BMJ Open Women's prepregnancy lipid levels and number of children: a Norwegian prospective population-based cohort study

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ABSTRACT

Objective To study prepregnancy serum lipid levels and the association with the number of children.

Design Prospective, population-based cohort.

Setting Linked data from the Cohort of Norway and the Medical Birth Registry of Norway.

Participants 2645 women giving birth to their first child during 1994–2003 (488 one-child mothers and 2157 women with ≥ 2 births) and 1677 nulliparous women.

Main outcome measures ORs for no and one lifetime pregnancy (relative to ≥ 2 pregnancies) obtained by multinomial logistic regression, adjusted for age at examination, education, body mass index (BMI), smoking, time since last meal and oral contraceptive use.

Results Assessed in quintiles, higher prepregnant triglyceride (TG) and TG to high-density lipoprotein (TG:HDL-c) ratio levels were associated with increased risk of one lifetime pregnancy compared with having ≥ 2 children. Compared with the highest quintile, women in the lowest quintile of HDL cholesterol levels had an increased risk of one lifetime pregnancy (OR 1.7, 95% CI 1.2 to 2.4), as were women with the highest low-density lipoprotein (LDL) cholesterol, TG and TG:HDL-c ratio quintiles (compared with the lowest) (OR 1.2, 95% CI 0.8 to 1.7; OR 2.2, 95% CI 1.5 to 3.2; and OR 2.2, 95% CI 1.5 to 3.2, respectively). Similar effects were found in women with BMI ≥ 25 and the highest LDL and total cholesterol levels in risk of lifetime nulliparity.

Conclusion Women with unfavourable prepregnant lipid profile had higher risk of having no or only one child. These findings substantiate an association between prepregnant serum lipid levels and number of children. Previously observed associations between low parity and increased cardiovascular mortality may in part be due to pre-existing cardiovascular disease lipid risk factors.

INTRODUCTION

Cardiovascular disease (CVD) is an important public health problem and remains the number one cause of death in women.¹ Reproductive history is important in evaluating health risks in women, as pregnancy may unmask a woman's predisposition for CVD.¹ Several studies have reported increased CVD mortality among women with no or only

Strengths and limitations of this study

- This is a large population-based study with data collected before pregnancy.
- Linkage with the Medical Birth Registry of Norway provided complete registration of total reproduction.
- Limitations include lack of data on family planning, dietary intake, duration of oral contraceptive use, APOE genotype, low-grade inflammation and thyroid status.
- Non-fasting lipid measurements were used; however, adjustments in our analyses for time since last meal did not change the results.

one lifetime pregnancy.^{2–4} Efforts to elucidate the association between the number of children and the risk of female CVD have been inconclusive.^{1–3} Proposed explanations are lifestyle risk factors associated with child-rearing,⁵ sex hormone fluctuations, protective effect of future pregnancies,³ lifestyle factors prior to conception such as elevated blood pressure and obesity,⁶ as well as metabolic irregularities triggered by gestation.¹ Detection of high-density lipoprotein (HDL) cholesterol and apolipoprotein B (ApoB) in follicular fluid from oocytes^{7,8} suggests a relation between lipids and female reproductive function. More recent studies have reported associations between lipids and fertility in both sexes.⁹ Low parity (as a feature of subfertility) and cardiovascular events may share common pathophysiological mechanisms.¹⁰

While the role of serum lipids in cardiometabolic health is well established, showing low HDL and high triglycerides (TGs) to be strong predictors of CVD,¹¹ their role in reproduction is uncertain. It is also uncertain whether women with no or one lifetime pregnancy have a higher CVD risk to begin with, or whether future pregnancies may reduce the CVD risk.

We pursued this question by exploring the extent to which prepregnant serum lipid levels of total, HDL and low-density lipoprotein (LDL) cholesterol, TG and TG:HDL-c ratio are associated with having no and one lifetime pregnancy.

MATERIALS AND METHODS

Study design and population

We used linked data from the Cohort of Norway (CONOR) and the Medical Birth Registry of Norway (MBRN). CONOR is a population-based collection of health data and blood samples provided by participants older than 20 years of age residing in several different regions in Norway during 1994–2003.¹² Our subset included women with no children at the time of examination with standardised measurements of height, weight and non-fasting lipids levels. Lifestyle factors were obtained through an extensive questionnaire that collected self-reported information on smoking, oral contraceptive (OC) use, self-reported status on receipt of social security benefits, attained level of education and various lifestyle factors.¹² Education in Norway consists of primary school (7 years), lower secondary school (3 years), upper secondary school (3 years) and higher education. The first 10 years are obligatory.

