



Prenatal folic acid supplementation and folate status in early pregnancy: ECLIPSES study

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Abstract

This research evaluates the prevalence of inadequate folate status in early pregnancy, the pattern of prenatal folic acid (FA) supplementation and associated factors in Spanish pregnant women from the ECLIPSES study, which included 791 participants prior gestational week 12. A cross-sectional evaluation of erythrocyte folate levels was performed at recruitment and used to calculate the prevalence of folate deficiency (erythrocyte folate < 340 nmol/l) and insufficiency (erythrocyte folate < 906 nmol/l). Sociodemographic and lifestyle data as well as information on prenatal FA supplementation were recorded. Descriptive and multivariate statistical analyses were performed. The prevalence of folate deficiency and insufficiency were 9.6% and 86.5%, respectively. Most of women used prenatal FA supplements, but only 6.3% did so as recommended. Supplementation with FA during the periconceptional period abolished folate deficiency and reduced folate insufficiency. Prenatal FA supplementation with ≥ 1000 $\mu\text{g}/\text{d}$ in periconceptional time and pregnancy planning increased erythrocyte folate levels. The main risk factor for folate insufficiency in early pregnancy was getting prenatal FA supplementation out of the periconceptional time (OR 3.32, 95% CI 1.02, 15.36), while for folate deficiency they were young age (OR 2.02, 95% CI 1.05, 3.99), and smoking (OR 2.39, 95% CI 1.30, 4.37). In addition, social and ethnic differences according to folate status were also identified. As conclusion, periconceptional FA use is crucial for achieving optimal folate levels in early pregnancy. Pregnancy planning should focus on young women, smokers, those with low consumption of folate-rich foods, low socio-economic status or from ethnic minorities.

Key words: Folic acid: Folate: Supplementation: Pregnancy planning: Prenatal care: ECLIPSES study

Some folate-dependent *in utero* processes occur very early in gestation, making the periconceptional period especially sensitive to maternal folate status^(1–4). Low prenatal folate concentration poses a public health problem, as it has been widely associated with poor pregnancy outcomes, including megaloblastic anaemia, pre-eclampsia, stillbirth, and pre-term delivery^(5–7). An inadequate maternal folate status may also have a harmful impact on offspring development, including neural tube defects⁽⁸⁾ but also neurodevelopmental disorders such as delayed cognitive abilities, hyperactivity and autism spectrum disorders^(9–14).

One of the major advances to address this concern has been prenatal folic acid (FA) supplementation. The WHO currently recommends 400 $\mu\text{g}/\text{d}$ of FA for childbearing-age women, especially those planning to get pregnant⁽¹⁵⁾. Accordingly, supplementation should begin at periconceptional time, which is from 12 weeks before conception to the first month of

gestation⁽¹⁵⁾. Benefits of prenatal FA supplementation in prevention of congenital disorders are well known^(3,4,8) and, additionally, they have also been observed in cases ranging from neurostructural defects to neurobehavioural and cognitive disorders in the offspring^(1,10,16,17).

Despite growing awareness of the need for correct prenatal folate status, the limited data available indicate that folate deficiency and insufficiency are 0–5%^(18–22) and 40–50%^(19,22), respectively, in developed countries. As for clinical significance and based on WHO recommendations, folate deficiency refers to depleted folate stores, while folate insufficiency indicates a high risk of neural tube defect despite having folate reserves^(19,22–24).

Furthermore, compliance of periconceptional FA supplementation remains low^(25,26), while many women start it during the first months of pregnancy, when damage may already have occurred as a result of an inadequate folate level, if any⁽²⁷⁾. Some sociodemographic and lifestyle characteristics have been

Abbreviations: FA, folic acid; GW, gestational week; SES, socio-economic status.

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identified as possible determinants of the prenatal use of FA and, in turn, of the maternal folate status in early pregnancy. In this sense, young maternal age, low educational level, immigrant status and unplanned pregnancy would be considered the main predictors of reduced use of FA supplements before becoming pregnant, according to previous studies^(26,28–33). Furthermore, smoking, not going to the gynaecologist, and having previous children have also been associated with failure to adhere to recommended prenatal FA supplementation⁽³⁰⁾.

Although the nutritional status of other macro and micronutrients is usually well studied in pregnant women, in Spain there are few data on the prevalence of inadequate folate status in early pregnancy. Moreover, adherence to prenatal FA supplementation by pregnant women in this country is not adequately understood either, despite its great importance for both mother and child. The objective of the present study was to describe the prevalence of folate deficiency and insufficiency at the end of the first trimester of gestation, the pattern of prenatal supplementation with FA, and their associated factors in a sample of healthy Spanish pregnant women.

Methods

Observational study nested in the ECLIPSES community randomised controlled trial, involving 791 pregnant women from Tarragona (Spain) recruited between 2013 and 2017. The aim of the ECLIPSES study was to evaluate the effectiveness of Fe supplements during pregnancy in different doses adjusted to the Hb levels of the first trimester. Participants were contacted in their primary care centres during the first routine visit with midwives before gestational week (GW) 12 and were allocated into three groups of Fe supplementation according to their Hb levels at GW12. Detailed information on ECLIPSES study was shown elsewhere⁽³⁴⁾. The classification of the participants applied in the parent study did not interfere with the current analyses because we evaluated here folate status at GW12, while Fe supplementation started from that moment. Thus, the sample was considered as a whole for the present analyses.

