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Maternal diet in pregnancy and child's respiratory outcomes: an individual participant data meta-analysis of 18000 children

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Shareable abstract (@ERSpublications) A suboptimal maternal diet in pregnancy, as defined by a higher inflammatory potential or low quality of the diet, does not play an important role in the development of respiratory diseases in childhood https://bit.ly/38dj3jU

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Abstract

Rationale Severe fetal malnutrition has been related to an increased risk of respiratory diseases later in life, but evidence for the association of a suboptimal diet during pregnancy with respiratory outcomes in childhood is conflicting. We aimed to examine whether a pro-inflammatory or low-quality maternal diet during pregnancy was associated with child's respiratory health.

Methods We performed an individual participant meta-analysis among 18326 mother–child pairs from seven European birth cohorts. Maternal pro-inflammatory and low-quality diets were estimated by energy-adjusted Dietary Inflammatory Index (E-DII) and Dietary Approaches to Stop Hypertension (DASH) scores. Preschool wheezing and school-age asthma were measured using questionnaires and lung function by spirometry.

Results After adjustment for lifestyle and sociodemographic factors, we observed that a higher maternal E-DII score (a more pro-inflammatory diet) during pregnancy was associated only with a lower forced vital capacity (FVC) in children (z-score difference -0.05, 95% CI -0.08– -0.02, per interquartile range increase). No linear associations of the maternal E-DII or DASH score with child's wheezing or asthma were observed. In an exploratory examination of the extremes, a very low DASH score (<10th percentile) (a very low dietary quality) was associated with an increased risk of preschool wheezing and a low forced expiratory volume in 1 s/FVC (z-score <-1.64) (OR 1.20, 95% CI 1.06–1.36 and z-score difference 1.40,

95% CI 1.06–1.85, compared to \geq 10th percentile), with corresponding population attributable risk fractions of 1.7% and 3.3%, respectively.

Conclusion The main results from this individual participant data meta-analysis do not support the hypothesis that maternal pro-inflammatory or low-quality diet in pregnancy are related to respiratory diseases in childhood.

Introduction

Asthma is a common disorder in childhood, and is associated with respiratory health problems in adulthood [1, 2]. Therefore, it is important to identify early-life modifiable risk factors. Fetal exposure to a suboptimal diet during pregnancy might affect the maturation of the lungs and immune system, leading to a lower lung function and a higher risk of wheezing and asthma in childhood [3]. Severe malnutrition in pregnancy has previously been associated with an increased risk of respiratory diseases later in life [4]. Studies examining maternal diet during pregnancy and childhood respiratory health mainly focused on the intake of specific nutrients or food groups [5]. However, examining the overall diet might take the interactions within the diet into account and be better translatable to dietary guidelines [6]. The energy-adjusted Dietary Inflammatory Index (E-DII) [7] and Dietary Approaches to Stop Hypertension (DASH) [8] provide dietary scores for the inflammatory potential and overall quality of the diet, respectively. Cohort studies showed that a higher maternal E-DII score in pregnancy was associated with a higher risk of an early wheeze trajectory and a lower mid-expiratory flow or a higher risk of asthma in childhood [9, 10]. The relationship of the DASH score with respiratory outcomes has been studied only in adults, where a DASH-promoting behavioural intervention seemed to improve asthma control [11]. To date, a pooled analysis across cohorts which examines the relationship of the inflammatory potential and overall quality of maternal diet during pregnancy with child's respiratory health is lacking.

We performed an individual participant data meta-analysis among 18326 children, participating in seven European birth cohort studies. We assessed the associations of maternal diet during pregnancy, as summarised by the E-DII and DASH score, with preschool wheezing, school-age asthma and lung function, and estimated the impact of these associations on the general population by calculating the population attributable fraction.

Methods

This meta-analysis was performed among seven European prospective birth cohorts participating in the ALPHABET consortium, which aims to examine the early-life nutritional programming of noncommunicable diseases (supplementary methods) [12, 13]. We included 18 326 mother–child pairs for the current analyses (supplementary methods).

