TNFR-related protein; IFN- γ : Interferon- γ ; IL: Interleukin; IQR: Interquartile range; LAG3: Lymphocyte activation gene 3; LAL: Limulus ameobocyte lysate; LPA: Lipid A; OR: Odds ratio; PPG: Peptidoglycans; sIgE: Specific IgE; Th: T helper; TLR: Toll-like receptor; Treg: Regulatory T-cell

Acknowledgements

Not applicable.

Funding

This study was funded by National Natural Science Foundation of China [81300014].

Availability of data and materials

Due to the privacy policy of patients, we can't openly release the dataset of questionnaire survey underlying the conclusions of the paper available in public. A truncated dataset after eliminating all potentially identifiable features may be provided on an individual request basis.

Authors' contributions

Jing Liu conceived and designed the research. Shan-shan Meng wrote and edited the manuscript. Shan-shan Meng, Rong Gao, Bing-di Yan, Jin Ren, Fei Wu, Peng Chen, Jie Zhang, Li-fang Wang, Yuan-ming Xiao, Jing Liu make data collection and analyzation. Jie Zhang, Jing Liu reviewed and revised it critically for important intellectual content. Jing Liu gave final approval of the version to be published. All authors read and approved the final manuscript.

Competing interests

The authors declare that they have no competing interests.

Consent for publication

Not applicable.

Ethics approval and consent to participate

The research protocol was approved by the ethics committee of the Second Hospital of Jilin University. Each patient gave their written informed consent before the survey.

Author details

¹Department of Respiratory Medicine, The Second Hospital of Jilin University, No.218, Ziqiang Street, Nanguan District, Changchun, China. ²Department of Critical Care Medicine, Zhongda Hospital, School of Medicine, Southeast University, Nanjing, China. ³Department of Obstetrics and Gynecology, The Second Hospital of Jilin University, Changchun, China. ⁴Department of Pediatrics, The Second Hospital of Jilin University, Changchun, China.