Midwives were responsible for data collection from the clinical history and questionnaires, and for their introduction into an electronic database. This was monitored by an external service to ensure correct data entry and security. The following information of participants at GW12 was extracted from questionnaires and taken into consideration for this work: age, parity, pregnancy planning, BMI, smoking, ethnicity, use of supplements, educational level and occupational status. Educational level and occupational status were used to calculate the familiar socio-economic status (SES).

Dietary assessment at GW12 was performed using a FFQ and then the women's degree of adherence to the Mediterranean diet was obtained, as it is considered a healthy food pattern with a large supply of foods rich in folate. Adherence to Mediterranean diet was calculated using an rMED score, a variation of the original Mediterranean diet score^(35,36) based on the intake of nine components of this diet. Each rMED component (apart from alcohol) was expressed in grams per 1000 kcal/d and was divided by tertiles of dietary intake. Each tertile was

assigned a value of 0, 1 and 2 points. Out of the nine components of the rMED, six scored positively (fruit, vegetables, legumes, cereals, fresh fish and seafood, and olive oil), and two scored negatively (total and processed meat and dairy products). Alcohol was scored as dichotomous variable: 0 when women consumed alcohol and 2 when women did not drink alcohol. The total rMED score ranged from 0 points (minimal adherence) to 18 points (maximum adherence) and pregnant women were classified into three categories accordingly: "low adherence" from 0 to 6 points, "medium adherence" from 7 to 10 points and "high adherence" from 11 to 18 points. Detailed calculation can be found in Jardí *et al.*⁽³⁷⁾.

Regarding supplementation with FA, women were asked whether or not they use FA supplements. If so, information on the daily dose they received and when they started taking it were self-reported. Regarding the dose of FA, women reported 400 or ≥ 1000 $\mu\text{g}/\text{d}$ (including 5000 and 10 000 $\mu\text{g}/\text{d}$), so we classified the dose of FA in these two groups in the subsequent analyses. As for time of initiation, we divided our population into two groups to be able to observe the effect of FA supplementation including or not the periconceptional time: (1) women who took FA from 12 weeks before conception up to 4 weeks gestation and (2) women who started the FA supplementation from 4 weeks of gestation onwards. It has to be clarified that not all women in or out of the periconceptional period took supplements for one entire period or the other but started the FA use in that period. Combining these data, the pattern of FA supplementation was described based on whether women used the recommended dose of FA (400 $\mu\text{g}/\text{d}$) or an excessive amount (≥ 1000 $\mu\text{g}/\text{d}$) in the periconceptional time or after. The total amount of FA taken from the supplements in GW12 according to pattern was calculated by multiplying the number of days taking FA supplements by the daily dose of FA used by each woman.

The use of self-reported FA supplements was validated, considering both the dose and the length of time that the participants had been taking it, by the concentration of erythrocyte folate in GW12. Despite some limitations of erythrocyte folate measurement concerning the risk of haemolysis and other analytical variables, this biomarker reflects tissue stores more closely than serum folate so its concentration is considered the most reliable indicator of folate level^(38–40). In addition, vitamin B₁₂ concentration was also measured since its interplay with folate. Both folate and vitamin B₁₂ concentrations were determined by using a chemiluminescence immunoassay (ADVIA Centaur, Siemens Healthcare Diagnostics Inc.). Then, erythrocyte folate concentration was calculated following this formula: (serum folate in haemolysed whole blood * dilution factor in haemolysis * 100)/haematocrit.

Folate status at GW12 was described as follows: folate deficiency or depleted folate stores when erythrocyte folate < 340 nmol/l; folate insufficiency or at elevated risk of neural tube defect when erythrocyte folate < 906 nmol/l^(23,24,41).

Statistical analyses

Student's *t* test and ANOVA were used to describe continuous variables (mean and SD), while the χ^2 test was used to compare



categorical variables (percentages). Multivariate regression models (multiple linear regressions and logistic regressions) were used to assess the association of possible determinant factors on maternal folate levels and folate status at GW12. Based on previous literature and the results of the bivariate analyses, the regression models were adjusted for the following variables: maternal age (<25 years, 25–34.99 years and ≥ 35 years), maternal initial BMI (underweight, BMI < 18.5 kg/m²; normal weight, BMI 18.5–24.99 kg/m²; overweight, BMI 25–29.99 kg/m²; and obesity, BMI ≥ 30 kg/m²), parity (yes and no), SES (low, middle and high), pregnancy planning (yes and no), maternal ethnicity (Caucasian or ethnic minorities), smoking (yes and no), adherence to Mediterranean diet and serum levels of ferritin and vitamin B₁₂ at GW12.

Additional exploration on whether some sociodemographic characteristics (age, SES and ethnic origin) and health-related behaviours (smoking and adherence to Mediterranean diet) were associated with pregnancy planning was performed.

The analyses showed in this work are secondary analyses from a randomised controlled trial, from which sample size was calculated. Considering a two-sided significance level of 0.05 and a specified sample size of 791 subjects, the study would have 88.5% and 94.5% power to detect differences of 4% in prevalence of folate insufficiency and folate deficiency, respectively.

All statistical analyses were performed using SPSS (version 25.0 for Windows; SPSS Inc.) and statistical significance was set at $P < 0.05$.