Maternal diet

Information obtained from food frequency questionnaires (FFQs) before or during pregnancy was used to generate the maternal E-DII and DASH scores (supplementary tables S1 and S2), as described previously (supplementary methods) [7, 13]. To control for the effect of the total energy intake, the E-DII, calculated per 1000 kcal of food consumed, was used. The E-DII in ALPHABET was generated from 20–28 dietary parameters, out of 44 possible parameters. A higher E-DII score characterises a more pro-inflammatory diet [7]. For the seven cohorts in the ALPHABET project, a DASH score was generated. This score was composed of eight food components, based mainly on the Fung method with a scoring system based on quintile rankings in each cohort [8, 13]. A lower DASH score characterises a lower dietary quality. For the main analyses, we used data collected at one time point, preferably in early pregnancy (first or second trimester) (Generation R, Lifeways, REPRO_PL, ROLO and the Southampton Women's Survey (SWS) cohorts), since this period is of specific importance for lung disease development later in life [14], or, if not available, in late pregnancy (third trimester) (Avon Longitudinal Study of Parents and Children (ALSPAC) and EDEN cohorts).

Respiratory health

Data on preschool wheezing and school-age asthma was mainly obtained from questions adapted from the International Study of Asthma and Allergies in Childhood questionnaire [15]. We defined preschool wheezing as "ever reported wheezing during the first 4 years of life" and school-age asthma as "asthma diagnosis reported between 5 and 10 years" [16]. Cohort-specific information is shown in the supplementary methods and supplementary table S1. All cohorts obtained lung function measures by spirometry according to the American Thoracic Society/European Respiratory Society guidelines [17]. Lung function measures included forced expiratory volume in 1 s (FEV₁), forced vital capacity (FVC), FEV₁/FVC ratio and forced expiratory flow at 25–75% of FVC (FEF_{25–75}), and were converted into sex-, age-, height- and ethnicity-adjusted z-scores based on the Global Lung Initiative reference values [18].

Covariates

Information on lifestyle- and sociodemographic-related confounders, intermediates and effect modifiers was mainly obtained by questionnaires or clinical examinations at the research centre (supplementary methods and table S1).

Statistical analyses

Dietary scores were analysed as continuous variables to study the linear associations, and additionally as dichotomous variables to explore the effect of the extremes. We first conducted one-stage meta-analyses by using multilevel linear regression models or multilevel logistic regression models to study the associations of the maternal E-DII and DASH scores with child's respiratory outcomes. In these models, individual participant data from all cohorts were combined and modelled simultaneously, taking into account clustering of participants within cohorts [19]. We included a random intercept at cohort level, which allows intercepts to vary across cohorts. More information on the used models is provided in the supplementary methods. As explorative analyses to examine the effect of an extreme adverse diet in pregnancy, we additionally studied the dichotomous relationships and examined the associations of a very high E-DII score (>90th percentile) or low DASH score (<10th percentile) with wheezing and asthma, and with lung function below the lower limit of normal (LLN) (\leq 5th percentile, equals z-score of -1.64). The highest and lowest 10th percentile cut-off for the dietary scores is a common epidemiological approach, in the absence of clinical cut-offs. If consistent associations were observed, we subsequently calculated the population attributable risk fraction based on the adjusted odds ratio and the prevalence of a high E-DII or low DASH score, which indicates the proportion of wheezing, asthma or lung function measures below the LLN attributable to a high E-DII or low DASH score [20]. We considered the linear confounder models as the main models and applied several additional analyses to these models, as described in the supplementary methods.

P-values are two-tailed and statistical significance was defined at p<0.05. We did not adjust for multiple testing, since respiratory outcomes are strongly interrelated [21]. Statistical analyses were performed using SPSS version 24.0 (IBM, Armonk, NY, USA) and RevMan version 5.3 (Nordic Cochrane Centre, Copenhagen, Denmark) and R version 3.6.1 ("mediation" package; www.r-project.org).

Ethical approvals

Specific cohort approvals are for ALSPAC by the ALSPAC ethics and law committee (IRB00003312) and local research ethics committees; informed consent for the use of data collected *via* questionnaires and clinics was obtained from participants following the recommendations of the ALSPAC ethics and law committee at the time; for EDEN by the ethics committee and the Commission Nationale Informatique et Liberté, the French data privacy institution; for Generation R by the medical ethical committee of the Erasmus Medical Center (Rotterdam, the Netherlands); for Lifeways by the University College Dublin research ethics committee and St Vincent's University Hospital research ethics committee; for REPRO_PL by the ethical committee of the Nofer Institute of Occupational Medicine (Łódź, Poland; decision numbers 7/2007, 3/2008, 22/2014); for ROLO by the ethics committee of the National Maternity Hospital (Dublin, Ireland); and for SWS by the Southampton and South West Hampshire research ethics committee.

