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Association Between Dietary Patterns Reflecting One-Carbon Metabolism Nutrients Intake Before Pregnancy and Placental DNA Methylation

Short title: OCM nutrients intake and placental DNA methylation

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ABSTRACT

Background: The preconception period represents an important window for fetal and epigenetic programming. Some micronutrients (B vitamins, choline, betaine, methionine) implicated in one-carbon metabolism (OCM) are essential for major epigenetic processes that take place in early pregnancy. However, few studies have evaluated the implication of the micronutrients in placental DNA methylation.

Objective: We investigated whether intake of OCM nutrients in the year before pregnancy was associated with placental DNA methylation in the EDEN mother–child cohort.

Design: Maternal dietary intake was assessed with a food-frequency questionnaire. Three dietary patterns, “varied and balanced diet”, “vegetarian tendency”, and “bread and starchy food”, were used to characterize maternal OCM dietary intake. The Illumina Infinium HumanMethylation450 BeadChip was used to measure placental DNA methylation of 573 women included in the analyses. We evaluated the association of dietary patterns with global DNA methylation. Then, we conducted an agnostic epigenome-wide association study (EWAS) and investigated differentially methylated regions (DMRs) associated with each dietary pattern.

Results: We found no significant association between the three dietary patterns and global DNA methylation or individual CpG sites. DMR analyses highlighted associations between the “varied and balanced” or “vegetarian tendency” pattern and DMRs located at genes previously implicated in functions essential for embryonic development, such as neurodevelopment. The “bread and starchy food” pattern was associated with regions related to genes whose functions involve various metabolic and cell synthesis-related processes.

Conclusions: In mainly well-nourished French women without major deficiencies, OCM intake before pregnancy was not associated with major variation in DNA methylation.

Keywords: epigenetics; birth cohort; methyl-donors; dietary patterns; maternal diet; preconception period; global DNA methylation; placental DNA methylation; epigenome-wide association study.

INTRODUCTION

The preconception period represents an important critical window for fetal and epigenetic programming (1). Epigenetics is defined as the study of the mechanisms that regulate gene expression without altered DNA sequence and appears to be particularly sensitive to environmental exposures during the early development (1,2). Micronutrients implicated in one-carbon metabolism (OCM) deserve special attention because the OCM cycle supports multiple physiological processes essential for human development, including DNA methylation (3). The latter depends upon the supply of dietary OCM nutrients, which are necessary to establish methylation patterns during embryogenesis and maintain them during cell proliferation (1). OCM nutrients are carriers, methyl-group donors (e.g., folates, choline, betaine, methionine), and cofactors of enzymes involved in the transfer reactions of methyl groups to DNA (namely vitamins B2, B6, and B12) (3).

In humans, a few studies have evaluated the association between dietary intake of OCM nutrients and DNA methylation in children. In Gambia, a study evaluated the association between OCM micronutrient periconceptional status, assessed in maternal plasma, and DNA methylation at metastable epialleles, measured from lymphocytes and hair follicles in infants postnatally. Infants conceived during the dry and wet seasons (associated with significant seasonal nutritional variations) had different levels of DNA methylation at a number of sites, which were correlated with micronutrient status (4). Another study reported maternal dietary and supplemental intake of methyl-group donors (folate, betaine, folic acid) in the preconception period associated with buccal epithelial-cell DNA methylation in 6-month-old infants. The identified genes were related to child growth, metabolism and appetite (5). A recent randomized controlled trial showed folic acid supplementation in late pregnancy associated with methylation of a region upstream of the imprinting controller ZFP57 in cord blood (6).

Adequate micronutrient intake is necessary to support placenta homeostasis and development (7). A few studies have investigated the role of OCM nutrients in DNA methylation variation in placental tissue. Two studies evaluated associations between choline intake or folic acid supplementation during pregnancy and placental DNA methylation by using a gene candidate approach but with a small sample size (8,9). Exposure to an inappropriate maternal nutritional environment in the preconception period could also result in placental epigenetic changes, including changes in DNA methylation levels. The early waves of demethylation and de novo methylation that occur after fertilization affect both the epiblast which develops into the embryo, and the trophoblast, which develops into the placenta (10). Nevertheless, it is still unclear to what extent the maternal intake of OCM nutrients might affect placental DNA methylation.

We hypothesized that maternal OCM nutrient intake in the preconception period, more specifically during the year before pregnancy, could affect placental function by changing DNA methylation at an early stage in placental development. The purpose of our study was to examine whether dietary patterns which characterize the variability of pre-pregnancy intake of micronutrients involved in the one-carbon metabolism process (B vitamins, methionine, choline, and betaine) were associated with placenta DNA methylation at delivery in the French EDEN mother-child cohort. As previously described in a recent publication (11), we found that three dietary patterns explained the main variability of OCM nutrient intake before pregnancy, in women from the EDEN study: the “varied and balanced diet,” “vegetarian tendency,” “bread and starchy food” patterns. Our objective was exploratory and to assess how the different dietary patterns reflecting availability of OCM nutrients may influence both global methylation and specific genes. We first assessed the association of dietary pattern with global DNA methylation by using two proxies: *Alu* elements and long interspersed nucleotide elements (LINE-1). Then, we investigated DNA methylation associations with

each dietary pattern by using an epigenome-wide approach at both the single-CpG site and regional levels.

