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Maternal intake of fatty acids during pregnancy and allergies in the offspring

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Abstract

Fatty acids (FA) are known to have a number of immunological effects and, accordingly, may play a role in the development of allergic diseases. We investigated the effect of maternal intake of FA during pregnancy on the risk of allergic rhinitis, wheeze and atopic eczema in children aged 5 years. The present study analysed data from the Finnish Type 1 Diabetes Prediction and Prevention Nutrition Study, a population-based birth cohort study with a 5-year follow-up. Complete information on maternal diet (assessed by a validated FFQ) and International Study of Asthma and Allergies in Childhood-based allergic outcomes was available for 2441 children. Cox proportional regression and logistic regression were used for the analyses. After adjusting for potential confounding variables, high maternal consumption of butter and butter spreads (hazard ratio (HR) 1.33; 95% CI 1.03, 1.71) and higher ratio of *n-6:n-3* FA (HR 1.37; 95% CI 1.07, 1.77) during pregnancy were associated with an increased risk of allergic rhinitis in the offspring by 5 years of age. High maternal intakes of total PUFA (HR 0.71; 95% CI 0.52, 0.96) and α -linolenic FA (HR 0.73; 95% CI 0.54, 0.98) were associated with a decreased risk of allergic rhinitis. However, these results lost their significance after adjustment for multiple comparisons. Overall, our data suggest that maternal consumption of butter, the ratio of *n-6:n-3* FA and intake of PUFA and α -linolenic FA during pregnancy may be potential determinants of allergic rhinitis in the offspring.

Key words: Allergic rhinitis: Atopic eczema: Children: Fatty acids: Maternal diet: Pregnancy: Wheeze

The increasing prevalence of allergic diseases in Western countries has partly been linked to changes in the modern diet⁽¹⁾, particularly changes in the intake of dietary fats⁽²⁾. Over the past few decades, consumption of foods rich in *n-6* PUFA has increased, whereas that of *n-3* PUFA has decreased. Arachidonic acid (AA), which is an *n-6* PUFA, is a precursor for pro-inflammatory eicosanoids, while *n-3* EPA, is a precursor for anti-inflammatory eicosanoids⁽³⁾. Changes in fatty acid consumption may shift the T-helper balance from type 1 to type 2, which, consequently, may increase the incidence of IgE-mediated allergic diseases^(3,4).

The results of studies on the association between dietary intake of fats and fatty acids (FA) during pregnancy and allergic diseases in the offspring are inconsistent. A recent Japanese cohort study has linked maternal intake of α -linolenic acid and DHA during pregnancy to a reduced risk of wheezing in 2-year-old children, while intake of *n-6* PUFA and linoleic acid was associated with an increased risk of eczema⁽⁵⁾. Maternal consumption of butter and margarine was not associated with atopy in the offspring in an Italian cohort study⁽⁶⁾. Maternal fish consumption decreased the risk of allergic diseases in the offspring^(6–10). In contrast, maternal

Abbreviations: AA, arachidonic acid; DIPP, Diabetes Prediction and Prevention; FA, fatty acids.

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fish consumption was not associated with asthma or asthma symptoms in the offspring in a Dutch follow-up study⁽¹¹⁾. Further research is required in this area to elucidate the putative role of dietary fats during pregnancy and early life in the development of allergies. In the present large cohort with a relatively long follow-up, we aimed at examining the effects of maternal intake of dietary fats and a wide range of FA during pregnancy on the development of allergic rhinitis, wheeze and atopic eczema in the offspring by the age of 5 years.

Materials and methods

Subjects

In 1994, the Finnish Type 1 Diabetes Prediction and Prevention (DIPP) study was started. It is a multidisciplinary population-based prospective cohort study that examines potential means to predict and prevent type 1 diabetes⁽¹²⁾. Newborn infants with human leucocyte antigen (HLA)-conferred susceptibility to type 1 diabetes are recruited from three university hospitals in Finland (Turku, Oulu and Tampere) and monitored at 3–12-month intervals for diabetes-associated antibodies, growth and environmental exposures. The present study was conducted according to the guidelines laid down in the Declaration of Helsinki and all procedures involving human subjects were approved by the ethics committees of the Universities of Oulu and Tampere. Written informed consent was obtained from the parents.

In September 1996 and October 1997, the DIPP Nutrition Study was started within the framework of the DIPP study in Oulu (Northern Finland) and Tampere (Southern Finland), respectively⁽¹³⁾, examining the relationship of maternal diet during pregnancy and lactation, and the child's diet during infancy to the development of type 1 diabetes, allergic diseases and asthma in childhood. At the age of 5 years, 3523 children who were still participating in the dietary follow-up (born between 20 October 1997 and 29 February 2004) were invited to take part in the allergy study by completing a questionnaire on the child's history of allergic symptoms. Altogether 3253 (92% of those invited) participated in the allergy study. Of these, complete information on maternal diet during pregnancy was available for 2441 children (75% of those who participated in the allergy study).

Dietary assessment

Maternal food consumption during the last month before the beginning of maternity leave, i.e. the 8th month of pregnancy, was assessed using a validated 181-item FFQ⁽¹⁴⁾. The FFQ specially reflected the diet of Finnish women of childbearing age. Mothers received the FFQ after delivery and recorded the foods they had eaten during the month preceding their pregnancy leave. The use of supplements during the whole pregnancy period was also recorded. The FFQ was returned to the study centre at the child's 3-month visit and was checked by a study nurse. Software

of the National Institute for Health and Welfare, Helsinki, Finland, was used to convert the food consumption data to daily nutrient intake, taking into account women's individual choices of fats used in food preparations and baking, and oils used for salad dressing⁽¹⁵⁾. The dietary exposures investigated in the present study include the main dietary fats (butter, margarine and oils) and a range of FA (SFA, MUFA, PUFA and their subsidiaries). In the validation of the FFQ against a 10 d food records, the energy-adjusted and attenuation-corrected correlation coefficients for these dietary exposures were as follows: total fat (0.48), butter (0.77), margarines (0.64) and oils (0.22), SFA (0.55), MUFA (0.34), PUFA (0.47), *n*-3 PUFA (0.39) and *n*-6 PUFA (0.49). Overall, 70% of the foods and 69% of the nutrients fell into the same or adjacent quintiles⁽¹⁴⁾.