Results

Subject characteristics

Table 1 shows the main characteristics, maternal dietary scores and child's respiratory outcomes of the cohorts, and supplementary tables S3 and S4 show the corresponding information on maternal and child-related baseline characteristics. The median age of the included children at lung function measurement was 8.6 years (95% range 5.4–10.2 years). Of all participants, 51.9% (n=8018) had preschool wheezing and 15.6% (n=2193) had school-age asthma. The correlation between the E-DII and DASH score was moderate (Pearson r range -0.49 to -0.60, p<0.001).

Maternal E-DII and DASH score and child's respiratory outcomes

Table 2 shows that after adjustment for confounders, only an association of a higher maternal E-DII score during pregnancy with a lower FVC in the children was observed (z-score difference -0.05, 95% CI -0.08--0.02). A lower DASH score was not associated with preschool wheezing, school-age asthma or lung function measures. We observed no consistent associations for both the maternal E-DII and DASH score with FEF₂₅₋₇₅ (results not shown).

When exploratively examining the extremes, we observed no associations of a very high maternal E-DII score (>90th percentile) with child's respiratory outcomes compared to a normal maternal E-DII score (\leq 90th percentile) after adjustment for confounders (figure 1). A very low DASH score (<10th percentile)

	ALSPAC (UK)	EDEN (France)	Generation R (the Netherlands)	Lifeways (Ireland)	REPRO_PL (Poland)	ROLO (Ireland)	SWS (UK)
Total participants	10130	843	4263	224	523	301	2042
Inclusion years	1990–1992	2003–2006	2002–2006	2001–2003	2007–2011	2007–2011	1998–2002
Pregnancy							
FFQ (GA in weeks) [#]	32	24–28 Birth	<24	12–16	20–24	≼28	PP 11 34
FFQ assessed period	LP	PP LP	EP	EP	EP	EP	PP EP LP
E-DII score	0.51±1.82	0.76±1.65	-0.43±1.10	-0.12±1.43	-1.10±1.54	0.12±1.74	0.27±1.49
DASH score	24.1±4.0	24.3±4.1	24.4±4.4	25.2±4.5	24.1±4.4	24.2±4.1	24.1±4.3
Preschool wheezing							
Participants	9313	840	2876	NA	370	NA	2037
Age (years) [#]	0–3.5	0-4	1–4	NA	1–2	NA	0–3
Yes % (n)	54.4 (5070)	36.8 (309)	49.7 (1429)	NA	18.4 (68)	NA	56.1 (1142)
School-age asthma							
Participants n	7506	842	3510	224	275	301	1421
Age (years) [#]	8	5–8	9	9	7–8	5	5
Yes % (n)	20.3 (1525)	12.1 (102)	8.9 (312)	5.4 (12)	6.2 (17)	7.6 (23)	14.2 (202)
Lung function							
Participants n	5766	838	3651	NA	264	NA	730
Age (years)	8.6 (8.3–9.5)	5.6 (5.4–6.0)	9.8 (9.4–10.7)	NA	7.2 (7.0–8.8)	NA	6.5 (6.2–6.9)
FEV ₁ (z-score)	-0.03±1.01	-0.70±1.45	0.17±0.98	NA	-0.32±1.74	NA	0.09±0.98
FVC (z-score)	-0.04±1.02	-1.00 ± 1.48	0.21±0.93	NA	-0.44±1.85	NA	0.15±1.06
FEV ₁ /FVC (z-score)	0.05±1.07	0.87±1.06	-0.11±0.95	NA	0.30±1.25	NA	-0.08±1.06
FEF ₂₅₋₇₅ (z-score)	-0.15±1.02	-0.39±1.09	0.43±1.08	NA	-0.14 ± 1.01	NA	-0.25±0.92

Data are presented as n, mean±sp, median (95% range) or valid percentages (absolute numbers). ALSPAC: Avon Longitudinal Study of Parents and Children; SWS: Southampton Women's Survey; FFQ: food frequency questionnaire; GA: gestational age; E-DII: energy-adjusted Dietary Inflammatory Index; DASH: Dietary Approaches to Stop Hypertension; FEV₁: forced expiratory volume in 1 s; FVC: forced vital capacity; FEF₂₅₋₇₅: forced expiratory flow at 25–75% of FVC; PP: pre-pregnancy; LP: late pregnancy (third trimester); EP: early pregnancy (first or second trimester); NA: not available. [#]: time period of questionnaire assessment.

was associated with a higher risk of preschool wheezing and an FEV₁/FVC below the LLN, and borderline associated with a higher risk of asthma (OR 1.20, 95% CI 1.06–1.36; z-score 1.40, 95% CI 1.06–1.85; OR 1.17, 95% CI 1.00–1.39, respectively, as compared to a DASH score \geq 10th percentile). The estimated proportions of wheezing, FEV₁/FVC below the LLN and asthma attributable to a low DASH score were 1.7%, 3.3% and 1.4%, respectively.