MATERIALS AND METHODS

1. Study

The EDEN mother-child study is a prospective cohort that aimed to evaluate the early, pre- and post-natal determinants of child health and development (12) (see Supporting information, data availability). Study participation was proposed to all women visiting the prenatal clinic before 24 weeks' gestation. Between 2003 and 2006, 2002 (53%) pregnant women aged 18 to 45 years old were recruited in two centres: Nancy and Poitiers hospitals. Exclusion criteria were multiple pregnancies, known diabetes before pregnancy, French illiteracy or planning to move out of the region within the next 3 years. All women gave their informed consent for inclusion before participation in the study. The study received approval from the ethics committee of Kremlin Bicêtre Hospital (02-70) and the Commission Nationale de l'Informatique et des Libertés, the French data privacy institution.

2. Dietary data and other information collected during pregnancy

The diet of women during the year before pregnancy was assessed retrospectively at inclusion (before 28 weeks' gestation; 15 weeks on average) by using a validated food-frequency questionnaire (13). A food composition database was not available to estimate the direct intake of some nutrients of interest (methionine, choline and betaine), so we used different sources to establish a food composition database and calculated individual nutrient intake following a methodology that we previously described (11). Three dietary patterns — “varied and balanced”, “vegetarian tendency”, and “bread and starchy food” — were identified using the reduced rank regression method to characterize OCM micronutrient intake (folates,

choline, betaine, methionine, vitamins B2, B6, B12) in the year before pregnancy (14). The three patterns explained 58% of the variability in OCM nutrient intake. For each woman, a score was calculated for each dietary pattern. The “varied and balanced diet” pattern was characterized by positive coefficients for the main OCM micronutrients, but lower coefficient for betaine. The “vegetarian tendency” pattern was characterized by high coefficients for vitamin B6, B9 and betaine to the detriment of other nutrients such as vitamin B12, choline, and methionine. The “bread and starchy food” pattern was mainly characterized by a high coefficient for betaine and negative coefficients for vitamins B2, B6, and B9 (11). The main characteristics of the patterns are presented in **Table 1**. Information on dietary supplementation in the 3 months before and during pregnancy was also collected by maternal self-completed questionnaires as consumption of mixed multivitamins/minerals or use of single micronutrients, among which we had information for vitamins B2, B6 and B9 (folic acid) as another source of OCM nutrient intake (11). This variable was defined in four categories: no supplementation, supplementation only before pregnancy, supplementation only during pregnancy, and supplementation before and during pregnancy.

Sociodemographic and pregnancy information were recorded by interviews with the mothers at 6 months of pregnancy and after delivery. We selected information about the recruitment centre (Poitiers vs Nancy), maternal age at delivery, BMI at conception (continuous) and maternal education level (at most lower secondary, upper secondary, post-secondary, or tertiary [bachelor’s, master’s, or doctoral level] classified according to the International Standard Classification of Education (15)). Smoking during pregnancy was classified into three groups to identify non-smokers, moderate, and heavy smokers (never, 1–9 cigarettes/day, ≥ 10 cigarettes/day, respectively).

3. Placental DNA methylation assessment

In total, 1301 placental samples were collected at delivery by a midwife or a technician of the study following a standardized protocol. Samples of approximately 5×5 mm were taken from the centre of the placenta on the fetal side and were immediately frozen at -80°C . DNA from placental samples was extracted using the QIAasympphony instrument (Qiagen, Germany). DNA concentration was determined by Nanodrop measurement and fluorescent quantification using Picogreen. No sample was discarded due to low DNA concentration. The DNA quality control and epigenome analysis steps were performed by the Centre National de Recherche en Génomique Humaine (Evry, France). From the 2002 women, 1301 placenta samples were collected. Placenta samples could not be collected from all women, either due to absence of maternal consent, emergency for the EDEN woman or in care unit, and lack of time. Finally, due to budgetary restrictions, in total, 668 children had methylation data available for the study. The DNA samples were plated onto 96- or 48-well plates. A total of nine plates containing 64 chips were used. These plates were processed in four batches. The ratios for child sex (boy/girl) and recruitment centre (Poitiers/Nancy) were balanced for each BeadChip (16). Illumina's Infinium HumanMethylation450 BeadChips were used to evaluate the methylation status of 485,512 individual CpG sites in the placental samples (Illuminas, San Diego, CA, USA) (17). Raw signals from 450K BeadChips were extracted by using GenomeStudio v2011.1 (Illumina) (16). The DNA methylation level of each CpG was calculated as the ratio of the intensity of the normalized fluorescent signals of the methylated alleles to the sum of the methylated and unmethylated alleles.