Received: 13 April 2016 Accepted: 13 September 2016

Published online: 20 September 2016

References

- Li F, Zhou Y, Li S, Jiang F, Jin X, Yan C, Tian Y, Zhang Y, Tong S, Shen X. Prevalence and risk factors of childhood allergic diseases in eight metropolitan cities in China: a multicenter study. *BMC Public Health*. 2011;11:437.
- Paaso EM, Jaakkola MS, Rantala AK, Hugg TT, Jaakkola JJ. Allergic diseases and asthma in the family predict the persistence and onset-age of asthma: a prospective cohort study. *Respir Res*. 2014;15:152.
- Gough H, Grabenhenrich L, Reich A, Eckers N, Nitsche O, Schramm D, Beschoner J, Hoffmann U, Schuster A, Bauer CP, Forster J, Zepp F, Lee YA, Bergmann RL, Bergmann KE, Wahn U, Lau S, Keil T, group MASs. Allergic multimorbidity of asthma, rhinitis, and eczema over 20 years in the German birth cohort MAS. *Pediatr Allergy Immunol*. 2015;26:431–7.
- Schaub B, Campo M, He H, Perkins D, Gillman MW, Gold DR, Weiss S, Lieberman E, Finn PW. Neonatal immune responses to TLR2 stimulation: influence of maternal atopy on Foxp3 and IL-10 expression. *Respir Res*. 2006;7:40.
- Schaub B, Liu J, Hoppler S, Haug S, Sattler C, Lluís A, Illi S, von Mutius E. Impairment of T-regulatory cells in cord blood of atopic mothers. *J Allergy Clin Immunol*. 2008;121:1491–9. 1499 e1–13.
- Loo EX, Shek LP, Goh A, Teoh OH, Chan YH, Soh SE, Saw SM, Kwek K, Gluckman PD, Godfrey KM, Chong YS, Lee BW, Van Bever HP. Atopic dermatitis in early life: evidence for at least three phenotypes? results from the GUSTO Study. *Int Arch Allergy Immunol*. 2015;166:273–9.
- Strickland DH, Holt PG. T regulatory cells in childhood asthma. *Trends Immunol*. 2011;32:420–7.
- Kwiatk M, Geca T, Krzyzanowski A, Malec A, Kwasniewska A. Peripheral dendritic cells and CD4+ CD25+ Foxp3+ regulatory T cells in the first trimester of normal pregnancy and in women with recurrent miscarriage. *PLoS One*. 2015;10:e0124747.
- Williams LM, Rudensky AY. Maintenance of the Foxp3-dependent developmental program in mature regulatory T cells requires continued expression of Foxp3. *Nat Immunol*. 2007;8:277–84.
- Shimizu J, Yamazaki S, Takahashi T, Ishida Y, Sakaguchi S. Stimulation of CD25 (+) CD4 (+) regulatory T cells through GITR breaks immunological self-tolerance. *Nat Immunol*. 2002;3:135–42.
- Li N, Workman CJ, Martin SM, Vignali DA. Biochemical analysis of the regulatory T cell protein lymphocyte activation gene-3 (LAG-3; CD223). *J Immunol*. 2004;173:6806–12.
- Nicolson KS, O'Neill EJ, Sundstedt A, Streeter HB, Minaee S, Wraith DC. Antigen-induced IL-10+ regulatory T cells are independent of CD25+ regulatory cells for their growth, differentiation, and function. *J Immunol*. 2006;176:5329–37.
- Schaub B, Liu J, Höppler S, Schleich I, Huehn J, Olek S, Wiczorek G, Illi S, von Mutius E. Maternal farm exposure modulates neonatal immune mechanisms through regulatory T cells. *J Allergy Clin Immunol*. 2009;123:774–82. e5.
- Lluís A, Depner M, Gaugler B, Saas P, Casaca VI, Raedler D, Michel S, Tost J, Liu J, Genuneit J, Pfefferle P, Roponen M, Weber J, Braun-Fahrlander C, Riedler J, Lauener R, Vuitton DA, Dalphin JC, Pekkanen J, von Mutius E, Schaub B. Protection against allergy: study in rural environments Study group. Increased regulatory T-cell numbers are associated with farm milk exposure and lower atopic sensitization and asthma in childhood. *J Allergy Clin Immunol*. 2014;133:551–9.
- Schaub B, Bellou A, Gibbons FK, German V, Campo M, He H, Liang Y, Gillman MW, Gold D, Weiss ST, Perkins DL, Finn PW. TLR2 and TLR4 stimulation differentially induce cytokine secretion in human neonatal, adult, and murine mononuclear cells. *J Interferon Cytokine Res*. 2004;24: 543–52.
- Global Initiative for Asthma: Global strategy for asthma management and prevention. Updated, 2011. (<http://ginasthma.org/>).