Ethical approval

The study was designed in agreement with the Declaration of Helsinki/Tokyo and was approved by Clinical Research Ethics Committee of the Jordi Gol University Institute for Primary Care Research (Institut d'Investigació en Atenció Primària; IDIAP), the Pere Virgili Health Research Institute (Institut d'Investigació Sanitària Pere Virgili; IISPV) and the Spanish Agency for Medicines and Medical Devices (Agencia Española del Medicamento y Productos Sanitarios; AEMPS). Signed, informed consent was obtained from all women participating in the study. The ECLIPSES study was registered at www.clinicaltrialsregister.eu as EudraCT number 2012-005480-28 and at www.clinicaltrials.gov with identification number NCT03196882.

Results

The overall prevalence of folate deficiency (erythrocyte folate <340 nmol/l) and folate insufficiency (erythrocyte folate <906 nmol/l) were 9.6% and 86.5%, respectively (Table 1). This table describes the patterns of prenatal FA supplementation performed by the women in the ECLIPSES study (n 791) and the prevalence according to them; we found that 32.5% of women were not supplemented with FA, while 6.8% of them took more than recommended dose. In addition, almost 60% of study population started the preventive FA supplementation after the first month of gestation (GW4); 54.4% did so using 400 µg/d and 3.8% using more than 1000 µg/d. Consequently, only 6.3% of

the women were optimally compliant with the use of prenatal FA supplements, starting to take 400 µg/d in the recommended time (from 12 weeks before conception). Table 1 also shows that maternal folate level at GW12 was significantly higher in women taking daily 400 µg or ≥ 1000 µg of FA in the periconceptional period (609.69 and 745.74 nmol/l, respectively) than in those using any of the doses after the first month of pregnancy (545.02 and 599.17 nmol/l, respectively). As expected, maternal folate concentration was higher in any of these cases, compared with women who did not receive preventive FA supplements (528.21 nmol/l). Supplementation with high doses of FA (≥ 1000 µg/d) did not lead to exceed the upper value of cut-off point for correct folate status (erythrocyte folate = 1020 nmol/l)⁽⁴²⁾. Regarding the prevalence of folate deficiency (erythrocyte folate <340 nmol/l), it was nil for those women who reported FA supplementation around conception, while it increased to 10.5% and 10% in those using, respectively, daily 400 µg and ≥ 1000 µg out of the recommended period and up to 10.9% in those not supplemented.

Table 2 shows how different sociodemographic characteristics influenced the time of initiation of FA supplementation. Caucasian women, those aged between 25 and 34.99 years, with a medium SES level and pregnancy, tended to initiate FA supplementation in the periconceptional period to a greater extent than their counterparts. Likewise, significantly higher percentages of women under 25 years of age with low SES and ethnicity other than Caucasian were found among those who did not supplement or started the supplementation out of the periconceptional period than among those who met the recommendation. In addition, a significant greater percentage of high adherence to Mediterranean diet was found in women who met the recommendations compared with the others.

After measuring maternal folate levels, we found that Caucasian women, those aged over 35 years, non-smokers, with planned pregnancy, middle or high SES, and a high adherence to Mediterranean diet showed higher folate concentrations than their counterparts (Table 3). Here, coming out the observed association between adherence to the Mediterranean diet and erythrocyte folate levels, we verified the dietary intake according to the degree of adherence. As it is shown in Table 4, the greater the adherence to the Mediterranean diet, the greater the daily consumption of fruits, vegetables, legumes, nuts and fish. These results reinforce the representativeness of the Mediterranean diet as a whole as a good source of folate and its use in this study.

Regarding the prevalence of folate deficiency, some of these factors were statistically significantly associated, the most notable being age, smoking, pregnancy planning and SES. Thus, the results showed that the prevalence of folate deficiency fell from 18.1% in women under 25 years of age to 7.5% in those over 35 years of age. Similar values were obtained for smokers and non-smokers, respectively. The prevalence was 8.4% in women who had planned pregnancy, compared with 14.6% in those who did not, and went from 16.4% in those with a low SES to 5.4% in those with high SES. Maternal characteristics with the most evident association with folate insufficiency were young age, the lack of pregnancy planning and non-Caucasian ethnicity (data not shown).



Table 1. Prevalence of inadequate folate status at GW12, in total sample and according to the pattern of prenatal FA supplementation

		% of total sample (n 791)	Amount of FA from supplements until GW12 (mg)		Erythrocyte folate levels (nmol/l)		Folate status (%)	
			Mean	SD	Mean	SD	Deficiency erythrocyte folate <340 nmol/l (n 76)	Insufficiency erythrocyte folate <906 nmol/l (n 684)
Total sample		100					9.60	86.50
No supplementation		32.50		528.21	182.16		10.90	84.80
In periconceptual time*	400 µg/d	6.30	40.74	10.92	609.69	178.76	0.00	80.05
	≥1000 µg/d	3.00	602.27	184.21	745.74	210.78	0.00	75.00
Out of the periconceptual time*	400 µg/d	54.40	33.60	9.40	545.02	153.04	10.50	87.40
	≥1000 µg/d	3.80	504.00	173.90	599.17	202.10	10.00	83.30

FA, folic acid; GW, gestational week.

* The periconceptual time is from 12 weeks before conception to GW4; out of the periconceptual time is from GW4 onwards.