Assessment of allergic outcomes

At the age of 5 years, families of the participating children completed a questionnaire modified from the International Study of Asthma and Allergies in Childhood questionnaire on the child's history of allergic symptoms^(16,17). Allergic rhinitis was defined as sneezing, nasal congestion or rhinitis other than with respiratory infections, accompanied by itching of the eye and tearing during the previous 12 months. Wheezing was defined as parental 'yes' answer on whether the child has had any of the following during the past 12 months: wheezy sound in respiration; wheezy sound in respiration in connection with physical activity; difficulties in respiration in the morning on waking up; wheezy respiration without having the sniffles or respiratory infection; dry cough at nights not associated with common colds or respiratory infections. Age of the onset of rhinitis and wheeze was indicated by the date of the doctor-made diagnosis. Atopic eczema was defined with a positive answer to the question: 'Has your child ever had atopic eczema?'

Sociodemographic and perinatal characteristics

Information on the child's sex, maternal age, maternal educational level and the number of siblings was recorded in a structured questionnaire completed by the parents after the delivery. Information on duration of gestation, birth weight and height, and maternal smoking during pregnancy was received from the Medical Birth Registries of Oulu and Tampere University Hospitals.

Statistical analysis

The distribution of maternal intake of *n*-3 and *n*-6 PUFA in the study population was analysed using the *t* test and ANOVA. Cox proportional hazards regression was used to estimate the risk of the onset of allergic rhinitis and wheeze (because we had the follow-up times for these outcomes), while logistic regression was used to estimate the prevalence of atopic eczema. The FA were energy adjusted by the residual method⁽¹⁸⁾ while total energy intake was included in the models containing the dietary fats. The dietary

variables were categorised in quarters, and the two middle quarters were combined and used as the reference category. *A priori*, we judge that the two middle quarters constitute those with average diet intake in the study population, thus using them as the reference category and comparing the other two extreme quarters seems logical. Selection of confounding covariates was based on their relationships with the exposure and/or the outcome and, in respect of nutrients, on earlier findings. In the adjusted models, the following covariates were included and categorised as shown in Table 1: sex of child, hospital of birth, duration of gestation, maternal age at delivery, maternal basic education, maternal smoking during pregnancy, mode of delivery, number of siblings at

the time of the child's birth, parental asthma, parental allergic rhinitis, pets at home by 1 year of age. A second adjusted model was computed for the FA in which potentially confounding nutrients, vitamin C, Zn, Se, vitamin D and vitamin E were included as additional covariates. Correction for multiple comparisons was performed using the Bonferroni adjustments. All analyses were computed using SPSS version 15.0 (SPSS, Inc., IL, USA).

Results

Participants in the allergy study were generally not systematically different from non-participants regarding the

Table 1. Distribution of maternal intake of *n*-3 and *n*-6 PUFA by the characteristics of the participating children (Mean values with their standard errors, *n* 2441)

Characteristics	Maternal intake of <i>n</i> -3 PUFA			Maternal intake of <i>n</i> -6 PUFA		
	Mean (g/d)	SD	<i>P</i> *	Mean (g/d)	SD	<i>P</i> *
Sex			0.752			0.186
Boys	3.2	1.2		10.7	3.7	
Girls	3.2	1.3		10.9	4.3	
Delivery hospital			0.129			0.013
Oulu (Northern Finland)	3.1	1.3		10.5	3.8	
Tampere (Southern Finland)	3.2	1.7		10.9	4.2	
Duration of gestation			0.634			0.139
Quarter: ≤38-90 weeks	3.1	1.2		10.5	3.7	
Quarter: 39.00-39.90 weeks	3.2	1.2		10.8	3.8	
Quarter: 40.00-40.70 weeks	3.2	1.2		10.9	4.0	
Quarter: >40.70 weeks	3.2	1.3		11.0	4.5	
Missing						
Maternal age			<0.001			0.355
<25 years	3.0	1.2		10.6	4.1	
25-29 years	3.1	1.2		10.7	4.2	
30-34 years	3.2	1.3		10.8	3.9	
≥35 years	3.3	1.3		11.0	3.8	
Maternal basic education			<0.001			0.016
Less than high school	3.0	1.3		10.5	4.2	
High school graduate	3.2	1.3		10.9	3.9	
Maternal smoking during pregnancy			0.947			0.015
No	3.2	1.3		10.7	3.8	
Yes	3.1	1.2		12.5	7.1	
No information	3.2	1.4		11.3	5.2	
Mode of delivery			0.923			0.888
Vaginal delivery	3.2	1.2		10.8	4.0	
Forceps or suction	3.2	1.2		10.8	3.5	
Caesarean section	3.1	1.4		10.9	4.5	
Number of siblings			0.006			0.020
None	3.1	1.2		10.5	3.8	
One	3.3	1.3		11.1	4.1	
Two or more	3.2	1.3		10.9	4.4	
Parental asthma			0.039			0.136
No	3.2	1.3		10.8	4.0	
Yes	3.1	1.3		10.6	4.1	
No information	2.7	1.0		9.5	3.5	
Parental allergic rhinitis			0.194			0.386
No	3.2	1.3		10.8	4.1	
Yes	3.1	1.3		10.8	3.9	
No information	3.4	1.3		11.3	4.8	
Atopic eczema by 6 months of age			0.318			0.656
No	3.2	1.3		10.8	4.1	
Yes	3.2	1.2		10.7	3.9	
Pets at home during the first year			0.587			0.090
No	3.2	1.3		10.7	3.9	
Yes	3.2	1.3		11.0	4.3	

* Based on the *t* test or ANOVA.

background characteristics of the study population either in absolute or relative terms, except for the place of birth: those from Southern Finland were slightly more likely not to take part in the study (10%) than those from Northern Finland (4%; $P < 0.001$). The mean total daily intake of energy for participating mothers was 11 386 (SD 3148) kJ. Protein accounted for 16% (mean intake 110 g; SD 31), fat for 35% (105 g; SD 34) and total carbohydrates for 49% (325 g; SD 95) of the total energy intake. The mean total daily intake of SFA was 45 (SD 17) g, MUFA 37 (SD 12) g, *n*-3 PUFA 3.2 (SD 1.3) g and *n*-6 PUFA 10.8 (SD 4.0) g. None of the women was taking fatty acid supplements during pregnancy.