Additional analyses

Additional adjustment for early growth factors, lower respiratory tract infections, child's body mass index (BMI) or child's E-DII score did not materially change the effects (results not shown). Further mediation analyses showed that early growth factors and child's E-DII score only explained 6.2% (95% CI 2.3-21.0%) and 17.8% (95% CI 2.6-48.0%) of the association of the E-DII score with FVC. We observed a consistent interaction between the maternal DASH score and child's sex (interaction terms p-value range <0.001–0.549), but not between maternal dietary scores and child's atopic predisposition. After stratification by sex, no consistent differences between boys and girls in the association of the maternal E-DII or DASH score with child's respiratory outcomes were observed (supplementary table S5). The two-stage random-effect meta-analyses indicated at most moderate heterogeneity (range I^2 0–52%) and similar effects as the one-stage meta-analyses (supplementary figures S2 and S3). When we examined the dietary scores per time period of assessment in pregnancy, directions of the associations with respiratory outcomes were similar for all time periods (supplementary table S6). Examining the associations of maternal dietary scores with lung function measures in age groups of children showed that among children aged \geq 8 years, a higher maternal E-DII score was associated with a lower FEV₁ and FVC, and a lower maternal DASH score with a lower FEV_1 (supplementary table S7). We repeated the main models restricted to complete cases, to mothers with a European birthplace/ethnic background, and excluding one **TABLE 2** Linear associations of maternal energy-adjusted Dietary Inflammatory Index (E-DII) and Dietary Approaches to Stop Hypertension (DASH) score with preschool wheezing and school-age asthma and lung function

	Preschool wheezing OR (95% CI)	School-age asthma OR (95% CI)	FEV ₁ z-score change (95% CI)	FVC z-score change (95% CI)	FEV1/FVC z-score change (95% CI)
Participants, n	15436	14079	11249	11249	11249
E-DII score, per IQR increase					
Basic model [#]	1.14 (1.09–1.20)**	1.07 (1.00-1.15)*	-0.03 (-0.05-0.00)	-0.04 (-0.070.01)*	0.02 (-0.01-0.05)
p-value	< 0.001	0.047	0.082	0.010	0.11
Confounder model [¶]	1.02 (0.97-1.07)	1.00 (0.93-1.07)	-0.03 (-0.06-0.00)	-0.05 (-0.080.02)**	0.03 (-0.00-0.06)
p-value	0.484	0.883	0.057	0.003	0.051
DASH score, per IQR decrease					
Basic model ⁺	1.15 (1.10-1.21)**	1.16 (1.08-1.24)**	-0.01 (-0.04-0.02)	-0.01 (-0.04-0.02)	-0.02 (-0.05-0.01)
p-value	< 0.001	< 0.001	0.421	0.865	0.122
Confounder model [¶]	1.04 (0.98-1.09)	1.06 (0.99-1.14)	-0.02 (-0.05-0.01)	-0.01 (-0.04-0.02)	-0.02 (-0.05-0.01)
p-value	0.180	0.123	0.250	0.506	0.170

Values are derived from multilevel logistic or linear regression models and reflect odds ratios or changes in z-scores with their corresponding 95% confidence intervals per interquartile range (IQR) increase in the E-DII score or per IQR decrease in the DASH score. Bold type represents statistical significance. FEV₁: forced expiratory volume in 1 s; FVC: forced vital capacity. \ddagger : adjusted for child's sex; \P : additionally adjusted for maternal body mass index, education, birthplace/ethnic background, smoking during pregnancy and parity and child's breastfeeding; \ddagger : adjusted for child's sex and maternal energy intake. \ddagger : p<0.05; \ddagger : p<0.01.

cohort at a time, and mainly observed similar sizes and directions of the effect estimates (supplementary tables S7, S8a and S8b). Excluding only the intervention arm of the ROLO study did not materially change our results (results not shown).

Discussion

In this individual participant data meta-analysis among 18326 children from seven European birth cohorts, we observed that only a more pro-inflammatory diet during pregnancy was associated with a lower FVC in childhood. When studying the extremes, a very low maternal dietary quality was associated with a higher risk of preschool wheezing and a FEV_1/FVC below the LLN in the children, and borderline higher risk of school-age asthma.