The MBRN has since 1967 recorded data on all deliveries in the country after 16th week of gestation.¹³ Based on mandatory notification, midwives and doctors report information using standard forms throughout pregnancy and at the time of delivery. The registry includes demographic information, mother's health prior and during pregnancy, complications in pregnancy and perinatal outcome. Using the unique national identification number given to all Norwegian citizens, each woman was linked to all her subsequent births (if any) after participating in CONOR. Women reporting no children in CONOR at the time of examination and with no valid records in the MBRN were considered having no pregnancies.

Women with baseline assessment of lifestyle factors in CONOR were linked to the MBRN. We defined one-child mothers as women being 6 years out from their first pregnancy and with no additional births registered in the MBRN.

Preconception measurements

Non-fasting blood samples were analysed on a Hitachi 911 Auto Analyzer (Hitachi, Mito, Japan).¹² Applied reagents were from Boehringer Mannheim (Mannheim, Germany). Serum concentrations of total cholesterol, HDL cholesterol and TG were analysed subsequent to sampling. The total cholesterol, HDL cholesterol levels and TGs were measured by an enzymatic method. The day-to-day coefficients of variation were 2.4% and 0.7%–1.3% for total cholesterol, HDL cholesterol and TG, respectively. To calculate LDL, we used the Friedewald formula¹⁴: total serum cholesterol minus HDL cholesterol

minus one-fifth of the TG concentration. LDL cholesterol levels were calculated only for participants with TG concentrations below 4.5 mmol/L.^{6 14} Accordingly, the TG:HDL-c ratio was expressed as mmol/L.

Trained personnel measured the height and weight, with the participants wearing light clothes and no shoes; measurements were taken as follows: height to the nearest 1.0 cm and weight to the nearest 0.5 kg. Body mass index (BMI) was calculated as weight in kilogram/(height in metres)².

Patients and public involvement

Patients or the public were not directly involved in this study. The detailed explanation of the recruitment process and the obtainment of written informed consent for CONOR were provided elsewhere.¹²

Statistical analyses

The characteristics of the analysed women were presented as means with SD for continuous data and as number with percentages for categorical data. Differences between nulliparous women, one-child mothers and mothers with two or more children, as well as prepregnant health status, were analysed by χ^2 tests and t-tests where appropriate. Linear associations across prepregnant lipid levels (in quintiles) for no and one lifetime pregnancy were assessed by p values for trend. ORs of no and one lifetime pregnancy by lipid levels and TG:HDL-c ratio, when compared with women with two or more pregnancies, were calculated using multinomial logistic regression and adjusted for mother's age at examination, level of education (categorised in <11 years and >11 years of education), smoking (current smoker: yes, no), time since last meal, OC use (now, previously, never) and BMI (linear term). To extend each woman's likelihood of completing her birth record, we separately examined women who were 7 years out from their first pregnancy. About 95% of Norwegian women will complete their second pregnancy within 7 years.⁴ To test the effect of (pregnant) BMI, we stratified the main analyses by BMI (<25 and ≥ 25). To avoid influence from age at first delivery on the number of children, we excluded women older than 34 years at the time of first delivery in a subanalysis. Additionally, we performed sensitivity analyses including only mothers who were 22–30 years old at the time of first delivery. Using presence of a partner (ever) as a proxy for exposure to pregnancy among nulliparous women, we performed logistic regression in a subanalysis (nulliparous vs women with ≥ 2 births) including only women with a reported partner (ever). All analyses were performed using the Statistical Package for Social Sciences (SPSS V.22.0 and V.23.0).

RESULTS

There were 4743 women with baseline assessment of lifestyle factors in CONOR (1994–2003) that were linked to the MBRN. We excluded 421 women with pregnancy

Figure 1 Norwegian women examined in the Cohort of Norway (CONOR) before conception of their first pregnancy and with linked data from the Medical Birth Registry of Norway (MBRN).

at the time of examination ($n=139$), unsure pregnancy status ($n=157$) and missing lipid assessments ($n=125$). Thus, 4322 women were included in the analyses (1677 nulliparous, 488 one-child mothers and 2157 women with ≥ 2 births; see figure 1). Subanalyses included only women with reported partners (228 nulliparous and 216 mothers with ≥ 2 births).

The characteristics of the included women are given in table 1. Nulliparous women were older at the time of examination, had higher BMI and were more frequent smokers compared with women with two or more births. A higher proportion of nulliparous women had >11 years

of education. One-child mothers had higher mean age both at examination and at delivery (29.5 vs 26.7 and 32.3 vs 29.9, respectively), were more often smokers and had lower education than mothers with ≥ 2 births. The mean BMI prior to pregnancy was higher in one-child mothers (24.2 vs 23.5), whereas the mean years from examination to first delivery were similar for the two groups. Women with no and one child were less frequent users of OCs at the time of examination compared with mothers with ≥ 2 births (27.4%, 34.6% and 48.9%, respectively).