Significantly higher mean daily intake of *n*-3 PUFA was seen among women over 35 years of age, women with high school degree, children with one sibling and children without parental asthma (Table 1). A higher intake of *n*-6 PUFA was seen among those from Southern Finland, women with high school degree, women who smoked during pregnancy and among children with one sibling. Duration of gestation for more than 40 weeks and having a pet at home during the first year of the child were associated with a decreased risk of atopic eczema, whereas children with parental asthma and rhinitis were at increased risk of atopic eczema. A decreased risk of allergic rhinitis was seen among girls, while children with parental asthma, those with parental rhinitis and those who had atopic eczema by 6 months of age appeared to have an increased risk of rhinitis. Increasing maternal age was associated with decreased risk, whereas increasing number of siblings increased the risk of wheeze (Table 2).

After adjusting for potential confounders, none of the dietary fats or FA was significantly associated with atopic eczema and wheeze. For allergic rhinitis, high maternal consumption of butter and butter spreads was associated with an increased risk of the emergence of allergic rhinitis in the offspring by the age of 5 years (Table 3). A higher ratio of *n*-6:*n*-3 FA during pregnancy was associated with an increased risk of allergic rhinitis (Table 4). On the other hand, a decreased risk of allergic rhinitis was observed with high intakes of total PUFA and α -linolenic FA (Table 4). These results did not change in the models adjusted additionally for the effect of vitamin C, Zn, Se, vitamin D and vitamin E (Table 5). However, after a Bonferroni correction for multiple comparisons, these results lost their statistical significance.

Discussion

The results from the present study show that high maternal consumption of butter and higher ratio of *n*-6:*n*-3 FA during pregnancy were associated with an increased risk of allergic rhinitis in the offspring by the age of 5 years. Conversely, high maternal intake of total PUFA and α -linolenic FA was associated with a decreased risk of rhinitis in the offspring. Adjustment for the effects of vitamin C, Zn, Se, vitamin D and vitamin E did not alter these results, but the results were no longer significant after correction for multiple comparisons.

As a prospective study, the determination of dietary intake before the emergence of the outcomes ensured minimising bias related to the selection of the subjects. The FFQ used in the present study was validated and is suitable for the assessment of dietary intake among childbearing Finnish women⁽¹⁴⁾. The rationale for the dietary assessment was to get a general picture of the diet of the mothers during the first 8 months of pregnancy. We assumed that by assessing the maternal diet during the month preceding the maternity leave (i.e. the 8th month of pregnancy), we obtained a more appropriate overall picture of the maternal diet during pregnancy than by assessing it over the period of the last month before delivery (the 9th month) when most of the mothers did not work outside the home any longer. Dietary measurement error is most consequential if a proportion of subjects are misclassified according to the exposure. There are three recommended validity estimates for nutrients of interest in epidemiological studies: Spearman's correlation coefficients should be above 0.5; more than 50% of subjects should be correctly classified; less than 10% of subjects should be grossly misclassified into thirds⁽¹⁹⁾. In terms of dietary fats, the greatest misclassifications were found for MUFA (9% of the subjects grossly misclassified). Although biomarker assessments represent a precise estimate of the FA, they are limited by being unable to assess the long-term nutrient status. In this case, the FFQ is beneficial, since long-term and habitual intake can be measured. By assessing maternal dietary intake during the 8th month of pregnancy, we aimed at estimating the overall picture of maternal diet during the first 8 months of pregnancy. Although some seasonal variations have been observed in maternal diet during pregnancy as a result of fluctuations in nutrient intakes⁽²⁰⁾, it has been suggested that such changes may be difficult to accurately detect with the rather imprecise dietary assessment methods currently available⁽²¹⁾. Consequently, our generalisation of maternal dietary intake during the 8th month to the whole pregnancy period may remain only a rough approximation. In order to improve the classification of dietary fats, the grouping and brand names of spreads in our FFQ are continuously checked and updated. Most importantly, the overall quintile notation was acceptable for all FA.

There is considerable heterogeneity in respiratory symptoms among children. Consequently, we found associations between FA and rhinitis but not with eczema and wheeze. We lack the knowledge on the mechanisms and the optimal timing of exposure to lower or higher FA levels to influence the development of the different allergic diseases. To exclude younger children having transient respiratory symptoms related to viral infections, the endpoints in the present study were measured in subjects aged 5 years. Because of the lack of data, the child's fat and/or fatty acid intake was not included into our models as a possible confounding factor; thus, we cannot totally rule out that the present results might not have been influenced by dietary intake during infancy. Previous findings have suggested that processes leading to allergic diseases can be initiated very early in the immune development^(22,23). The complex relationship between early nutrition and allergic outcomes emphasises

Table 2. Relationship of characteristics of the participating children (*n* 2441) to the incidence of atopic eczema, allergic rhinitis and wheeze by 5 years of age (Number of participants, percentages, odds ratios, hazard ratios and 95 % confidence intervals)