Comparison with previous studies

To our best knowledge, our study is the first individual participant meta-analysis of prospective birth cohorts to examine the associations of the maternal E-DII score with child's respiratory outcomes. Previous studies showed that a higher E-DII score during pregnancy or in childhood was associated with a higher risk of early wheezing, wheezing trajectories or asthma, and a lower FEF_{25-75} , but not with other lung function measures or in high-risk children only [9, 10, 22]. Differences between results of these studies and our meta-analysis might be due to other definitions of respiratory outcomes. Asthma is difficult to diagnose in children aged <5 years, and the wheezing pathogenesis including the role of specific viruses in the development of a lower lung function and asthma might differ between age periods [23, 24]. Therefore, we used both preschool wheezing and school-age asthma as outcomes. The association of the E-DII with a lower FVC did not attenuate after additional adjustment for lower respiratory tract infections. However, further studies on the effect of the maternal E-DII score on harmonised longitudinal asthma-symptom phenotypes in the children are needed.

Our study showed no linear associations of the maternal DASH score with child's respiratory outcomes, but a very low-quality diet, defined by a very low DASH score capturing the intake of multiple food groups, was associated with a higher risk of wheezing and airway obstruction. A Mediterranean diet in pregnancy partly overlaps with the high DASH score diet (DASH diet) and has been associated with a lower risk of wheezing, whereas other dietary patterns, defined based on principal component analysis, were not associated with respiratory outcomes [25, 26]. The advantage of the DASH diet, as compared to these approaches, is that it might better reflect the dietary habits in a non-Mediterranean population and is easy to translate into public health guidelines [6].

Interpretation of the results

The E-DII score takes many food parameters into account, of which the main pro-inflammatory components are trans fat, saturated fat and cholesterol, and the main anti-inflammatory components are

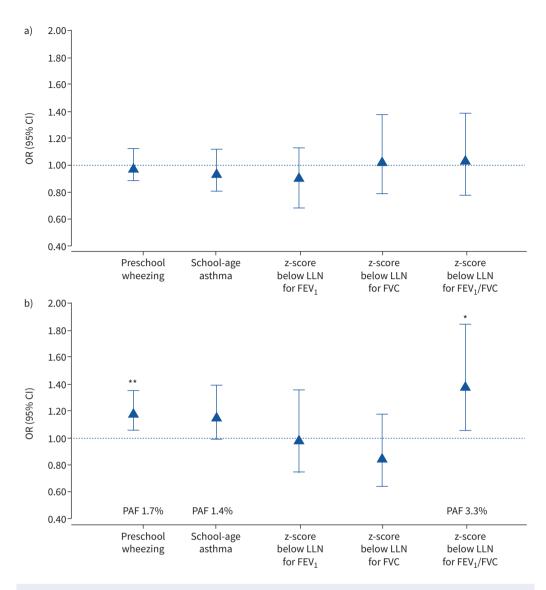


FIGURE 1 Associations of a) high energy-adjusted Dietary Inflammatory Index (E-DII) and b) low Dietary Approaches to Stop Hypertension (DASH) score in pregnancy with preschool wheezing and school-age asthma and lung function. Values are derived from multilevel logistic regression models and reflect odds ratios or changes in z-scores with 95% confidence intervals as compared to the reference group (\leq 90th percentile for the E-DII score and \geq 10th percentile for the DASH score). The population attributable risk fractions (PAFs) indicate the proportion of preschool wheezing, school-age asthma or forced expiratory volume in 1 s (FEV₁)/ forced vital capacity (FVC) below the lower limit of normal (LLN) attributable to a low DASH score. LLN is defined as z-score for lung function outcome <1.64. The models are adjusted for maternal body mass index, education, birthplace/ethnic background, smoking during pregnancy and parity, and child's sex and breastfeeding, and the models with DASH as exposure are additionally adjusted for maternal energy intake. *: p<0.05; **: p<0.01.

nutrients derived from fruits and vegetables and n-3 fatty acids [7]. Underlying mechanisms might be that a high-fat maternal diet leads to fetal lung inflammation and remodelling, which could make the lungs more susceptible to developing asthma later in childhood [27]. Obesity is another factor that is associated with inflammation [28], and this might be the reason why the associations with wheezing and asthma attenuated after adjustment for lifestyle factors including maternal BMI. In addition, an indirect effect through early growth factors may play a role in the association of the maternal E-DII score with child's FVC as shown by the moderate percentage of change of the effect estimates [29]. However, the effect of the association of the maternal E-DII score with child's FVC was small, and might therefore reflect a subclinical change or chance finding.