17. Brozek JL, Bousquet J, Baena-Cagnani CE, Bonini S, Canonica GW, Casale TB, van Wijk RG, Ohta K, Zuberbier T, Schunemann HJ, Global A, Asthma European N, Grading of Recommendations Assessment D, Evaluation Working G. Allergic rhinitis and its impact on asthma (ARIA) guidelines: 2010 revision. *J Allergy Clin Immunol*. 2010;126:466–76.
18. Baron SE, Cohen SN, Archer CB, British Association of D, Royal College of General P. Guidance on the diagnosis and clinical management of atopic eczema. *Clin Exp Dermatol*. 2012;37 Suppl 1:7–12.
19. Global Initiative for Asthma. Global strategy for the diagnosis and management of asthma in children 5 years and younger. 2009. (Assessed May 2009, at <http://ginasthma.org/>).
20. Schram D, Doekes G, Boeve M, Douwes J, Riedler J, Ublagger E, von Mutius E, Budde J, Pershagen G, Nyberg F, Alm J, Braun-Fahrlander C, Waser M, Brunekreef B, Group PS. Bacterial and fungal components in house dust of farm children, Rudolf Steiner school children and reference children—the PARSIFAL study. *Allergy*. 2005;60:611–8.
21. Prescott SL, Clifton V. Asthma and pregnancy: emerging evidence of epigenetic interactions in utero. *Curr Opin Allergy Clin Immunol*. 2009;9:417–26.
22. Dick S, Friend A, Dynes K, AlKandari F, Doust E, Cowie H, Ayres JG, Turner SW. A systematic review of associations between environmental exposures and development of asthma in children aged up to 9 years. *BMJ Open*. 2014;4:e006554.
23. Karvonen AM, Hyvarinen A, Gehring U, Korppi M, Doekes G, Riedler J, Braun-Fahrlander C, Bitter S, Schmid S, Keski-Nisula L, Roponen M, Kaulek V, Dalphin JC, Pfefferle PI, Renz H, Buchele G, von Mutius E, Pekkanen J, Group PS. Exposure to microbial agents in house dust and wheezing, atopic dermatitis and atopic sensitization in early childhood: a birth cohort study in rural areas. *Clin Exp Allergy*. 2012;42:1246–56.
24. Liu J, Radler D, Illi S, Klucker E, Turan E, von Mutius E, Kabesch M, Schaub B. TLR2 polymorphisms influence neonatal regulatory T cells depending on maternal atopy. *Allergy*. 2011;66:1020–9.
25. Ege MJ, Bieli C, Frei R, van Strien RT, Riedler J, Ublagger E, Schram-Bijkerk D, Brunekreef B, van Hage M, Scheynius A, Pershagen G, Benz MR, Lauener R, von Mutius E, Braun-Fahrlander C, Parsifal Study t. Prenatal farm exposure is related to the expression of receptors of the innate immunity and to atopic sensitization in school-age children. *J Allergy Clin Immunol*. 2006;117:817–23.
26. Conrad ML, Ferstl R, Teich R, Brand S, Blumer N, Yildirim AO, Patrascan CC, Hanuszkiewicz A, Akira S, Wagner H, Holst O, von Mutius E, Pfefferle PI, Kirschning CJ, Garn H, Renz H. Maternal TLR signaling is required for prenatal asthma protection by the nonpathogenic microbe *Acinetobacter lwoffii* F78. *J Exp Med*. 2009;206:2869–77.
27. Brand S, Teich R, Dicke T, Harb H, Yildirim AO, Tost J, Schneider-Stock R, Waterland RA, Bauer UM, von Mutius E, Garn H, Pfefferle PI, Renz H. Epigenetic regulation in murine offspring as a novel mechanism for transmaternal asthma protection induced by microbes. *J Allergy Clin Immunol*. 2011;128:618–25. e1-7.
28. Thorburn AN, McKenzie CI, Shen S, Stanley D, Macia L, Mason LJ, Roberts LK, Wong CHY, Shim R, Robert R, Chevalier N, Tan JK, Marino E, Moore RJ, Wong L, McConville MJ, Tull DL, Wood LG, Murphy VE, Mattes J, Gibson PG, Mackay CR. Evidence that asthma is a developmental origin disease influenced by maternal diet and bacterial metabolites. *Nat Commun*. 2015;6.
29. Kalliomaki M, Salminen S, Poussa T, Isolauri E. Probiotics during the first 7 years of life: a cumulative risk reduction of eczema in a randomized, placebo-controlled trial. *J Allergy Clin Immunol*. 2007;119:1019–21.
30. Kuitunen M, Kukkonen K, Juntunen-Backman K, Korpela R, Poussa T, Tuure T, Haahtela T, Savilahti E. Probiotics prevent IgE-associated allergy until age 5 years in cesarean-delivered children but not in the total cohort. *J Allergy Clin Immunol*. 2009;123:335–41.
31. Victor JR, Fusaro AE, Duarte AJD, Sato MN. Preconception maternal immunization to dust mite inhibits the type I hypersensitivity response of offspring. *J Allergy Clin Immunol*. 2003;111:269–77.
32. Victor JR, Muniz BP, Fusaro AE, de Brito CA, Taniguchi EF, Duarte AJD, Sato MN. Maternal immunization with ovalbumin prevents neonatal allergy development and up-regulates inhibitory receptor Fc gamma R1b expression on B cells. *BMC Immunol*. 2010;11.

Submit your next manuscript to BioMed Central and we will help you at every step:

- We accept pre-submission inquiries
- Our selector tool helps you to find the most relevant journal
- We provide round the clock customer support
- Convenient online submission
- Thorough peer review
- Inclusion in PubMed and all major indexing services
- Maximum visibility for your research

Submit your manuscript at
www.biomedcentral.com/submit