Table 2. Time of initiation of prenatal folic acid supplementation according to maternal characteristics

	% Total sample (n 791)	Time of initiation of folic acid supplementation			P (A–B)	P (A–C)	P (B–C)
		None (A) (n 257)	In periconceptual time* (B) (n 74)	Out of the periconceptual time* (C) (n 460)			
		%	%	%			
Total sample	100	32.49	9.36	58.15			
Age (years)					0.150	0.480	0.030
<25	14.70	13.60	5.40	16.70			
25–34.99	63.30	63.40	67.60	62.60			
≥35	22.00	23.00	27.00	20.70			
Smoking at recruitment, yes	17.80	19.50	17.00	17.60	0.720	0.400	0.900
Parity, yes	60.10	59.80	59.50	60.40	0.960	0.860	0.870
Pregnancy planning, yes	80.00	76.90	85.10	76.50	0.008	0.950	0.100
Initial BMI					0.470	0.080	0.400
Underweight	1.60	2.70	0.90	2.70			
Normal weight	57.80	53.30	59.30	63.50			
Overweight	26.40	30.40	24.80	23.00			
Obesity	14.20	13.60	15.00	10.80			
Familiar SES					0.010	0.570	0.010
Low	16.20	18.30	4.10	17.00			
Middle	67.00	63.80	74.30	67.60			
High	16.80	17.90	21.60	15.40			
Ethnic origin					0.001	0.950	<0.001
Caucasian	82.60	80.80	97.30	81.00			
Ethnic minorities	17.40	19.20	2.70	19.00			
Adherence to Mediterranean diet					0.003	0.880	0.020
Low-middle	66.80	67.70	54.10	68.30			
High	33.20	32.30	45.90	31.70			

BMI, body mass index; SES, socio-economic status.

* Periconceptual time is from 12 weeks before conception to GW4; out of the periconceptual time is from GW4 onwards.

 Data are expressed in %. P-values for comparisons between groups result from χ^2 test.

Multivariate adjusted analyses (Table 5) showed that, compared with the optimal pattern of FA supplementation which is daily 400 µg in a periconceptual period, both the lack of supplementation ($\beta = -47.55$, 95 % CI -98.51 , 3.41) and the use of daily 400 µg after GW4 ($\beta = -62.47$, 95 % CI -111.68 , -13.25) reduced in a great extent the concentration of maternal folate at GW12. On the contrary, the use of ≥ 1000 µg/d of FA increased maternal folate levels but only when it was taking during the recommended period ($\beta = 137.92$, 95 % CI 57.36 , 218.49). As for the effect on folate status, compared with the use of daily 400 µg of

FA in the periconceptual time, its use after GW4 increased the risk of folate insufficiency (adjusted OR = 3.32, 95 % CI 1.02, 15.36), while the use of ≥ 1000 µg in the recommended period seemed to reduce it by 75 % (adjusted OR = 0.23, 95 % CI 0.05, 1.13). On the contrary, no effect of the prenatal FA supplementation pattern was found on the risk of folate deficiency.

The most notable effects of sociodemographic factors on erythrocyte folate concentrations and folate status were as follows: age under 25 years, compared with ages between 25 and 34.99 years, greatly reduced maternal folate levels

Table 3. Erythrocyte folate in early pregnancy, according to maternal characteristics and pattern of prenatal folic acid supplementation (Mean values and standard deviations)

	% total sample (n 791)	Erythrocyte folate levels (nmol/l)			P
		Mean	SD		
Total sample	100	547.96	173.31		
Age (years)					<0.001
<25	14.70	475.62	160.80		
25–34.99	63.30	557.05	168.05		
≥35	22.00	569.99	184.72		
Smoking at recruitment					0.007
Yes	17.80	512.31	187.74		
No	82.20	555.69	169.19		
Parity					0.980
Yes	60.10	548.08	175.00		
No	39.90	547.77	171.30		
Pregnancy planning					<0.001
Yes	80.00	561.98	175.67		
No	20.00	491.78	151.48		
Initial BMI					0.850
Underweight	1.60	555.33	113.11		
Normal weight	57.80	550.62	182.69		
Overweight	26.40	538.88	166.28		
Obesity	14.20	553.18	152.49		
Familiar SES					<0.001
Low	16.20	501.39	168.98		
Middle	67.00	549.57	174.73		
High	16.80	586.33	162.31		
Ethnic origin					0.040
Caucasian	82.60	554.15	175.90		
Ethnic minorities	17.40	519.49	156.43		
Adherence to Mediterranean diet					0.004
Low-Middle	66.80	535.54	167.64		
High	33.20	572.89	181.95		

BMI, body mass index; SES, socio-economic status.

Table 4. Dietary intake (g/d) of folate-rich foods according to adherence to the Mediterranean diet (Mean values and standard deviations)

	Adherence to Mediterranean diet				P
	Low-middle		High		
	Mean	SD	Mean	SD	
Fruits	149.89	109.22	219.08	129.66	<0.001
Vegetables	65.17	38.69	93.06	50.78	<0.001
Legumes	12.89	10.13	18.32	12.26	<0.001
Nuts	2.52	3.40	3.35	3.96	0.005
Cereals	155.81	67.17	165.32	72.93	0.075
Dairy products	343.20	184.36	255.35	151.16	<0.001
Total meat	118.39	51.76	104.62	38.28	<0.001
Red and processed meat	62.26	34.55	52.76	24.62	<0.001
Fish	36.67	27.79	55.94	30.76	<0.001
Eggs	16.75	12.20	16.87	10.44	0.899

($\beta = -58.56$, 95 % CI -95.13 , -21.98) and doubled the risk of folate deficiency in GW12 (adjusted OR = 2.02, 95 % CI 1.05, 3.99); similarly, smoking reduced folate levels ($\beta = -43.23$, 95 % CI -75.96 , -10.50) and increased the risk of folate deficiency more than twice (adjusted OR = 2.39, 95 % CI 1.30, 4.37). Multivariate analyses also confirmed that maternal folate levels in GW12 were positively associated with pregnancy

planning increased, which protects against folate insufficiency (adjusted OR = 0.12, 95 % CI 0.01, 0.80). Additionally, increasing serum levels of vitamin B₁₂ had a slight but statistically significant protective effect against inadequate folate status (Table 5).