A DASH diet is mainly characterised by a high intake of fruits and vegetables, which are rich in antioxidants, and a low intake of added sugars and sodium [13]. Antioxidants might make the lungs less vulnerable to oxidative stress, and thereby may reduce the risk of asthma and airway obstruction [30]. The DASH diet has also been shown to lower blood pressure [31]. A higher blood pressure in pregnancy, which might reflect poorer vascular health, has been associated with a higher risk of wheezing and asthma and a lower FEV₁/FVC in children [32, 33]. We only observed in our explorative analyses that a very low DASH score was associated with a higher risk of preschool wheezing, airway obstruction and borderline with asthma. However, we were not able to take the DASH score of child's current diet into consideration. Additionally, the effect of maternal diet on child's respiratory outcomes may differ between different periods of pregnancy. Since lung development already starts in the fourth week of pregnancy, adverse exposures in early pregnancy are considered to be specifically important for lung disease development later in life [14]. Further research is needed to understand the effect of maternal diet at different gestational ages during pregnancy and in different periods of early life after birth on lung development across the life course.

Although previous studies showed that high maternal intake of single nutrients including vitamin D and n-3 fatty acids may be beneficial for child's respiratory health [34, 35], we observed no consistent association of the maternal E-DII or DASH score with respiratory outcomes. This suggests that specific supplements may have greater importance than a balanced diet for asthma development.

The moderate correlation between the E-DII and DASH scores suggest that these scores partly represent different factors of the diet. The scores differ in concept as the E-DII is mainly nutrient based and focuses on the inflammatory effects of the diet, whereas the DASH score defines the overall quality of the diet based on food components. Our hypothesis for the effect of maternal diet on child's respiratory outcomes was based on a population with an extremely adverse diet [4]. The distribution of maternal diet during pregnancy in Western countries might be within optimal ranges, and any potential adverse effect might lay in the extremes. Therefore, we studied the extremes by using a common epidemiological cut-off approach, the highest and lowest 10th percentiles, since clinical cut-offs are lacking. Categorisation is prone to bias and our analyses are explorative and should be considered as hypothesis-generating. Therefore, results suggesting that the associations of an adverse diet with clinically relevant respiratory outcomes only exist in those exposed to an extremely adverse diet should be carefully interpreted. In addition, if we assume that these relationships are causal, the average proportions of wheezing, asthma and an FEV₁/FVC below the LLN attributable to a low DASH are tenuous. Targeting maternal diet, in addition to other lifestyle and sociodemographic factors, to improve child's respiratory outcomes could be the subject for future intervention trials, but in a population of mothers with an extremely adverse diet only.

Strengths and limitations

A major strength of this meta-analysis is the use of individual participant data. This resulted in a large sample size and enabled us to harmonise the data, and to reduce the risk of publication bias. However, some limitations do apply. First, dietary scores as well as wheezing and asthma were defined based on questionnaires, which could have led to reporting errors. In addition, FFQs may not adequately assess the intake of specific nutrients, such as sodium, a component of the DASH score or the specific food parameters for the E-DII score. Furthermore, missing data in the FFQs might have biased the estimation of the dietary scores. Clearly, we cannot know the effect of foods eaten that are not on the FFQ. Nevertheless, most cohorts used validated questionnaires [13, 15]. Second, although the dietary scores were calculated in all cohorts according to the same methods, there were differences in the included food parameters, length and content of the FFQs, assessed time periods in pregnancy and assessment years. However, two-stage meta-analyses gave similar results and showed limited heterogeneity between the cohort estimates. Although none of the cohorts had information on all 44 possible parameters for the E-DII score, a previous validation study showed that an DII score based on 28 parameters had a good predictive ability, and an additional study showed that a score based on 17 parameters was related to inflammatory markers [36, 37]. Thus, our E-DII score gives a valid, if imprecise, estimation of the inflammatory potential of the diet. We were not able to take potential changes in a mother's diet due to seasonal variation or food aversions into consideration, and we did not have information for all cohorts on the exact gestational age at which diet was assessed. However, FFQs are considered an adequate method to measure the usual dietary intake over an extensive period of time, and dietary patterns are suggested not to change much during pregnancy [38, 39]. Thus, our dietary measurements across cohorts were appropriate and support our findings. Third, although the participating cohorts were carefully selected based on a priori power calculations, data availability, and spread throughout Europe, most participants come from two cohorts and have a European birthplace/ethnic background. Therefore, results may not be generalisable to mothers in other geographical regions. Fourth, we did not measure changes in the associations of maternal diet with child's respiratory outcomes over time. Fifth, to date, no validated method to calculate child's DASH score is available. Although results remained similar after adjustment for child's BMI, potential mediating effect of child's DASH score cannot be fully ruled out. Last, we adjusted for major potential confounders, but, as in all observational studies, residual confounding due to unmeasured or insufficiently harmonised factors, such as other sociodemographic factors, environmental pollution, the use of supplements or medication in pregnancy or the duration of breastfeeding remains an issue. Future randomised controlled intervention trials might minimise the risk of confounding factors influencing the results, but should be carefully considered given the absence of a consistent association in our current study.