Characteristics	Frequency (all)		Atopic eczema endpoints		Atopic eczema		Allergic rhinitis endpoints		Allergic rhinitis		Wheeze endpoints		Wheeze	
	<i>n</i>	%	<i>n</i>	%	OR	95 % CI	<i>n</i>	%	Hazard ratio	95 % CI	<i>n</i>	%	Hazard ratio	95 % CI
Sex						<i>P</i> =0.857				<i>P</i> =0.004				<i>P</i> =0.841
Boys	1272	52	480	38	1		212	17	1		358	29	1	
Girls	1169	48	446	38	1.02	0.86, 1.20	147	13	0.73	0.59, 0.91	292	25	0.98	0.80, 1.20
Delivery hospital						<i>P</i> =0.450				<i>P</i> =0.207				<i>P</i> =0.337
Oulu (Northern Finland)	855	35	341	40	1		119	14	1		212	25	1	
Tampere (Southern Finland)	1586	65	585	37	0.88	0.74, 1.05	240	15	1.16	0.92, 1.45	438	28	1.11	0.90, 1.36
Duration of gestation						<i>P</i> =0.004				<i>P</i> =0.713				<i>P</i> =0.591
Quarter: ≤ 38.90 weeks	597	25	235	39	1		79	13	1		171	29	1	
Quarter: 39.00–39.90 weeks	565	23	205	36	0.86	0.68, 1.09	85	15	1.12	0.82, 1.51	153	28	0.85	0.65, 1.11
Quarter: 40.00–40.70 weeks	614	25	265	43	1.16	0.92, 1.46	99	16	1.18	0.88, 1.59	152	25	0.85	0.64, 1.13
Quarter: > 40.70 weeks	646	27	216	33	0.78	0.61, 0.98	95	15	1.06	0.79, 1.44	179	27	0.92	0.70, 1.21
Missing	19	1	5	26			1	5			4	21		
Maternal age						<i>P</i> =0.091				<i>P</i> =0.561				<i>P</i> <0.001
< 25 years	361	15	131	36	1		57	16	1		117	33	1	
25–29 years	871	36	352	40	1.20	0.93, 1.55	127	15	0.94	0.68, 1.31	227	26	0.65	0.49, 0.88
30–34 years	751	31	289	38	1.10	0.84, 1.42	114	15	1.12	0.79, 1.59	179	24	0.68	0.50, 0.93
≥ 35 years	458	19	154	33	0.89	0.67, 1.19	61	13	0.95	0.64, 1.41	127	28	0.44	0.30, 0.63
Maternal basic education						<i>P</i> =0.926				<i>P</i> =0.124				<i>P</i> =0.415
Less than high school	834	34	315	38	1		132	16	1		219	27	1	
High school graduate	1558	64	594	38	1.01	0.85, 1.20	220	14	0.84	0.67, 1.05	414	27	1.09	0.88, 1.35
Missing	49	2	17	35			7	12			17	35		
Maternal smoking during pregnancy						<i>P</i> =0.452				<i>P</i> =0.640				<i>P</i> =0.761
No	2177	89	830	38	1		325	15	1		581	27	1	
Yes	29	1	8	28	0.61	0.27, 1.38	6	21	1.46	0.64, 3.31	8	28	1.39	0.50, 3.89
No information	194	8	77	40	1.06	0.78, 1.43	24	12	0.95	0.63, 1.44	49	26	1.08	0.74, 1.58
Missing	41	2	11	27			4	10			12	29		
Mode of delivery						<i>P</i> =0.336				<i>P</i> =0.245				<i>P</i> =0.234
Vaginal delivery	1959	80	758	39	1		279	14	1		508	26	1	
Forceps or suction	158	6	57	36	0.89	0.63, 1.25	33	21	1.38	0.94, 2.01	46	30	1.13	0.76, 1.68
Caesarean section	309	13	107	35	0.84	0.65, 1.08	47	15	1.00	0.73, 1.39	94	31	1.27	0.96, 1.68
Missing	15	1	4	27			0	0			2	13		
Number of siblings						<i>P</i> =0.427				<i>P</i> =0.863				<i>P</i> =0.001
None	1122	46	427	38	1		177	16	1		314	28	1	
One	750	31	296	39	1.08	0.89, 1.30	101	14	0.93	0.72, 1.21	181	24	1.28	1.01, 1.63
Two or more	546	22	197	36	0.93	0.75, 1.15	79	16	0.95	0.70, 1.28	150	28	1.68	1.27, 2.24
Missing	23	1	6	26			2	9			5	22		
Parental asthma						<i>P</i> <0.001				<i>P</i> =0.038				<i>P</i> =0.998
No	2004	82	716	36	1		269	13	1		486	25	1	
Yes	381	16	185	49	1.70	1.36, 2.12	81	21	1.39	1.08, 1.79	149	40	1.00	0.80, 1.25
No information	33	1	15	45	1.56	0.78, 3.15	5	15	1.28	0.56, 1.28	8	24	1.04	0.29, 3.69
Missing	23	1	10	43			4	17			7	32		
Parental allergic rhinitis						<i>P</i> <0.001				<i>P</i> <0.001				<i>P</i> =0.178
No	797	33	227	28	1		61	8	1		150	19	1	
Yes	1512	62	650	43	1.92	1.59, 2.31	280	19	2.32	1.74, 3.08	462	31	0.90	0.71, 1.13
No information	109	4	43	39	1.68	1.11, 2.54	16	15	1.65	0.92, 2.99	35	33	1.44	0.82, 2.55
Missing	23	1	6	26			2	9			3	14		

Table 2. Continued

Characteristics	Frequency (all)		Atopic eczema endpoints		Allergic rhinitis endpoints		Allergic rhinitis		Wheeze endpoints		Wheeze	
	n	%	n	%	n	%	Hazard ratio	95% CI	n	%	Hazard ratio	95% CI
Atopic eczema by 6 months of age												
No	2007	82	514	26	239	12	1	1	489	25	1	P=0.287
Yes	384	16	384	100	118	31	2.71	2.17, 3.39	147	39	1.13	0.90, 1.41
Missing	50	2	28	56	2	4			14	30		
Pets at home during the first year												
No	1647	67	653	40	252	15	1	1	458	28	1	P=0.182
Yes	783	32	272	35	107	14	0.81	0.68, 0.96	192	25	1.15	0.94, 1.43
Missing	11	1	1	9	0	0			0	0		
Total	2441	100	926	100	359	100			650	100		

the need for consistent and thorough assessment of maternal diets throughout the duration of pregnancy and the child's diet throughout early years. However, duration of exclusive and total breast-feeding has not been related to allergic outcomes in our cohort.

After adjusting for multiple comparisons, the results no longer achieved statistical significance. Therefore, we suggest caution in the interpretation of the present findings. However, we note that the hypotheses linking each of the FA and allergies are mixed, and the precise mechanisms of the role of fatty acids on the occurrence of allergies are not clearly elucidated. Consequently, we believe that each of the associations between the studied FA and the allergic outcomes should be judged on its own merit. Furthermore, the limited and conflicting reports in this area of research signify that reproducibility of these results in further studies would be crucial to rule out any chance finding.

The genetic susceptibility to type 1 diabetes of our study subjects may constrain the generalisability of the present results. Type 1 diabetes is associated with T helper cell 1 (Th1) immune responses, while allergic diseases are related to T helper cell 2 (Th2)-type responses⁽²⁴⁾. In addition to the complexities surrounding the components of the immune system that relate to the generation of Th1 and Th2 responses, it is inconclusive how type 1 diabetes and allergies may be associated⁽²⁵⁾. However, the cumulative incidence of allergies in the present study paralleled that observed in the general Finnish population^(17,26). Furthermore, our subjects were unselected in terms of history of allergy, thus minimising bias towards more selection of children with a positive history for allergic diseases. Participants in the present allergy study were generally not systematically different from non-participating women regarding the background characteristics, suggesting that differential loss to follow-up was minimised in the present study.