We found that prevalence of smoking and low-middle rather than high adherence to Mediterranean diet were higher when there was no pregnancy planning. The results also showed that pregnancy planning was more common among Caucasian women, those between the ages of 25 and 34.99 years, and middle SES compared with their counterparts (online Supplementary Table 1).

Discussion

Given the scarcity of available data in Spain on the prevalence of inadequate folate status in early gestation as well as on the compliance of prenatal FA supplementation, the present study provides valuable information on these issues and associated factors in a sample of healthy Spanish pregnant women.

The prevalence of folate deficiency and insufficiency found in our study population (9.6 % and 86.5 %, respectively) was much higher than that in neighbouring countries, since available data indicate that they were around 0–5 % and 40–50 %, respectively^(18–22).

Regarding compliance with FA supplementation at any time until the end of the first trimester, the percentage of women using FA in this study was similar to or higher than in other countries in Europe. Thus, participants reporting prenatal FA use represented the 55 % in a large European multicentre study evaluating about 23 000 women⁽⁴⁴⁾, 60.5 % in an Italian study with more than 2000 participants⁽²⁸⁾ and 65.7 % in a Norwegian cohort including 811 healthy pregnant⁽⁴⁵⁾.

Focusing on timing, current global evidence indicates suboptimal use of periconceptional FA supplements, with many countries reporting that fewer than 50 % of women started it before conception⁽²⁶⁾. Recent studies in England⁽⁴⁶⁾ and Norway⁽⁴⁵⁾ found 30–31 % of women taking preconceptionally FA supplements, while lower percentages were found in Italy (23.5 %)⁽²⁸⁾ and Spain (19.2 %)⁽³⁰⁾. We believe that the much lower percentage of periconceptional use of FA found in our study (9.30 %) could be due to the relevant number of participants (~20 %) from ethnic minorities in our population sample; as explained below, it could influence health care awareness^(32,33,47,48). An interesting finding that stands out is the importance of using FA supplements during periconceptional time in relation to the prevalence of folate deficiency. Our observation that the percentage of women with folate deficiency at GW12 was comparable between the group without supplementation and that of women supplemented from GW4 onwards, regardless the dose of FA, allows us to hypothesise that FA supplementation after the conceptional time is useless in improving maternal folate status in early pregnancy.

Based on our findings, and in agreement with many studies, we highlight the central role of pregnancy planning in the optimal adherence to prenatal FA supplementation and the correct folate status^(28–31,49,50). We have found that it takes some time from the start of FA use until good folate stores built up, so pregnancy

Table 5. Association between the pattern of prenatal folic acid supplementation and other characteristics on their erythrocyte folate levels and folate status at GW12†

Erythrocyte folate levels (nmol/l)		
Independent variables	β	95 % CI
Pattern of FA supplementation (none)*	-47.55	-98.51, 3.41
Pattern of FA supplementation (400 μ g/d out of the periconceptual time)*	-62.47	-111.68, -13.25
Pattern of FA supplementation (≥ 1000 μ g/d in periconceptual time)*	137.92	57.36, 218.49
Pattern of FA supplementation (≥ 1000 μ g/d out of periconceptual time)*	15.86	-59.93, 91.64
Maternal age (<25 years v. 25–34.99 years)	-58.56	-95.13, -21.98
Maternal age (≥ 35 years v. 25–34.99 years)	0.05	-29.72, 29.81
Smoking (yes v. no)	-43.23	-75.96, -10.50
Pregnancy planning (yes v. no)	43.68	12.22, 75.15
Serum levels of vitamin B ₁₂ at GW12 (pg/ml)	0.11	0.01, 0.21

Folate insufficiency (erythrocyte folate <906 nmol/l)		
Independent variables	Adjusted OR	95 % CI
Pattern of FA supplementation (none)*	1.37	0.34, 5.47
Pattern of FA supplementation (400 μ g/d out of the periconceptual time)*	3.32	1.02, 15.36
Pattern of FA supplementation (≥ 1000 μ g/d in periconceptual time)*	0.23	0.05, 1.13
Pattern of FA supplementation (≥ 1000 μ g/d out of periconceptual time)*	0.88	0.13, 6.08
Pregnancy planning (yes v. no)	0.12	0.01, 0.80

Folate deficiency (erythrocyte folate <340 nmol/l)		
Independent variables	Adjusted OR	95 % CI
Pattern of FA supplementation (none)*	1.38	0.25, 5.39
Pattern of FA supplementation (400 μ g/d out of the periconceptual time)*	1.32	0.47, 9.58
Pattern of FA supplementation (≥ 1000 μ g/d in periconceptual time)*	0.80	0.24, 2.65
Pattern of FA supplementation (≥ 1000 μ g/d out of periconceptual time)*	0.75	0.32, 4.18
Maternal age (<25 years v. 25–34.99 years)	2.02	1.05, 3.99
Maternal age (≥ 35 years v. 25–34.99 years)	1.06	0.53, 2.11
Pregnancy planning (yes v. no)	0.58	0.32, 1.04
Smoking (yes v. no)	2.39	1.30, 4.37
Serum levels of vitamin B ₁₂ at GW12 (pg/ml)	0.99	0.99, 1.00

GW, gestational week; FA, folic acid.