Conclusion

A more pro-inflammatory diet of mothers during pregnancy was only related to a lower FVC in childhood. Both the inflammatory potential and quality of the diet were not consistently related to wheezing or asthma in childhood. The main results from this individual participant data meta-analysis do not support the hypothesis that maternal pro-inflammatory or low-quality diet in pregnancy are related to respiratory diseases in childhood.

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References

- 1 Asher MI, Montefort S, Björkstén B, *et al.* Worldwide time trends in the prevalence of symptoms of asthma, allergic rhinoconjunctivitis, and eczema in childhood: ISAAC phases one and three repeat multicountry cross-sectional surveys. *Lancet* 2006; 368: 733–743.
- 2 Bui DS, Lodge CJ, Burgess JA, *et al.* Childhood predictors of lung function trajectories and future COPD risk: a prospective cohort study from the first to the sixth decade of life. *Lancet Respir Med* 2018; 6: 535–544.
- **3** Devereux G, Seaton A. Diet as a risk factor for atopy and asthma. *J Allergy Clin Immunol* 2005; 115: 1109–1117.
- 4 Lopuhaä CE, Roseboom TJ, Osmond C, *et al.* Atopy, lung function, and obstructive airways disease after prenatal exposure to famine. *Thorax* 2000; 55: 555–561.
- 5 Beckhaus AA, Garcia-Marcos L, Forno E, *et al.* Maternal nutrition during pregnancy and risk of asthma, wheeze, and atopic diseases during childhood: a systematic review and meta-analysis. *Allergy* 2015; 70: 1588–1604.
- 6 Hu FB. Dietary pattern analysis: a new direction in nutritional epidemiology. Curr Opin Lipidol 2002; 13: 3–9.
- 7 Shivappa N, Steck SE, Hurley TG, *et al.* Designing and developing a literature-derived, population-based dietary inflammatory index. *Public Health Nutr* 2014; 17: 1689–1696.