We performed the data correction, quality control and normalization steps using the *meffil* R package as described by Min et al (18). In the quality control report, we checked variation due to technical artefacts, outliers, poor-quality samples and control probes, the ratio

of unmethylated/methylated signal and extreme dye-bias. We excluded samples showing evidence of extreme dye-bias, probes targeting the sex chromosomes, probes with a bead count <3 in $>10\%$ of samples as well as probes having a detection p-value greater than 0.01 in $>10\%$ of samples, which led to 666 samples and 473 362 CpG sites measured for analyses.

Samples were then normalized by using functional normalization implemented in *meffil*. The main parameter of this method, the number of control probe principal components, was set to 15. This was determined to be the number that maximizes the variance explained in the data (18). Probe intensity quantiles were then adjusted by removing variance encoded in the principal components and batch effects, as fixed and random effects, respectively. Finally, probe intensities were adjusted to conform to the adjusted quantiles. Methylation levels or β -values at CpG sites were estimated as ratios between normalized probe intensities representing methylated signals and the sum of methylated and unmethylated signals. Normalization reports were generated with *meffil*, evaluating associations between potential batch variables with normalized data. This approach identified batches and technical artefacts that were not completely resolved by using functional normalization and for which it was necessary to include the batch variable as covariate in the association studies.

Global methylation was also evaluated by measuring methylation in four CpG sites of repetitive *Alu* and LINE-1 elements by using a previously published pyrosequencing methylation assay (19). More than one-third of DNA methylation is estimated to occur in repetitive elements. Methylation of these repetitive elements can serve as a surrogate marker for global genomic DNA methylation. We then used the median percentage methylation of the four CpG sites for analyses.

4. Estimation of placental cellular composition

Cellular composition of biological samples has been described as an important source of variability in epigenetic epidemiological studies. Placenta is a heterogeneous tissue comprising many cell types of differing development origins (20). DNA methylation levels are known to vary considerably between cell types, trophoblasts and mesenchymal components of the placenta (21). We used a method called reference-free adjustment for cell-type composition (ReFACTor) that is based on principal component analysis (PCA) for correcting cell type heterogeneity (22). This method has been described as improving estimates of the cell-type composition, power and control for false discoveries in epigenome-wide association study (EWAS) (22). One study described the maternal-fetal interface and cellular organization of placenta: seven cell types were identified on the fetal side (23). To be conservative, ReFACTor was used with default parameter settings, including the default number of cell types set to 10. The algorithm is robust to the selection of this parameter set to 10 (22). To avoid ReFACTor generating components encoding sex differences, we included this variable as a covariate in the ReFACTor model.

5. Statistical analyses

5.1 Descriptive analyses

Characteristics of the population with available data on dietary patterns and placental DNA methylation are described with mean $\pm$ SD and number (%).

5.2 Global DNA methylation

We examined the association between the three dietary patterns characterizing the variability in OCM micronutrient intake in the year before pregnancy (“varied and balanced”, “vegetarian tendency” and “bread and starchy food”), OCM intake as supplements and the *Alu* and LINE-1 elements as proxies of overall methylation level. Associations were

investigated by using robust linear regression, accounting for potential confounders and other adjustment factors identified from the literature: child sex, maternal age at delivery (continuous), study centre (Nancy/Poitiers), smoking during pregnancy (never, 1–9 cigarettes/day, ≥ 10 cigarettes/day) and gestational age (continuous). We also adjusted for technical factors related to methylation measurements (batch effect) and estimated cell-type proportions. We also considered vitamin supplementation as an adjustment covariate. The interaction between OCM dietary patterns and maternal supplement intake before pregnancy (yes/no) was tested.

5.3 EWAS

Using the *meffil* package, we ran epigenome-wide analyses to identify CpG associations with each of the three dietary patterns. The dietary patterns are independent factors by construction and thus can be considered separately in the models. We performed analyses in three nested models, namely, a “no covariate model” (no adjustment), a “main model” (adjusted for child sex, study center, maternal age, smoking during pregnancy, gestational age, batch effect and cellular composition), and a “full model” (main model + adjustment for “surrogate variables” (SVA)). Indeed, in this last model, we used surrogate variable analysis to remove other potential sources of heterogeneity (24). Finally, we adjusted these analyses for vitamin supplementation before and during pregnancy. We used Bonferroni correction to derive a conservative p-value threshold of 1.0×10^{-7} ($0.05/473362$ independent tests) to determine sites showing strong evidence of association with our exposure, dietary patterns (25). To reduce the influence of potential outliers, we excluded data points below the 25th percentile minus 3 interquartile ranges or above the 75th percentile plus 3 interquartile ranges for each probe. Analyses with additional adjustment for maternal education and parity were performed. Since these adjustments had no notable effect on associations with DNA

methylation, either globally or at individual CpG sites, these factors were not considered in the full model.

As a supplementary analysis, we also tested the association between global DNA methylation and EWAS approaches with vitamin supplementation before and during pregnancy as an exposure, adjusted for the three dietary patterns in the “full” model. Statistical analyses were performed with R v3.6.2 and a Bonferroni-corrected $p < 0.05$ was considered significant.