* Reference category for pattern of FA supplementation: 400 μ g/d in periconceptual time.

† Adjusted for: maternal age, smoking, parity, pregnancy planning, pattern of FA supplementation, hormonal contraception use, maternal initial BMI, socio-economic status, maternal ethnicity, adherence to Mediterranean diet, and serum levels of ferritin and vitamin B₁₂ at GW12.

planning provides that time. We suggest, in addition, that pregnancy planning underlies the observed association between some other sociodemographic factors and the degree of compliance of FA supplementation in early pregnancy. Thus, younger women in our study have been found to be more likely to start FA supplementation after the first month of pregnancy which supports former findings^(26,28,29) and we hypothesise it is probably due to the lack of pregnancy planning in most cases. This makes them more likely to have low prenatal folate concentrations and a greater risk of inadequate folate status^(30,31,50). Similarly and supporting our findings, previous studies have repeatedly identified social and ethnic inequalities in regard to antenatal care and maternal folate status^(32,33). Women of ethnic minorities tend to neglect sexual and reproductive health more than Caucasian women, including prenatal use of FA supplements^(31,45,51–53). Sometimes, they have to deal with low SES and educational level, which can lead to less healthy lifestyle, according to previous knowledge^(31,45). In addition, certain reluctance towards gynaecological and prenatal care has been identified in women from ethnic minorities or born in foreign countries, either for religious, cultural or linguistic reasons^(32,33,47,48). Such situations hinder pregnancy planning and correct compliance with FA supplementation^(32,33,47,48).

Blood folate concentration used to be lower in smokers^(54–56) and a recent systematic review confirmed the detrimental effects of tobacco exposure on folate levels specifically during pregnancy⁽⁵⁷⁾. One of the main postulated mechanisms by which smoking contributes to lowering folate concentrations and to increase the risk of inadequate folate status is the poor eating habits of smokers, who tend to consume less folate-rich foods such as fruits, vegetables and nuts^(54–56,58,59). This suggests that a diet rich in plant-based foods is highly recommended even before pregnancy to achieve optimal folate levels⁽⁶⁰⁾. Thereby, since nuts are not usually consumed so widely, it is especially interesting to highlight its beneficial role in relation to folate status and promote their incorporation into the prenatal diet.

The extensive data collection on sociodemographic characteristics, clinical and obstetrical information, and lifestyle strengthens the findings of this study. Available data on both the dose and the time of initiation of prenatal FA supplementation allowed us to know if women were following international recommendations. As erythrocyte folate reflected tissue stores during the previous 3–4 months, its measurement in GW12 accounts for maternal folate status at the periconceptual time. However, the risk of blood haemolysis and the high sensitivity of erythrocyte folate to some analytical variables could difficult the

measurement procedure. In addition, some limitations have to be considered. Our findings have to be interpreted with caution given that the not-so-large sample size of the present study, despite being common in population-based studies, may limit the generalisability of the results. Furthermore, causal relationships could not be established due to the cross-sectional design.

In conclusion, study findings emphasise the importance of following the recommendation of starting FA supplementation at the periconceptional period to achieve optimal folate levels in early pregnancy. Although pregnancy planning is an accepted guideline, many women still do not adhere to it, so we continue to highlight its crucial role in routine obstetric visits for women who wish to become pregnant in order to strengthen public health strategies aimed to getting good pregnancy outcomes. These strategies should target young women, smokers, poorly adhere to Mediterranean diet which means less consumption of plant-based foods, and those especially vulnerable, such as those with low SES or those belonging to ethnic minorities.