The proportion of diabetes at first delivery in one-child mothers was higher than in women with two or more

Table 1 Characteristics of 4322 Norwegian women in the Cohort of Norway, 1994–2003, with no, 1 or ≥ 2 children

Mean values	1677 no child	488 one child	2157 ≥ 2 children
Age (SD) at examination	30.5 (2.1)	29.5 (5.2)	26.7 (4.0)
Age (SD) at first delivery	–	32.3 (4.9)	29.9 (3.8)
Years (SD) from examination to first pregnancy	–	3.7 (2.1)	4.1 (2.3)
BMI (SD) at examination*	24.8 (5.1)	24.2 (4.5)	23.5 (3.4)
OC use*			
Now	455 (27.4)	168 (34.6)	1047 (48.9)
Previously	724 (43.5)	239 (49.2)	779 (36.4)
Never	484 (29.1)	79 (16.3)	317 (14.8)
Smoking at examination*			
Yes	537 (32.2)	182 (37.4)	462 (21.5)
No	1132 (67.8)	304 (62.6)	1685 (78.5)
Education*			
<11 years	312 (18.8)	127 (26.3)	300 (14.1)
≥ 11 years	1344 (81.2)	356 (73.7)	1834 (85.9)

Values are numbers (percentages) unless stated otherwise.

*Missing data on smoking: 8 nulliparous, 2 one-child mothers and 10 women with ≥ 2 children; BMI, 10 nulliparous; education, 21 nulliparous, 5 one-child mothers and 23 women with ≥ 2 children; OC use: 4 nulliparous, 2 one-child mothers and 14 women with ≥ 2 children.

BMI, body mass index; OC, oral contraceptive.

Table 2 ORs with 95% CI for no and one lifetime pregnancy (reference: women with ≥ 2 pregnancies) by prepregnant lipid quintiles in 4322 women in the Cohort of Norway, 1994–2003

Lipid quintiles* in mmol/L	n (%)	n ^a (%)	n ^b (%)	N	Nulliparous OR (95% CI)	One-child mothers OR (95% CI)	P for trend
LDL cholesterol†							0.82
<2.42	449 (56.9)	261 (33.0)	80 (10.1)	790	1.0 (reference)	1.0 (reference)	
2.43–2.84	433 (52.3)	309 (37.4)	85 (10.3)	827	1.1 (0.9 to 1.4)	1.0 (0.7 to 1.4)	
2.85–3.24	454 (52.2)	325 (37.4)	90 (10.4)	869	1.0 (0.8 to 1.3)	0.9 (0.7 to 1.3)	
3.25–3.76	426 (48.0)	353 (39.8)	108 (12.2)	887	1.1 (0.7 to 1.2)	1.1 (0.8 to 1.6)	
>3.77	391 (41.9)	421 (45.1)	121 (13.0)	933	1.2 (0.7 to 1.1)	1.2 (0.8 to 1.7)	
Total cholesterol							0.26
<4.19	432 (55.0)	259 (33.0)	94 (12.0)	785	1.0 (reference)	1.0 (reference)	
4.20–4.61	456 (54.6)	304 (36.4)	75 (9.0)	835	1.2 (0.9 to 1.5)	0.8 (0.5 to 1.1)	
4.62–5.0	434 (52.2)	306 (36.8)	91 (11.0)	831	1.0 (0.8 to 1.3)	0.9 (0.6 to 1.2)	
5.1–5.63	442 (45.7)	415 (43.0)	109 (11.3)	966	1.3 (1.0 to 1.7)	1.0 (0.7 to 1.4)	
>5.64	393 (43.4)	393 (43.4)	119 (13.1)	905	1.2 (0.9 to 1.6)	1.0 (0.7 to 1.4)	
TG							0.01
<0.66	429 (48.6)	372 (42.2)	81 (9.2)	882	1.0 (reference)	1.0 (reference)	
0.67–0.86	447 (49.8)	350 (39.0)	100 (11.1)	897	0.9 (0.7 to 1.1)	1.2 (0.8 to 1.7)	
0.87–1.09	455 (54.4)	294 (35.1)	88 (10.5)	837	0.9 (0.7 to 1.2)	1.3 (0.9 to 1.9)	
1.10–1.45	452 (53.6)	303 (35.9)	88 (10.4)	843	1.0 (0.8 to 1.3)	1.4 (1.0 to 2.0)	
>1.46	373 (43.3)	358 (41.6)	130 (15.1)	861	1.1 (0.9 to 1.5)	2.2 (1.5 to 3.2)	
HDL cholesterol							0.18
<1.20	326 (47.5)	263 (38.3)	97 (14.1)	686	1.0 (0.8 to 1.3)	1.7 (1.2 to 2.4)	
1.21–1.40	271 (44.7)	260 (42.9)	75 (12.4)	606	1.0 (0.8 to 1.2)	1.2 (0.8 to 1.7)	
1.41–1.60	634 (53.0)	431 (36.1)	130 (10.9)	1195	1.0 (0.7 to 1.3)	1.2 (0.9 to 1.6)	
1.61–1.84	443 (49.7)	356 (40.0)	92 (10.3)	891	0.9 (0.7 to 1.2)	1.1 (0.8 to 1.5)	
>1.85	483 (51.2)	367 (38.9)	94 (10.0)	944	1.0 (reference)	1.0 (reference)	