The immunological properties of FA have been elaborated^(3,27), particularly that of the *n*-3 and *n*-6 families, which represent the most widely studied FA. While *n*-6 PUFA may possess pro-inflammatory properties, *n*-3 PUFA are thought to act as anti-inflammatory mediators^(27,28). These effects may be prominent during fetal and early life, emphasising the consequences of maternal intake of FA during pregnancy^(29,30). In the present study, while both *n*-6 and *n*-3 PUFA were not appreciably related to the outcomes, total PUFA and α -linolenic FA, however, were associated with a decreased risk of rhinitis. In addition to our present findings, our earlier report has shown an unclear association between maternal intake of PUFA during pregnancy and allergen-specific IgE sensitisation in the offspring in a different subject series of this cohort⁽³¹⁾. In a Japanese study, maternal intake of α -linolenic acid during pregnancy was associated with a reduced risk of wheezing in 2-year-old children⁽⁵⁾. Also, consistent with the present findings, no association was seen between any of the FA and suspected eczema in the Japanese children at the age of 3–4 months⁽³²⁾. The younger age of children in the Japanese cohort makes it likely that their wheezing symptoms may be a result of viral infections, which are usually unrelated to

Table 3. Association between maternal consumption of dietary fats during pregnancy and prevalence of atopic eczema and incidence of allergic rhinitis and wheeze in 5-year-old children (Hazard ratios (HR), odds ratios and 95% confidence intervals)

Food variables	Atopi eczema				Allergic rhinitis				Wheeze				
	Unadjusted†		Adjusted‡		Unadjusted†		Adjusted‡		Unadjusted†		Adjusted‡		
	HR	95 % CI	HR	95 % CI	OR	95 % CI	OR	95 % CI	HR	95 % CI	HR	95 % CI	
Butter and butter spreads													
First quarter (<4.72 g/d)	0.91	0.74, 1.11	0.94	0.76, 1.17	1.10	0.85, 1.43	1.07	0.81, 1.42	1.07	0.84, 1.35	1.09	0.84, 1.41	
Second and third quarters (4.7–19.82 g/d)	1		1		1		1		1		1		
Fourth quarter (19.83–128.38 g/d)	1.09	0.89, 1.34	1.05	0.85, 1.30	1.42*	1.11, 1.81	1.33*	1.03, 1.71	0.84	0.67, 1.06	0.80	0.62, 1.03	
Industrial fats§													
First quarter (<4.84 g/d)	0.96	0.78, 1.17	0.94	0.76, 1.17	0.81	0.62, 1.05	0.76	0.58, 1.00	1.02	0.81, 1.29	0.95	0.74, 1.23	
Second and third quarters (4.85–10.85 g/d)	1		1		1		1		1		1		
Fourth quarter (10.86–115.93 g/d)	0.94	0.75, 1.17	0.95	0.76, 1.20	0.91	0.69, 1.20	0.89	0.67, 1.18	1.21	0.94, 1.56	1.16	0.89, 1.52	
Margarines													
Non-consumers	1		1		1		1		1		1		
Consumers (0.06–120 g/d)	0.92	0.78, 1.09	0.92	0.77, 1.09	0.88	0.72, 1.09	0.89	0.72, 1.11	1.03	0.85, 1.26	1.04	0.84, 1.29	
Oils													
First quarter (<7.25 g/d)	0.98	0.80, 1.20	0.91	0.73, 1.13	0.92	0.71, 1.19	0.86	0.66, 1.12	0.87	0.68, 1.10	0.88	0.68, 1.14	
Second and third quarters (7.25–16.37 g/d)	1		1		1		1		1		1		
Fourth quarter (16.38–145.00 g/d)	0.97	0.80, 1.19	1.01	0.81, 1.26	0.79	0.61, 1.03	0.81	0.61, 1.07	0.98	0.78, 1.24	1.05	0.82, 1.35	

* $P < 0.05$.

† Total energy included in the model.

‡ Adjusted for total energy, sex of child, hospital of birth, duration of gestation, maternal age at delivery, maternal basic education, maternal smoking during pregnancy, mode of delivery, number of siblings at the time of the child's birth, parental asthma, parental allergic rhinitis, pets at home by 1 year of age.

§ Industrial fats: industrial fat mixes and animal fats (grease, suet, lard).

Table 4. Association between maternal fatty acid intake during pregnancy and prevalence of atopic eczema and incidence of allergic rhinitis and wheeze in 5-year-old children (Hazard ratios (HR), odds ratios and 95% confidence intervals)