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Supplementary material

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References

1. Naninck EFG, Stijger PC & Brouwer-Brolsma EM (2019) The importance of maternal folate status for brain development and function of offspring. *Adv Nutr* **10**, 502–519.
2. Sato K (2020) Why is folate effective in preventing neural tube closure defects? *Med Hypotheses* **134**, 109429.
3. Imbard A, Benoist JF & Blom HJ (2013) Neural tube defects, folic acid and methylation. *Int J Environ Res Public Health* **10**, 4352–4389.
4. Greenberg JA, Bell SJ, Guan Y, *et al.* (2011) Folic acid supplementation and pregnancy: more than just neural tube defect prevention. *Rev Obstet Gynecol* **4**, 52–59.
5. Molloy AM, Kirke PN, Brody LC, *et al.* (2008) Effects of folate and vitamin B₁₂ deficiencies during pregnancy on fetal, infant, and child development. *Food Nutr Bull* **29**, S101–S111.
6. Talaulikar VS & Arulkumaran S (2011) Folic acid in obstetric practice: a review. *Obstet Gynecol Surv* **66**, 240–247.
7. Green R & Datta Mitra A (2017) Megaloblastic anemias: nutritional and other causes. *Med Clin North Am* **101**, 297–317.
8. Martinez H, Weakland AP, Bailey LB, *et al.* (2018) Improving maternal folate status to prevent infant neural tube defects: working group conclusions and a framework for action. *Ann N Y Acad Sci* **1414**, 5–19.
9. Devilbiss EA, Gardner RM, Newschaffer CJ, *et al.* (2015) Maternal folate status as a risk factor for autism spectrum disorders: a review of existing evidence. *Br J Nutr* **114**, 663–672.
10. Gao Y, Sheng C, Xie R-H, *et al.* (2016) New perspective on impact of folic acid supplementation during pregnancy on neurodevelopment/autism in the offspring children – a systematic review. *PLOS ONE* **11**, e0165626.
11. McGarel C, Pentieva K, Strain JJ, *et al.* (2015) Emerging roles for folate and related B-vitamins in brain health across the lifecycle. *Proc Nutr Soc* **74**, 46–55.
12. Scholtz W, Jones AA, Phillips DI, *et al.* (2012) Lower maternal folate status in early pregnancy is associated with childhood hyperactivity and peer problems in offspring. *J Child Psychol Psychiatry* **100**, 130–134.
13. Zou R, El Marroun H, Cecil C, *et al.* (2021) Maternal folate levels during pregnancy and offspring brain development in late childhood. *Clin Nutr* **40**, 3391–3400.
14. Steenweg-de Graaff J, Roza SJ, Steegers EAP, *et al.* (2012) Maternal folate status in early pregnancy and child emotional and behavioral problems: the generation R study. *Am J Clin Nutr* **95**, 1413–1421.
15. World Health Organization (2016) *WHO Recommendation on Antenatal Care for Positive Pregnancy Experience*. Switzerland: WHO.
16. DeVilbiss EA, Magnusson C, Gardner RM, *et al.* (2017) Antenatal nutritional supplementation and autism spectrum disorders in the Stockholm youth cohort: population based cohort study. *BMJ* **4273**, j4273.
17. Iglesias-Vázquez L, Canals J & Arijá V (2019) Review and meta-analysis found that prenatal folic acid was associated with a 58 % reduction in autism but had no effect on mental and motor development. *Acta Paediatr* **108**, 600–610.
18. O'Malley EG, Cawley S, Kennedy RAK, *et al.* (2018) Maternal anaemia and folate intake in early pregnancy. *J Public Health* **40**, e296–e302.
19. Herter-Aeberli I, Wehrli N, Bärlocher K, *et al.* (2020) Inadequate status and low awareness of folate in Switzerland—a call to strengthen public health measures to ensure sufficient intakes. *Nutrients* **12**, 1–13.
20. Ars CL, Nijis IM, Marroun HE, *et al.* (2019) Prenatal folate, homocysteine and vitamin B₁₂ levels and child brain volumes, cognitive development and psychological functioning: the generation R study. *Br J Nutr* **122**, Suppl. 1, S1–S9.