5.4 Differentially methylated regions

Finally, we identified differentially methylated regions (DMRs) by using the *dmrff* package in R, which identifies differentially methylated regions by combining EWAS summary statistics (from the “full model”) for nearby CpGs (26). Significant differentially methylated regions were defined by the following criteria: 1) within one differentially methylated region, the distance between two neighboring probes is at most 500 base pairs; 2) the adjusted Bonferroni p-value associated with regions is < 0.05 , and 3) the regions were considered relevant if they included at least two CpG sites with beta values in the same direction and significantly different from 0 (nominal p-value < 0.05). After identification of the DMR regions, the integrative gene database Genecards (27) and the encyclopedia of DNA elements ENCODE (28) were used to describe functions of identified genes or regions.

6. Sample selection

Among the 2002 women included in the EDEN study, 1301 placental samples were collected at birth and 668 methylome analyses were performed. After the previously described steps of quality control, normalisation and exclusion due to missing data in our exposure variable

(dietary patterns), 573 women were included for analyses. The selection of the population is presented in **Figure 1**.

RESULTS

1. Population characteristics

The characteristics of the 573 women selected for analysis are summarized in **Table 2**. Mean maternal age was 29 years (± 5) and about 30% of mothers smoked during pregnancy. Only 8.6% of women received vitamin and/or mineral supplementation both within the 3 months before pregnancy and during pregnancy.

2. Global DNA methylation

The mean (SD) percentage methylation was 16.2 (± 1.0) for *Alu* and 26.1 (± 1.9) for LINE-1. The estimated correlation between *Alu* and LINE-1 sequences was 0.08 ($p=0.04$). Associations between dietary patterns and global DNA methylation are shown in **Table 3**. We found no association between each dietary pattern and overall DNA methylation levels of *Alu* and LINE-1 elements, after adjustment for different confounding factors. We found no significant interactions between vitamin supplementation before pregnancy and dietary pattern scores in association with *Alu* and LINE-1 methylation levels ($p_{\text{interaction}} > 0.05$). Results were similar after adjustment for cell-type heterogeneity (**Table 3**). However, women who had vitamin supplementation before and during pregnancy showed higher methylation level on the *Alu* repetitive element, with a positive regression coefficient of 0.40 ($p=0.005$), which was not confirmed with the LINE-1 elements.

3. EWAS results

In the unadjusted model, the “varied and balanced” pattern was not associated with placental DNA methylation. Adjustment for potential confounding variables previously described (“full model”) and cell-type estimation did not result in significant association between the dietary pattern scores and placental DNA methylation at any single site among the 557 mother–child pairs ($p > 1.0 \times 10^{-7}$), as shown in the Manhattan plot (**Figure 2**). In the adjusted model, including surrogate variables, no significant CpG site was found. We also showed the results by using a quantile-quantile plot, showing the observed distribution of the p-values against the expected p-values under the null hypothesis of no association (**Figure 3**). Results were similar after adjusting for vitamin supplementation (**Supplemental Figure 1**). Similarly, we did not find evidence for an association between the other dietary patterns “vegetarian tendency” or “bread and starchy food” or vitamin supplementation (considered as a binary variable: supplemented or not) and placental DNA methylation (**Supplemental Figure 2**).

4. Identification of differentially methylated regions

The three dietary patterns were related to several DMRs, which are composed of CpG sites that are differentially methylated compared with the surrounding region. All DMRs contained at least two CpGs sites with a p-value < 0.05 . The “varied and balanced” pattern was significantly related to a DMR located on chromosome 3 and containing eight CpG sites (**Table 4**). This region was associated with the *EPM2A Interacting Protein 1* (*EPM2AIP1*) and *MutL Homolog 1* (*MLH1*) genes. The “vegetarian tendency” pattern was associated with two DMRs, located on chromosomes 19 and 11, containing respectively 6 and 5 CpG sites. DMRs were associated with *Zinc Finger Protein 562* (*ZNF562*) and *Fatty Acyl-CoA Reductase 1* (*FAR1*) genes, respectively (**Table 4**). The “bread and starchy” pattern was related to three DMRs, located on chromosomes 10, 8 and 5 and containing 5, 8, and 3 CpG sites, respectively (**Table 4**). The three DMRs most strongly associated with the “bread and

starchy” pattern were located near the *Arsenite Methyltransferase (AS3MT)* gene, *Gasdermin D (GSDMD)* gene and *Solute Carrier Family 25 Member 46 (SLC25A46)* gene respectively. After additional adjustment for vitamin supplementation, DMR analyses showed similar results for the “bread and starchy food” pattern. One additional region was found located near the *Zinc Finger Protein 175 (ZNF175)* gene (**Table 5**). For the “varied and balanced” pattern, we found an association with a DMR located on chromosome 9 and in a gene related to neurologic development, the *Neural Proliferation, Differentiation and Control 1 (NPDC1)* (**Table 5**). We found consistent association between the “vegetarian tendency” pattern and regions located on *FAR1*. Additionally, we observed a tendency for an association with a DMR associated with the *Delta-Like Canonical Notch Ligand 1 (DLL1)* (**Table 5**).