- 8 Fung TT, Rimm EB, Spiegelman D, *et al.* Association between dietary patterns and plasma biomarkers of obesity and cardiovascular disease risk. *Am J Clin Nutr* 2001; 73: 61–67.
- 9 Hanson C, Rifas-Shiman SL, Shivappa N, *et al.* Associations of prenatal dietary inflammatory potential with childhood respiratory outcomes in Project Viva. *J Allergy Clin Immunol Pract* 2020; 8: 945–952.
- **10** Chen LW, Lyons B, Navarro P, *et al.* Maternal dietary inflammatory potential and quality are associated with offspring asthma risk over 10-year follow-up: the Lifeways Cross-Generation Cohort Study. *Am J Clin Nutr* 2020; 111: 440–447.
- 11 Ma J, Strub P, Lv N, *et al.* Pilot randomised trial of a healthy eating behavioural intervention in uncontrolled asthma. *Eur Respir J* 2016; 47: 122–132.
- 12 Phillips CM, Chen LW, Heude B, *et al.* Dietary inflammatory index and non-communicable disease risk: a narrative review. *Nutrients* 2019; 11: 1873.
- **13** Aubert AM, Forhan A, de Lauzon-Guillain B, *et al.* Deriving the Dietary Approaches to Stop Hypertension (DASH) score in women from seven pregnancy cohorts from the European ALPHABET consortium. *Nutrients* 2019; 11: 2706.
- 14 Mullassery D, Smith NP. Lung development. Semin Pediatr Surg 2015; 24: 152–155.
- **15** Asher MI, Keil U, Anderson HR, *et al.* International Study of Asthma and Allergies in Childhood (ISAAC): rationale and methods. *Eur Respir J* 1995; 8: 483–491.
- 16 Sonnenschein-van der Voort AM, Arends LR, de Jongste JC, et al. Preterm birth, infant weight gain, and childhood asthma risk: a meta-analysis of 147,000 European children. J Allergy Clin Immunol 2014; 133: 1317–1329.
- 17 Miller MR, Hankinson J, Brusasco V, et al. Standardisation of spirometry. Eur Respir J 2005; 26: 319–338.
- **18** Quanjer PH, Stanojevic S, Cole TJ, *et al.* Multi-ethnic reference values for spirometry for the 3–95-yr age range: the global lung function 2012 equations. *Eur Respir J* 2012; 40: 1324–1343.
- 19 Debray TP, Moons KG, Abo-Zaid GM, *et al.* Individual participant data meta-analysis for a binary outcome: one-stage or two-stage? *PLoS One* 2013; 8: e60650.
- 20 Flegal KM, Graubard BI, Williamson DF. Methods of calculating deaths attributable to obesity. Am J Epidemiol 2004; 160: 331–338.
- 21 Rothman KJ. No adjustments are needed for multiple comparisons. Epidemiology 1990; 1: 43–46.
- 22 Han YY, Forno E, Shivappa N, *et al.* The Dietary Inflammatory Index and current wheeze among children and adults in the United States. *J Allergy Clin Immunol Pract* 2018; 6: 834–841.
- 23 Martinez FD, Wright AL, Taussig LM, *et al.* Asthma and wheezing in the first six years of life. *N Engl J Med* 1995; 332: 133–138.
- 24 van Meel ER, den Dekker HT, Elbert NJ, *et al.* A population-based prospective cohort study examining the influence of early-life respiratory tract infections on school-age lung function and asthma. *Thorax* 2018; 73: 167–173.
- 25 Shaheen SO, Northstone K, Newson RB, *et al.* Dietary patterns in pregnancy and respiratory and atopic outcomes in childhood. *Thorax* 2009; 64: 411–417.
- 26 Chatzi L, Torrent M, Romieu I, *et al.* Mediterranean diet in pregnancy is protective for wheeze and atopy in childhood. *Thorax* 2008; 63: 507–513.
- 27 Heyob KM, Mieth S, Sugar SS, et al. Maternal high-fat diet alters lung development and function in the offspring. Am J Physiol Lung Cell Mol Physiol 2019; 317: L167–L174.
- 28 Timpson NJ, Nordestgaard BG, Harbord RM, *et al.* C-reactive protein levels and body mass index: elucidating direction of causation through reciprocal Mendelian randomization. *Int J Obes* 2011; 35: 300–308.
- 29 Sen S, Rifas-Shiman SL, Shivappa N, *et al.* Dietary inflammatory potential during pregnancy is associated with lower fetal growth and breastfeeding failure: results from Project Viva. *J Nutr* 2016; 146: 728–736.
- 30 Caramori G, Papi A. Oxidants and asthma. *Thorax* 2004; 59: 170–173.
- **31** Appel LJ, Moore TJ, Obarzanek E, *et al.* A clinical trial of the effects of dietary patterns on blood pressure. *N Engl J Med* 1997; 336: 1117–1124.
- **32** Tang JR, Karumanchi SA, Seedorf G, *et al.* Excess soluble vascular endothelial growth factor receptor-1 in amniotic fluid impairs lung growth in rats: linking preeclampsia with bronchopulmonary dysplasia. *Am J Physiol Lung Cell Mol Physiol* 2012; 302: L36–L46.
- 33 Wilmink FA, den Dekker HT, de Jongste JC, *et al.* Maternal blood pressure and hypertensive disorders during pregnancy and childhood respiratory morbidity: the Generation R Study. *Eur Respir J* 2018; 52: 1800378.
- 34 Bisgaard H, Stokholm J, Chawes BL, *et al.* Fish oil-derived fatty acids in pregnancy and wheeze and asthma in offspring. *N Engl J Med* 2016; 375: 2530–2539.
- **35** Wolsk HM, Chawes BL, Litonjua AA, *et al.* Prenatal vitamin D supplementation reduces risk of asthma/ recurrent wheeze in early childhood: a combined analysis of two randomized controlled trials. *PLoS One* 2017; 12: e0186657.
- 36 Shivappa N, Steck SE, Hurley TG, et al. A population-based dietary inflammatory index predicts levels of C-reactive protein in the Seasonal Variation of Blood Cholesterol Study (SEASONS). Public Health Nutr 2014; 17: 1825–1833.

- **37** Shivappa N, Hébert JR, Rietzschel ER, *et al.* Associations between dietary inflammatory index and inflammatory markers in the Asklepios Study. *Br J Nutr* 2015; 113: 665–671.
- 38 Willett W. Nutritional epidemiology: issues and challenges. *Int J Epidemiol* 1987; 16: 312–317.
- **39** Crozier SR, Robinson SM, Godfrey KM, *et al.* Women's dietary patterns change little from before to during pregnancy. *J Nutr* 2009; 139: 1956–1963.