21. Schulpis K, Spiropoulos A, Gavriili S, *et al.* (2004) Maternal – neonatal folate and vitamin B₁₂ serum concentrations in Greeks and in Albanian immigrants. *J Hum Nutr Diet* **17**, 443–448.
22. Vandevijvere S, Amsalkhir S, Van Oyen H, *et al.* (2012) Determinants of folate status in pregnant women: results from a national cross-sectional survey in Belgium. *Eur J Clin Nutr* **66**, 1172–1177.
23. World Health Organization (2015) Guideline: optimal serum and red cell folate concentrations in women of reproductive age for prevention of neural defects. <https://apps.who.int/iris/handle/10665/161988>
24. de Benoist B (2008) Conclusions of a WHO technical consultation on folate and vitamin B₁₂ deficiencies. *Food Nutr Bull* **29**, S238–S244.
25. McNulty B, Pentieva K, Marshall B, *et al.* (2011) Women's compliance with current folic acid recommendations and achievement of optimal vitamin status for preventing neural tube defects. *Hum Reprod* **26**, 1530–1536.
26. Ray JG, Singh G & Burrows RF (2004) Evidence for suboptimal use of periconceptional folic acid supplements globally. *BJOG* **111**, 399–408.
27. Toriello HV (2005) Folic acid and neural tube defects. *Genet Med* **7**, 283–284.
28. Nilsen RM, Leoncini E, Gastaldi P, *et al.* (2016) Prevalence and determinants of preconception folic acid use: an Italian multicenter survey. *Ital J Pediatr* **42**, 65.
29. Morin P, De Wals P, Noiseux M, *et al.* (2002) Pregnancy planning and folic acid supplement use: results from a survey in Quebec. *Prev Med* **35**, 143–149.
30. Navarrete-Muñoz EM, Monzó DG, De La Hera MG, *et al.* (2010) Folic acid intake from diet and supplements in a population of pregnant women in Valencia, Spain. *Med Clin* **135**, 637–643.
31. Stockley L & Lund V (2008) Use of folic acid supplements, particularly by low-income and young women: a series of systematic reviews to inform public health policy in the UK. *Public Health Nutr* **11**, 807–821.
32. Alderliesten ME, Vrijkotte TGM, Van Der Wal MF, *et al.* (2007) Late start of antenatal care among ethnic minorities in a large cohort of pregnant women. *BJOG* **114**, 1232–1239.
33. Rowe RE, Garcia J & Davidson LL (2004) Social and ethnic inequalities in the offer and uptake of prenatal screening and diagnosis in the UK: a systematic review. *Public Health* **118**, 177–189.
34. Arija V, Fargas F, March G, *et al.* (2014) Adapting iron dose supplementation in pregnancy for greater effectiveness on mother and child health: protocol of the ECLIPSES randomized clinical trial. *BMC Pregnancy Childbirth* **14**, 33.
35. Trichopoulou A, Costacou T, Bamia C, *et al.* (2003) Adherence to a Mediterranean diet and survival in a Greek population. *N Engl J Med* **348**, 2599–2608.
36. Trichopoulou A, Kouris-Blazos A, Wahlqvist ML, *et al.* (1995) Diet and overall survival in elderly people. *BMJ* **311**, 1457.
37. Jardí C, Aparicio E, Bedmar C, *et al.* (2019) Food consumption during pregnancy and post-partum. ECLIPSES study. *Nutrients* **11**, 2447.
38. McNeely M (1987) Folic acid. In *Methods in Clinical Chemistry*, pp. 539–542 [A Pesce and L Kaplan, editors]. St. Louis: CV Mosby.
39. Brewster M (1989) Vitamins. In *Clinical Chemistry: Theory, Analysis, and Correlation*, pp. 543–568 [L Kaplan and A Pesce, editors]. St. Louis: CV Mosby.
40. West AA, Caudill MA, Bailey LB (2020) Folate. In *Present Knowledge in Nutrition*, pp. 239–255 [B Marriott, D Birt, V Stallings, A Yates, editors]. United States: Academic Press.
41. World Health Organization (2015) Serum and red blood cell folate concentrations for assessing folate status in populations. <https://apps.who.int/iris/handle/10665/162114>.
42. LaGow B (2007) *PDR Lab Advisor: A Comprehensive Point-of-Care Guide for Over 600 Lab Tests*. Montvale, NJ, United States: Thomson PDR.
43. Rogers LM, Cordero AM, Pfeiffer CM, *et al.* (2018) Global folate status in women of reproductive age: a systematic review with emphasis on methodological issues. *Ann N Y Acad Sci* **1431**, 35–57.
44. Bitzer J, Von Stenglin A & Bannemerschult R (2013) Women's awareness and periconceptional use of folic acid: data from a large European survey. *Int J Womens Health* **5**, 201–213.
45. Kinnunen TI, Sletner L, Sommer C, *et al.* (2017) Ethnic differences in folic acid supplement use in a population-based cohort of pregnant women in Norway. *BMC Pregnancy Childbirth* **17**, 143.
46. Bestwick JP, Huttly WJ, Morris JK, *et al.* (2014) Prevention of neural tube defects: a cross-sectional study of the uptake of folic acid supplementation in nearly half a million women. *PLOS ONE* **9**, e89354.
47. Van Eijsden M, Van Der Wal MF & Bonsel GJ (2006) Folic acid knowledge and use in a multi-ethnic pregnancy cohort: the role of language proficiency. *BJOG* **113**, 1446–1451.
48. Alomair N, Alageel S, Davies N, *et al.* (2020) Factors influencing sexual and reproductive health of Muslim women: a systematic review. *Reprod Health* **17**, 33.
49. Bixenstine PJ, Cheng TL, Cheng D, *et al.* (2015) Association between preconception counseling and folic acid supplementation before pregnancy and reasons for non-use. *Matern Child Health J* **19**, 1974–1984.
50. Navarrete-Muñoz EM, Valera-Gran D, García de la Hera M, *et al.* (2015) Use of high doses of folic acid supplements in pregnant women in Spain: an INMA cohort study. *BMJ Open* **5**, e009202.
51. Timmermans S, Jaddoe VWV, Mackenbach JP, *et al.* (2008) Determinants of folic acid use in early pregnancy in a multi-ethnic urban population in the Netherlands: the generation R study. *Prev Med* **47**, 427–432.
52. Manniën J, De Jonge A, Cornel MC, *et al.* (2013) Factors associated with not using folic acid supplements preconceptionally. *Public Health Nutr* **17**, 2344–2350.
53. Baraka MA, Steurbaut S, Leemans L, *et al.* (2011) Determinants of folic acid use in a multi-ethnic population of pregnant women: a cross-sectional study. *J Perinat Med* **39**, 685–692.
54. Yila TA, Araki A, Sasaki S, *et al.* (2016) Predictors of folate status among pregnant Japanese women: the Hokkaido study on environment and children's health, 2002–2012. *Br J Nutr* **115**, 2227–2235.
55. Erdemir EO & Bergstrom J (2007) Effect of smoking on folic acid and vitamin B12 after nonsurgical periodontal intervention. *J Clin Periodontol* **34**, 1074–1081.
56. Okumura K & Tsukamoto H (2011) Folate in smokers. *Clin Chim Acta* **412**, 521–526.
57. Tuenter A, Bautista Nino PK, Vitezova A, *et al.* (2018) Folate, vitamin B₁₂, and homocysteine in smoking-exposed pregnant women: a systematic review. *Matern Child Nutr* **2018**, e12675.
58. Segura R, Javierre C, Lizarraga MA, *et al.* (2006) Other relevant components of nuts: phytosterols, folate and minerals. *Br J Nutr* **99**, S36–S44.
59. Ros E (2010) Health benefits of nut consumption. *Nutrients* **2**, 652–682.
60. Koebnick C, Heins UA, Hoffmann I, *et al.* (2001) Folate status during pregnancy in women is improved by long-term high vegetable intake compared with the average Western diet. *J Nutr* **131**, 733–739.