DISCUSSION

To our knowledge, our study is the first epigenome-wide analysis of placental DNA methylation of dietary patterns characterizing OCM nutrient intake in the year before pregnancy for young women of childbearing age. We found no association between the different dietary patterns and global DNA methylation or individually analysed CpG sites in the placental tissue.

We previously showed in the EDEN cohort, that women with disadvantaged social status and possibly nutritional deficiencies were under-represented (11), which may explain in part why we did not find stronger associations of prenatal diet with placental DNA methylation. We evaluated the impact of pre-pregnancy maternal diet, which is relevant for early phenomena such as implantation and placenta development. However, other environmental and maternal factors during pregnancy affecting the placenta development may have masked the effects of earlier factors such as diet before pregnancy. Also, dietary effects on DNA methylation may be tissue-specific, and different results could be found in other

tissues (cord blood, buccal cells) (20). Finally, little is known about the potentially complex association between methyl-group donors obtained by diet and global DNA methylation. One study evaluated the dietary intake of OCM nutrients (vitamin B12, betaine, choline and folate) during the periconception period and second trimester of pregnancy and found no association with LINE-1 methylation in cord blood (29). A randomized trial showed that children born from healthy mothers who received folic acid supplementation in the second and third trimesters of pregnancy had lower global LINE-1 methylation in cord blood than women who received a placebo (30). Another study of 24 pregnant women did not find an association between folic acid supplementation during pregnancy and LINE-1 methylation in cord blood (31).

However, we identified associations with several DMRs located on genes that could be implicated in functions related to development. The DMR associated with the “varied and balanced” pattern was located in *EPM2AIP1* and *MLH1* genes. *EPM2AIP1* is a protein coding gene, involved in regulating glycogen synthesis and potential diseases associated with its function includes Lafora disease, a rare form of neurodegeneration and severe progressive epilepsy in adolescence (32,33). *MLH1* maintains the integrity of the genome by correcting mismatches that arise from DNA replication errors or damage (34). After adjustment for vitamin supplementation, the same pattern was associated with another DMR located in *NPDC1*, involved in neural cell proliferation and differentiation (35). The strengths of associations with DMRs were relatively low with p-values close to the significance threshold, which explains why some DMRs were identified after additional covariate adjustment. The “varied and balanced” pattern is characterized by intake of the main micronutrients, such as choline and vitamin B9 (or folate), that are known to play an essential role in neurological development. A recent study reported a positive association between serum folate concentrations in adults aged above 65 years and increased DNA methylation on *MLH1* (36).

The “vegetarian tendency” pattern was associated with a DMR located in *ZNF562*, a gene that may be involved in transcriptional regulation, but few data have been published to date. Another DMR negatively associated with this pattern was located on *FAR1* that encodes a protein required for lipid metabolism and fatty acid reduction. Mutations in this gene have been associated with neurological conditions such as severe intellectual disability, epilepsy, microcephaly, congenital cataracts, growth retardation and spasticity (37). After taking into account vitamin supplementation, we found another associated DMR located on *DLL1*, which is required for embryonic development, and plays a role in inter-cellular communication and cell fate decision processes in human lymphopoiesis (38). One study reported a novel differentially methylated region in placental samples from children with autism disorder, corresponding probably to a fetal brain enhancer near the *DLL1* locus (39). *DLL1* is also a transmembrane ligand protein of NOTCH receptors, which are involved in many aspects of vascular development and angiogenesis (40). Several studies have identified dysregulated expression of Notch family members in placentas with preeclampsia (40). As previously described, the “vegetarian tendency” pattern was characterized by a high intake of vitamin B9, which plays an important role in brain and placental development (41). One study reported low folate concentrations in early pregnancy associated with increased risk of preeclampsia (42).

The “bread and starchy food” pattern was associated with DMRs located in genes with varied functions, involved in arsenic metabolism (43), regulation of epithelial proliferation and cell differentiation in the skin and gastrointestinal tract, particularly as shown in one animal study focusing on the *GSDMD* gene (44). Finally, the *SLC25A46* gene may play a role in mitochondria, and mutation of this gene results in neurodegenerative diseases (45). However, interpretation of these results must be considered with caution without additional information on the relation between the identified DMRs and related gene expression.

Finally, vitamin supplementation before and during pregnancy was associated with significantly increased global DNA methylation as assessed by the *Alu* repetitive element. Women who initiated supplementation before pregnancy and followed this during pregnancy may have higher intake of OCM nutrients than their peers. High intake could result in increased methylation level, but this association was not confirmed with the LINE-1 elements. The correlation between the two proxies of global DNA methylation was lower than in other publications (0.08 vs > 0.20), which may explain this difference (46).

Strengths and limitations

This study provides new insights into the characteristics of the diet of women of childbearing age before pregnancy in light of its contribution to the combined intake of OCM nutrients involved in DNA methylation. The placenta represents a relevant tissue, as the main organ of nutritional exchange between the mother and fetus during pregnancy. An important strength is that we had information on global DNA methylation (*Alu* and LINE-1 elements), which few studies have evaluated in placenta so far. Despite a lack of power that could explain the lack of an association in our analyses, particularly with individual CpG sites, the sample size is one of the largest in studies of placental DNA methylation (10).