Fatty acids	Atopic eczema				Allergic rhinitis				Wheeze			
	Unadjusted		Adjusted model I†		Unadjusted		Adjusted model I†		Unadjusted		Adjusted model I†	
	OR	95% CI	OR	95% CI	HR	95% CI	HR	95% CI	HR	95% CI	HR	95% CI
Total fatty acids												
First quarter (<196.85 g/d)	0.89	0.72, 1.09	0.84	0.68, 1.04	0.93	0.71, 1.20	0.92	0.70, 1.20	1.02	0.80, 1.28	1.03	0.81, 1.33
Second and third quarters (196.85–280.83 g/d)	1		1		1		1		1		1	
Fourth quarter (280.84–299.25 g/d)	1.03	0.85, 1.26	1.04	0.84, 1.28	0.96	0.75, 1.24	0.98	0.75, 1.27	0.99	0.79, 1.25	0.95	0.74, 1.21
SFA												
First quarter (<82.24 g/d)	1.01	0.83, 1.24	0.99	0.80, 1.23	1.07	0.83, 1.38	1.06	0.82, 1.39	1.00	0.80, 1.26	0.99	0.79, 1.27
Second and third quarters (82.24–118.56 g/d)	1		1		1		1		1		1	
Fourth quarter (118.57–145.84 g/d)	1.04	0.85, 1.27	1.03	0.84, 1.27	1.10	0.85, 1.41	1.06	0.82, 1.37	0.89	0.70, 1.12	0.80	0.63, 1.03
Trans-fatty acids												
First quarter (<3.10 g/d)	0.95	0.78, 1.16	0.84	0.68, 1.04	1.00	0.77, 1.30	0.95	0.72, 1.25	0.96	0.76, 1.21	1.03	0.80, 1.31
Second and third quarters (3.10–4.18 g/d)	1		1		1		1		1		1	
Fourth quarter (4.19–7.73 g/d)	0.78*	0.62, 0.99	1.08	0.88, 1.33	1.17	0.91, 1.49	1.14	0.89, 1.46	0.85	0.67, 1.07	0.83	0.64, 1.06
MUFA												
First quarter (<78.05 g/d)	0.94	0.77, 1.15	0.90	0.73, 1.11	0.89	0.68, 1.15	0.89	0.68, 1.17	0.94	0.74, 1.18	0.98	0.77, 1.25
Second and third quarters (78.05–93.52 g/d)	1		1		1		1		1		1	
Fourth quarter (93.53–104.16 g/d)	0.98	0.81, 1.20	0.98	0.80, 1.21	0.95	0.74, 1.22	0.96	0.74, 1.24	1.02	0.81, 1.29	1.08	0.85, 1.38
PUFA												
First quarter (<22.90 g/d)	1.02	0.83, 1.25	0.94	0.76, 1.17	0.90	0.70, 1.16	0.82	0.63, 1.07	0.93	0.74, 1.17	0.94	0.73, 1.21
Second and third quarters (22.90–32.46 g/d)	1		1		1		1		1		1	
Fourth quarter (32.47–59.37 g/d)	1.00	0.82, 1.22	1.03	0.83, 1.27	0.75*	0.57, 0.97	0.73*	0.55, 0.96	1.01	0.80, 1.28	1.08	0.84, 1.40
Ratio of n-6 to n-3 PUFA												
First quarter	0.97	0.80, 1.19	0.99	0.80, 1.22	1.08	0.84, 1.40	1.09	0.84, 1.43	0.90	0.71, 1.13	0.85	0.66, 1.10
Second and third quarters	1		1		1		1		1		1	
Fourth quarter	0.94	0.77, 1.15	1.01	0.82, 1.24	1.28*	1.00, 1.64	1.35*	1.05, 1.73	1.14	0.90, 1.43	1.09	0.85, 1.40
n-6 PUFA												
First quarter (<19.32 g/d)	1.03	0.85, 1.26	0.97	0.78, 1.20	0.92	0.71, 1.18	0.85	0.66, 1.11	1.01	0.80, 1.27	1.01	0.79, 1.30
Second and third quarters (19.32–24.50 g/d)	1		1		1		1		1		1	
Fourth quarter (24.51–49.23 g/d)	1.10	0.90, 1.35	1.15	0.93, 1.43	0.86	0.66, 1.12	0.84	0.64, 1.11	1.15	0.92, 1.45	1.21	0.94, 1.55
Linoleic fatty acid												
First quarter (<18.64 g/d)	1.05	0.86, 1.28	0.98	0.79, 1.21	0.93	0.72, 1.19	0.86	0.66, 1.12	1.00	0.80, 1.26	1.01	0.79, 1.30
Second and third quarters (18.64–23.87 g/d)	1		1		1		1		1		1	
Fourth quarter (23.87–47.75 g/d)	1.14	0.93, 1.39	1.17	0.95, 1.45	0.85	0.65, 1.10	0.83	0.63, 1.09	1.14	0.91, 1.44	1.20	0.94, 1.53
Linoleic fatty acid, conjugated												
First quarter (<0.38 g/d)	0.96	0.79, 1.18	0.93	0.75, 1.15	0.98	0.76, 1.27	0.98	0.75, 1.27	1.09	0.87, 1.38	1.19	0.93, 1.52
Second and third quarters (0.38–0.54 g/d)	1		1		1		1		1		1	
Fourth quarter (0.54–0.75 g/d)	1.01	0.83, 1.23	1.01	0.82, 1.24	0.99	0.77, 1.27	0.96	0.74, 1.25	0.93	0.73, 1.18	0.88	0.69, 1.14
γ-Linolenic fatty acid												
First quarter (<0.11 g/d)	0.90	0.73, 1.10	0.91	0.73, 1.12	1.01	0.78, 1.30	1.07	0.82, 1.39	1.14	0.91, 1.44	1.20	0.94, 1.53
Second and third quarters (0.11–0.16 g/d)	1		1		1		1		1		1	
Fourth quarter (0.16–0.98 g/d)	1.05	0.86, 1.28	1.05	0.86, 1.30	1.04	0.81, 1.33	1.08	0.83, 1.39	0.94	0.74, 1.19	0.96	0.74, 1.24
Arachidonic fatty acid												
First quarter (<0.14 g/d)	1.03	0.85, 1.27	1.02	0.82, 1.26	0.82	0.63, 1.07	0.86	0.66, 1.14	0.88	0.69, 1.13	0.95	0.73, 1.23
Second and third quarters (0.14–0.20 g/d)	1		1		1		1		1		1	
Fourth quarter (0.20–0.44 g/d)	1.04	0.86, 1.27	1.02	0.83, 1.25	1.04	0.81, 1.32	1.04	0.81, 1.34	0.85	0.68, 1.06	0.93	0.73, 1.17
n-3 PUFA												
First quarter (<5.55 g/d)	1.08	0.88, 1.32	1.03	0.84, 1.27	0.91	0.71, 1.17	0.89	0.69, 1.15	0.89	0.71, 1.12	0.98	0.77, 1.25
Second and third quarters (5.55–7.34 g/d)	1		1		1		1		1		1	

Fatty acids and allergies in the offspring

Table 4. Continued

Fatty acids	Atopic eczema				Allergic rhinitis				Wheeze			
	Unadjusted		Adjusted model I†		Unadjusted		Adjusted model I†		Unadjusted		Adjusted model I†	
	OR	95% CI	OR	95% CI	HR	95% CI	HR	95% CI	HR	95% CI	HR	95% CI
Fourth quarter (7.35–11.28 g/d)	0.94	0.77, 1.15	0.93	0.75, 1.14	0.78	0.60, 1.02	0.82	0.63, 1.08	0.95	0.75, 1.21	1.11	0.85, 1.43
<i>n</i> -3 PUFA from fish‡												
First quarter (<0.38 g/d)	1.16	0.95, 1.42	1.16	0.93, 1.43	0.99	0.77, 1.28	0.99	0.76, 1.28	1.15	0.92, 1.45	1.08	0.85, 1.38
Second and third quarters (0.38–0.98 g/d)	1		1		1		1		1		1	
Fourth quarter (0.99–5.50 g/d)	0.96	0.79, 1.18	0.97	0.79, 1.20	0.91	0.71, 1.17	0.94	0.72, 1.22	0.95	0.75, 1.20	0.95	0.74, 1.22
<i>n</i> -3 PUFA from plants												
First quarter (<4.79 g/d)	1.02	0.83, 1.24	0.97	0.78, 1.19	1.02	0.80, 1.30	0.97	0.75, 1.25	0.86	0.69, 1.08	0.94	0.74, 1.19
Second and third quarters (4.79–6.27 g/d)	1		1		1		1		1		1	
Fourth quarter (6.28–8.97 g/d)	0.93	0.76, 1.14	0.92	0.74, 1.14	0.80	0.61, 1.04	0.79	0.60, 1.05	0.88	0.69, 1.11	0.94	0.73, 1.21
α -Linolenic fatty acid												
First quarter (<4.45 g/d)	1.08	0.89, 1.32	1.05	0.86, 1.30	1.06	0.83, 1.35	1.01	0.79, 1.29	0.79	0.63, 0.99	0.87	0.69, 1.11
Second and third quarters (4.45–6.40 g/d)	1		1		1		1		1		1	
Fourth quarter (6.40–8.90 g/d)	1.01	0.83, 1.23	0.99	0.80, 1.23	0.74*	0.56, 0.97	0.74*	0.56, 0.99	0.87	0.69, 1.10	0.95	0.74, 1.22
EPA												
First quarter (<0.07 g/d)	1.09	0.89, 1.34	1.07	0.86, 1.33	1.02	0.79, 1.32	0.99	0.76, 1.29	1.05	0.83, 1.32	1.01	0.79, 1.29
Second and third quarters (0.07–0.23 g/d)	1		1		1		1		1		1	
Fourth quarter (0.23–1.25 g/d)	0.96	0.78, 1.17	0.94	0.76, 1.16	1.00	0.78, 1.29	0.99	0.76, 1.28	0.97	0.77, 1.23	0.98	0.76, 1.26
DHA												
First quarter (<0.19 g/d)	1.21	0.99, 1.48	1.18	0.95, 1.46	1.00	0.78, 1.29	0.94	0.72, 1.22	1.10	0.87, 1.38	1.06	0.83, 1.36
Second and third quarters (0.19–0.60 g/d)	1		1		1		1		1		1	
Fourth quarter (0.60–3.5 g/d)	0.98	0.81, 1.20	0.98	0.79, 1.20	0.94	0.73, 1.21	0.93	0.72, 1.21	0.94	0.74, 1.19	0.93	0.73, 1.20