However the placenta is a complex and heterogeneous tissue in its formation, and genes may be differentially expressed depending on the location of the sampled region (20), which may be a source of variability and thus noise for biological analyses. Moreover, even though we used ReFACTor method to account for cellular heterogeneity, we cannot exclude the possibility of residual confounding. One of the perspectives would be to compare results using the recently developed cell estimation references in the placenta (47).

Another limitation of nutritional epidemiology is the lack of precision in measuring diet in all its complexity. Information on food consumption was self-reported by women during

pregnancy, which might be a source of declarative bias. Biomarkers of nutritional intake of OCM nutrients would be necessary to confirm these results. Unfortunately, these markers are not easy to collect during preconception unless specific studies have been set up during this period. One recent study meta-analysed results from two population-based birth cohort studies in Northern Europeans and reported significant associations between maternal plasma folate during pregnancy and DNA methylation in cord blood (48). We did not find any associations with the CpG sites linked to genes identified in this study, however we did not specifically investigate folate intake, and none of the dietary patterns was designed to proxy folate intake. Nutrients involved in the one-carbon metabolism pathway include a range of biological processes and complex interactions that need further investigation. Furthermore, we attempted to account for many potential confounders including technical factors and cell heterogeneity, but we cannot rule out confounding by genetics or other unmeasured environmental factors that may influence DNA methylation and dietary patterns (maternal stress, diabetes or food contaminant) (10). Finally, although we had information on vitamin supplementation, the data remain imprecise, which is a limitation of this study. We did not have information on the duration of supplementation or at which trimester during which women took supplements. Moreover, women may have had supplementation of one or more vitamins, so interpreting the individual effect of vitamin supplementation on DNA methylation is difficult. We did not attempt replication in this project because to our knowledge, there were no other studies with available data on the combined OCM nutrients intake before pregnancy and placental DNA methylation. However, replication in other cohorts with different nutritional intakes would be helpful to confirm these results.

CONCLUSION

In our sample of French women without major nutritional deficiencies, we did not find any major effect of OCM nutrient intake or vitamin supplementation on global DNA methylation or on CpG sites analysed separately. However, our results highlighted an association between the “varied and balanced” or “vegetarian tendency” dietary pattern and DMRs located in genes involved in various functions related to embryonic development, such as neurodevelopment. DMR analyses showed the “bread and starchy food” pattern associated with methylation of genes with various functions including metabolic or cell synthesis-related processes. Replication of these results in other studies is essential to evaluate the role of OCM nutrients in DNA methylation. Other studies of the functional implications of the genes potentially involved are necessary to understand how they may affect child development.

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Table 1: Characteristics of the three dietary patterns resulting from the reduced rank regression method, described in details in (11)

	Varied and balanced	Vegetarian tendency	Bread and starchy food
High intake of main food groups*	Low-fat milk, meat, liver, fish, eggs and egg dishes, cereals, mixed vegetables, chicory, leeks and cabbage, and broccoli	Fruits, chicory, mixed vegetables, cereals, and bread	Bread, sandwiches and rice, pasta and other grains
Low intake of food groups*	Snacks and confectionery and sugar-sweetened beverages	Meat and liver	Low-fat milk, fruits, fruit juices, and sugar-sweetened beverages
Explained variation in nutrients intake ¹	35%	14.4%	8.3%
Association with OCM nutrients ²			
Positive coefficient	B vitamins, choline, and methionine	B6 and B9 vitamins and betaine	Betaine
Negative coefficient	Betaine	Vitamin B12, choline and methionine	Vitamins B2, B6, and B9

*High and low intake of main food groups with factor loading ≥ 0.20 .

¹Explained variation in the overall nutrient variables from the reduced-rank regression

²Association of the nutrient variables with the three dietary patterns.

Table 2: Characteristics of women selected for analysis (N=573)

Sociodemographic data	Missing data n (%)	Population*
Study centre	0 (0)	-
Poitiers		251 (44)
Nancy		322 (56)
Maternal age (years)	0 (0)	29.2 ± 5
Maternal education level	3 (0.5)	-
Lower secondary school		151 (26.5)
Upper secondary school		100 (17.5)
Post-secondary		123 (21.6)
Tertiary		196 (34.4)
Pregnancy		
Gestational age (weeks)	0 (0)	39.3 ± 1.5
Primiparous	1 (0.15)	258 (45)
Smoking during pregnancy	16 (2.8)	-
No		390 (70)
1–9 cigarettes/day		138 (24.8)
At least 10 cigarettes/day		29 (5.2)
Vitamin supplementation	3 (0.5)	
No		339 (59.5)
Before pregnancy		53 (9.3)
During pregnancy		129 (22.6)
Before and during pregnancy		49 (8.6)
BMI at conception, kg/m²	10 (1.7)	22.9 ± 4.2
Child sex	0 (0)	-
Boy		302 (52.7)
Girl		271 (47.3)

*Descriptive data n (%) and mean ± SD. BMI, body mass index

Table 3: Association between dietary patterns or vitamin supplementation and global DNA methylation in different adjusted models

Exposure	<i>Alu</i> (16.2 ± 1.0) *				<i>LINE-1</i> (26.1 ± 1.9) *			
	β^1 (SE)	P-value	β^2 (SE)	P-value	β^1 (SE)	P-value	β^2 (SE)	P-value
Dietary patterns								
Varied and balanced	-0.02 (0.03)	0.60	-0.02(0.03)	0.52	-0.10 (0.07)	0.16	-0.07 (0.07)	0.31
Vegetarian tendency	0.02 (0.04)	0.69	0.03 (0.04)	0.50	-0.13 (0.08)	0.11	-0.10 (0.08)	0.18
Bread and starchy food	0.03 (0.04)	0.52	0.03 (0.04)	0.41	-0.06 (0.09)	0.52	-0.05 (0.09)	0.58
Vitamin supplementation								
No supplementation	Reference		Reference		Reference		Reference	
Before pregnancy only	-0.14 (0.14)	0.29	-0.14 (0.14)	0.29	0.10 (0.29)	0.74	0.06 (0.28)	0.82
During pregnancy only	0.15 (0.09)	0.11	0.16 (0.09)	0.08	0.18 (0.20)	0.37	0.23 (0.20)	0.24
Before and during pregnancy	0.40 (0.14)	0.005	0.37 (0.14)	0.009	-0.07 (0.30)	0.80	-0.10 (0.29)	0.74

Median percentage of the four CpG sites in Alu and long interspersed nucleotide elements [LINE-1] as proxies of overall methylation level

Data are regression coefficients for an increase in one score unit for each dietary pattern.

*mean percentage methylation level

¹ Adjusted models for study center, maternal age, child sex, smoking during pregnancy, gestational age, batch effect and vitamin supplementation (model for dietary pattern), and adjusted mutually for the three dietary patterns (model for vitamin supplementation).

²Additional adjustment for estimated cellular composition.

P<0.05. SE: standard error

Table 4: Dietary patterns in the year before pregnancy and differentially methylated regions (DMRs)¹ of placental DNA in the EDEN cohort

Exposure	Number of CpG sites	Chr	Position	Gene(s)	CpG site	DMR direction of association and individual CpG site β -value ²	P-value ³
Varied and balanced	8	Chr 3	37034730-37034956	<i>EPM2AIP1</i> , <i>MLH1</i>		Positive	0.003
					cg10769891	0.0002466	0.05
					cg19132762	0.0000365	0.81
					cg23658326	0.0001607	0.04
					cg11600697	-0.0002116	0.49
					cg21490561	-0.0001209	0.45
					cg00893636	-0.0000941	0.68
					cg03192963	0.0000985	0.54
					cg06791151	-0.0000509	0.58
Vegetarian tendency	6	Chr 19	9785647-9785919	<i>ZNF562</i>		Positive	4.6 10^{-18}
					cg10755961	0.0005334	0.04
					cg24874111	-0.0006368	0.09
					cg18587364	-0.0003679	0.46
					cg20631204	-0.0002906	0.40
					cg14073063	0.0001544	0.53
					cg14308647	0.0003846	0.03
	5	Chr 11	13690122-13690194	<i>FAR1</i>		Negative	9.5 10^{-10}
					cg02831587	0.0003768	0.04
					cg26240185	-0.0000984	0.17
					cg03326059	-0.0000360	0.49
					cg10572969	-0.0000200	0.77
					cg24412501	0.0003739	0.04

Bread and starchy food	5	Chr 10	104628959-104629151	<i>AS3MT</i>	Negative	0.04
				cg16989281	-0.0040595	0.01
				cg00086670	0.0000005	0.99
				cg25143771	-0.0008883	0.53
				cg08856941	-0.0008247	0.07
				cg24592962	-0.0011958	0.0004
	8	Chr8	144635309-144635547	<i>GSDMD</i>	Negative	0.002
				cg22012530	-0.0021065	0.03
				cg01134012	0.0004952	0.44
				cg18198896	-0.0004702	0.04
				cg14721632	-0.0011329	0.27
				cg23090207	-0.0002147	0.64
				cg12686110	-0.0005133	0.06
				cg15541193	-0.0004292	0.23
				cg09317036	-0.0006335	0.003
	3	Chr 5	110075158-110075285	<i>SLC25A46</i>	Positive	0.0008
				cg13505794	-0.0029363	0.01
				cg13911629	-0.0000855	0.85
				cg27366305	-0.0014410	0.003

¹ DMRs determined with the use of *dmrff* (26). All models were adjusted for child sex, study center, maternal age, gestational age, smoking during pregnancy, batch effect, cell-type distribution estimated and SVA variables.

² Coefficient associated with DMR or direction of the association. β -value reported for individual CpG sites.

³ Adjusted p-value associated with DMR and no-adjusted p-value for CpG sites.