* $P < 0.05$.

† Adjusted for sex of child, hospital of birth, duration of gestation, maternal age at delivery, maternal basic education, maternal smoking during pregnancy, mode of delivery, number of siblings at the time of the child's birth, parental asthma, parental allergic rhinitis, pets at home by 1 year of age.

‡ *n*-3 PUFA from fish is a summary variable including all *n*-3 fatty acids from fish and fish products.

Table 5. Association between maternal fatty acid intake during pregnancy and prevalence of atopic eczema and incidence of allergic rhinitis and wheeze in 5-year-old children

(Hazard ratios (HR), odds ratios and 95% confidence intervals)

	Atopic eczema		Allergic rhinitis		Wheeze	
	Adjusted model II†		Adjusted model II†		Adjusted model II†	
	OR	95% CI	HR	95% CI	OR	95% CI
Total fatty acids						
First quarter (< 196.85 g/d)	0.84	0.67, 1.05	0.94	0.71, 1.24	1.09	0.84, 1.41
Second and third quarters (196.85–280.83 g/d)	1		1		1	
Fourth quarter (280.84–299.25 g/d)	1.03	0.83, 1.28	0.96	0.73, 1.25	0.93	0.72, 1.19
SFA						
First quarter (< 82.24 g/d)	1.00	0.80, 1.24	1.08	0.83, 1.41	1.0	0.78, 1.29
Second and third quarters (82.24–118.56 g/d)	1		1		1	
Fourth quarter (118.57–145.84 g/d)	1.0	0.81, 1.24	1.00	0.76, 1.31	0.77	0.59, 1.00
Trans-fatty acids						
First quarter (< 3.10 g/d)	0.84	0.68, 1.05	0.98	0.74, 1.29	1.08	0.84, 1.40
Second and third quarters (3.10–4.18 g/d)	1		1		1	
Fourth quarter (4.19–7.73 g/d)	1.07	0.87, 1.32	1.09	0.85, 1.41	0.80	0.62, 1.03
MUFA						
First quarter (< 78.05 g/d)	0.90	0.72, 1.12	0.90	0.68, 1.19	1.02	0.79, 1.32
Second and third quarters (78.05–93.52 g/d)	1		1		1	
Fourth quarter (93.53–104.16 g/d)	0.98	0.79, 1.23	0.97	0.73, 1.27	1.08	0.83, 1.40
PUFA						
First quarter (< 22.90 g/d)	0.92	0.73, 1.16	0.81	0.61, 1.07	0.92	0.70, 1.21
Second and third quarters (22.90–32.46 g/d)	1		1		1	
Fourth quarter (32.47–59.37 g/d)	1.03	0.81, 1.31	0.71*	0.52, 0.96	1.09	0.82, 1.44
Ratio of n-6 to n-3 PUFA						
First quarter	1.00	0.80, 1.23	1.10	0.84, 1.44	0.86	0.66, 1.11
Second and third quarters	1		1		1	
Fourth quarter	1.01	0.82, 1.25	1.37*	1.07, 1.77	1.08	0.84, 1.38
n-6 PUFA						
First quarter (< 19.32 g/d)	0.92	0.74, 1.16	0.82	0.62, 1.10	1.0	0.76, 1.32
Second and third quarters (19.32–24.50 g/d)	1		1		1	
Fourth quarter (24.51–49.23 g/d)	1.19	0.94, 1.50	0.83	0.62, 1.12	1.17	0.90, 1.54
Linoleic fatty acid						
First quarter (< 18.64 g/d)	0.94	0.75, 1.18	0.85	0.64, 1.12	1.0	0.76, 1.32
Second and third quarters (18.64–23.87 g/d)	1		1		1	
Fourth quarter (23.87–47.75 g/d)	1.21	0.96, 1.52	0.81	0.60, 1.09	1.15	0.88, 1.51
Linoleic fatty acid, conjugated						
First quarter (< 0.38 g/d)	0.94	0.76, 1.16	1.00	0.77, 1.30	1.24	0.97, 1.59
Second and third quarters (0.38–0.54 g/d)	1		1		1	
Fourth quarter (0.54–0.75 g/d)	0.99	0.79, 1.22	0.91	0.70, 1.19	0.84	0.65, 1.10
γ-Linolenic fatty acid						
First quarter (< 0.11 g/d)	0.91	0.73, 1.12	1.06	0.82, 1.38	1.22	0.95, 1.56
Second and third quarters (0.11–0.16 g/d)	1		1		1	
Fourth quarter (0.16–0.98 g/d)	1.04	0.84, 1.28	1.04	0.80, 1.35	0.97	0.74, 1.27
Arachidonic fatty acid						
First quarter (< 0.14 g/d)	0.98	0.78, 1.23	0.80	0.60, 1.07	0.98	0.74, 1.31
Second and third quarters (0.14–0.20 g/d)	1		1		1	
Fourth quarter (0.20–0.44 g/d)	1.06	0.85, 1.33	1.13	0.86, 1.48	0.87	0.68, 1.11
n-3 PUFA						
First quarter (< 5.55 g/d)	1.03	0.82, 1.28	0.84	0.64, 1.10	0.97	0.75, 1.25
Second and third quarters (5.55–7.34 g/d)	1		1		1	
Fourth quarter (7.35–11.28 g/d)	0.91	0.73, 1.14	0.85	0.64, 1.14	1.14	0.87, 1.49
n-3 PUFA from fish‡						
First quarter (< 0.38 g/d)	1.13	0.90, 1.42	0.91	0.69, 1.20	1.08	0.84, 1.39
Second and third quarters (0.38–0.98 g/d)	1		1		1	
Fourth quarter (0.99–5.50 g/d)	1.00	0.80, 1.26	1.05	0.80, 1.40	0.97	0.73, 1.28
n-3 PUFA from plants						
First quarter (< 4.79 g/d)	0.96	0.77, 1.19	0.95	0.73, 1.24	0.95	0.74, 1.22
Second and third quarters (4.79–6.27 g/d)	1		1		1	
Fourth quarter (6.28–8.97 g/d)	0.90	0.72, 1.12	0.78	0.58, 1.04	0.93	0.72, 1.21
α-Linolenic fatty acid						
First quarter (< 4.45 g/d)	1.05	0.85, 1.30	1.01	0.78, 1.30	0.87	0.67, 1.12
Second and third quarters (4.45–6.40 g/d)	1		1		1	
Fourth quarter (6.40–8.90 g/d)	0.98	0.78, 1.22	0.73*	0.54, 0.98	0.95	0.73, 1.23
EPA						
First quarter (< 0.07 g/d)	1.04	0.83, 1.31	0.90	0.68, 1.19	1.0	0.77, 1.29
Second and third quarters (0.07–0.23 g/d)	1		1		1	
Fourth quarter (0.23–1.25 g/d)	0.97	0.77, 1.22	1.12	0.84, 1.48	1.03	0.78, 1.36