Table 5: Dietary patterns in the year before pregnancy and differentially methylated regions (DMRs)¹ of placental DNA, after adjustment for vitamin supplementation

Exposure	Number of CpG sites	Chr	Position	Gene	CpG site	DMR direction of association and individual CpG site β -value ²	P-value ³
Varied and balanced	3	Chr 9	139940640-139941067	<i>NPDC1</i>		Positive	0.02
					cg22952834	0.0002152	0.031
					cg12611723	0.0014960	0.0001
					cg17902858	0.0010973	0.0004
Vegetarian tendency	4	Chr 6	170599869-170600060	<i>DLL1</i>		Negative	5.3.10 ⁻⁷
					cg08273501	-0.0002168	0.31
					cg12024311	-0.0000268	0.79
					cg27139933	-0.0000134	0.97
					cg00474209	0.0012425	0.004
	3	Chr 6	170599826-170599840	<i>DLL1</i>		Negative	7.0.10 ⁻³
					cg03394401	0.0038978	0.008
					cg14497851	-0.0000644	0.55
					cg03815571	0.0005043	0.40
	2	Chr 11	13690122-13690157	<i>FAR1</i>		Negative	3.7.10 ⁻⁹
					cg02831587	0.0003705	0.04
					g10572969	-0.0001112	0.12
	3	Chr 11	13690160-13690194	<i>FAR1</i>		Negative	7.8.10 ⁻³
					cg03326059	-0.0000295	0.57
					cg10572969	-0.0000240	0.73
					cg24412501	0.0003721	0.05

Bread and starchy food	5	Chr 10	104628959-104629151	<i>AS3MT</i>	cg16989281	Negative	0.02
					cg0086670	-0.0040765	0.01
					cg25143771	0.0000536	0.94
					cg08856941	-0.0007242	0.61
					cg24592962	-0.0008231	0.07
						-0.0012140	0.0003
	8	Chr8	144635309-144635547	<i>GSDMD</i>	cg22012530	Negative	0.004
					cg01134012	-0.0020483	0.03
					cg18198896	0.0005035	0.43
					cg14721632	-0.0004714	0.04
					cg23090207	-0.0010857	0.29
					cg12686110	-0.0002096	0.65
					cg15541193	-0.0005028	0.06
					cg09317036	-0.0004105	0.25
						-0.0006211	0.003
	3	Chr 5	110075158-110075285	<i>SLC25A46</i>	cg13505794	Positive	0.01
					cg13911629	-0.0029720	0.01
					cg27366305	-0.0001567	0.74
						-0.0013743	0.006
	4	Chr 19	52074293-52074348	<i>ZNF175</i>	cg18379413	Positive	0.006
					cg18708075	0.0127613	0.003
					cg26605406	0.0129549	0.02
					cg01382982	0.0095819	0.21
						0.0075372	0.26

¹ DMRs determined with the use of dmrff (26). All models were adjusted for child sex, study center, maternal age, gestational age, smoking during pregnancy, batch effect, cell-type distribution estimated, surrogate variable analysis variables and vitamin supplementation.

² Coefficient associated with DMR or direction of the association. β -value reported for individual CpG sites.

³ Adjusted p-value associated with DMR and no-adjusted p-value for CpG sites.

Figure 1: Flow-chart of population selection

Figure 2: Manhattan plots of epigenome-wide association results for “varied and balanced pattern” within the placenta. The red dashed line represents Bonferroni correction significance threshold ($p = 1.0 \times 10^{-7}$). The direction of the associations is reported with negative effects below zero and positive effects above zero.

Model adjusted for study center, maternal age, child sex, gestational age, smoking during pregnancy, batch effect, cellular composition and surrogate variable analysis variables.

Figure 3: Quantile-quantile plot for regression model between scores on the “varied and balanced diet” in the year before pregnancy and placental DNA methylation.

Model adjusted for study center, maternal age, child sex, gestational age, smoking during pregnancy, batch effect, cellular composition, and SVA variables.

Supporting information

Data availability

The EDEN individual-level data have restricted access owing to ethical and legal conditions in France. They are available upon request from the EDEN steering committee at etude.eden@inserm.fr and through collaborations with the principal investigators of EDEN. Questionnaires are also available upon request at the same address.

Supplemental Figure 1: Quantile-quantile (QQ) plot for regression model between scores on the “varied and balanced diet” pattern in the year before pregnancy and placental DNA methylation, additionally adjusted for vitamin supplementation.

Model adjusted for study center, maternal age, child sex, gestational age, smoking during pregnancy, batch effect, cellular composition, SVA variables and vitamin supplementation.

Vegetarian tendency pattern

Bread and starchy food pattern

Expected $-\log_{10}$ p-value

Vitamin supplementation

Supplemental Figure 2: QQ plots for regression models between the “vegetarian tendency” or “bread and starchy food” patterns* in the year before pregnancy, vitamin supplementation** and placental DNA methylation.

* Model adjusted for study center, maternal age, child sex, gestational age, smoking during pregnancy, batch effect, cellular composition, surrogate variable analysis variables.

** Model adjusted for the previous factors and the three dietary patterns.