Table 5. Continued

	Atopic eczema		Allergic rhinitis		Wheeze	
	Adjusted model II†		Adjusted model II†		Adjusted model II†	
	OR	95% CI	HR	95% CI	OR	95% CI
DHA						
First quarter (<0.19 g/d)	1.15	0.92, 1.45	0.85	0.64, 1.12	1.04	0.80, 1.34
Second and third quarters (0.19–0.60 g/d)	1		1		1	
Fourth quarter (0.60–3.5 g/d)	1.00	0.80, 1.26	1.05	0.79, 1.39	0.95	0.72, 1.26

* $P < 0.05$.

† In addition to the covariates in model I, adjusted for energy-adjusted maternal intake of vitamin C, Zn, Se, vitamin D and vitamin E.

‡ n -3 PUFA from fish is a summary variable including all n -3 fatty acids from fish and fish products.

actual allergic manifestations. The role of supplementation with n -3 and n -6 PUFA for primary prevention of allergic outcomes has provided inconclusive evidence^(33–37).

Recently, it has been suggested that the ratio of n -6: n -3 FA, rather than the intake of their absolute amount, may represent a more important indicator of optimum health^(38,39). Although previous studies have been inconsistent^(4,39), the results from the present study, however, show that a higher ratio of n -6: n -3 FA was associated with an increased risk of rhinitis in the offspring by the age of 5 years. While this finding is in line with the results of an Australian study reporting an enhanced risk of asthma in children aged 8 years with a higher ratio of n -6: n -3 FA during pregnancy⁽³⁹⁾, it is at odds with that of a recent Japanese study reporting no association⁽³²⁾. In a previous US cohort, cord blood ratios of AA:EPA and linoleic acid: α -linolenic acid were positively associated with eczema and wheeze, respectively, while the ratio of α -linolenic acid: n -3 PUFA was inversely associated with wheeze in 30- to 42-month-old toddlers⁽⁴⁰⁾. In another US study, higher n -3 EPA, n -6 AA and a reduced ratio of AA:EPA were associated with reduced allergen-stimulated interferon- γ levels in neonates⁽⁴¹⁾.

We found an increased risk of rhinitis with high maternal consumption of butter. It has been hypothesised that margarines may increase the risk of allergies, whereas butter may decrease the risk⁽²⁾. The present results of no association between consumption of margarines and the risks of allergies did not support this proposition. In addition, our data on butter are in contrast to the hypothesised role of butter in the development of allergies, and also contradictory to the present observation on the effect of the ratio of n -6: n -3 FA on the risk of rhinitis, considering that n -6 FA are particularly derived from margarines. We lack the explanation for these disparities, but as earlier noted, it may highlight the complex mechanisms relating FA to the development of allergies, especially during early life. In a German cohort, maternal consumption of margarine but not butter during pregnancy was associated with an increased risk of allergies in the offspring⁽⁴²⁾, whereas an Italian retrospective cohort study found no association between either maternal consumption of butter or margarine and atopy in the offspring⁽⁶⁾. Oily fish is a major source of n -3 PUFA, and its consumption

during pregnancy has been related to a reduced risk of asthma⁽⁷⁾, eczema^(9,10), house dust mite, wheeze⁽⁹⁾ and allergic sensitisation^(6,8) in the offspring. In contrast, in an 8-year longitudinal follow-up study from The Netherlands, maternal fish consumption during pregnancy was not associated with asthma or any asthma symptom in the offspring⁽¹¹⁾. We did not have a measure of oily fish in the present study; therefore, we were unable to assess its association with the endpoints.

Conclusion

The present data indicate that maternal consumption of butter, the ratio of n -6: n -3 PUFA, and intake of PUFA and α -linolenic during pregnancy may be potential determinants of the development of allergic rhinitis in the offspring. In the light of discordant reports, more evidence is needed to confirm and further define the putative role of dietary butter and FA in the current increasing prevalence of allergic diseases.

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for 5-year-olds. M. L. participated in the data analysis, interpretation, preparation and review of the manuscript and revising it critically for important intellectual content. S. A. and C. K.-K. were responsible for the data acquisition and preparation, interpretation of the results and critically reviewing the manuscript. R. V., J. I., O. S. and M. K. were responsible for data collection in the clinics and for paediatric expertise in this study, and review of the manuscript and revising it critically for important intellectual content.

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