The estimates were obtained by multinomial logistic regression and adjusted for age at examination, educational level, smoking, time since last meal, oral contraceptive use and body mass index (linear term).

Number of women: with ≥ 2 children (n, reference group), nulliparous women (n^a), one-child mothers (n^b) and total number of women within the category (N).

*Quintiles calculated on a total sample prior to pregnancy.

†Missing data within lipids on 16 cases of LDL and 2 cases of TG.

HDL, high-density lipoprotein; LDL, low-density lipoprotein; TG, triglycerides.

births (1.4% vs 0.9%, $p=0.30$). Polycystic ovary syndrome (PCOS) was rare and we only had three cases in our material. A significantly higher number of one-child mothers had in vitro fertilisation (IVF) in their first pregnancy (7.2% vs 2.6% in women with ≥ 2 births; $p<0.001$) (data not shown). This latter finding remained after excluding mothers older than 34 years at first delivery.

ORs with 95% CIs for no and one lifetime pregnancy (vs ≥ 2 lifetime pregnancies) by lipid levels (in quintiles) are presented in table 2 and figure 2. Significant trends in ORs for one lifetime pregnancy across TG and TG:HDL-c ratio quintiles were observed (p trend=0.01). The OR for having one lifetime pregnancy for women with the highest TG quintile compared with the lowest quintile was 2.2 (95% CI 1.5 to 3.2). The ORs for having

one lifetime pregnancy for women with TG:HDL-c ratio levels in the two highest quintiles were 1.7 (95% CI 1.2 to 2.5) and 2.2 (95% CI 1.5 to 3.2), respectively, compared with the lowest quintile. There were no significant trends for LDL cholesterol, total cholesterol or HDL cholesterol, although the ORs of one lifetime pregnancy for the lowest HDL quintile were 1.7 (95% CI 1.2 to 2.4) and for the highest LDL quintile 1.2 (95% CI 0.8 to 1.7). We found no increased risk of being nulliparous by serum lipid levels except for the highest LDL and total cholesterol levels, and these estimates were not persuasive (OR 1.2 (95% CI 0.9 to 1.6) and 1.2 (95% CI 0.9 to 1.5), respectively). Truncation of data to extend the time for each woman to complete her birth record (to 7 years) did not appreciably alter the results, neither did

Figure 2 ORs with 95% CI for no and one lifetime pregnancy (reference: women with ≥ 2 pregnancies) by TG:HDL-c ratio quintiles in 4322 women in the Cohort of Norway, 1994–2003. The estimates were obtained by multinomial logistic regression and adjusted for age at examination, educational level, smoking, time since last meal, oral contraceptive use and body mass index (linear term). TG:HDL-c, triglycerides to high-density lipoprotein ratio.

exclusion of women older than 34 years at the time of first delivery nor the additional restriction of our analyses to mothers aged 22–30 years at first pregnancy. The similar effects of prepregnant lipids as in one-child mothers were observed when subanalyses (nulliparous vs ≥ 2 births) were performed on women who had a partner (as a proxy for ever being exposed to pregnancy). For women with partner, the risk of having no children was increased among the women in the highest quintiles of TG and TG:HDL-c ratio (compared with the lowest quintiles) and also for those in the lowest HDL quintile (compared with the highest) (OR 1.9, 95% CI 0.9 to 4.2; OR 2.0, 95% CI 1.0 to 4.1; and OR 1.6, 95% CI 0.7 to 3.6, respectively).

Stratified analyses by BMI are presented in table 3. In women with BMI ≥ 25 there were significant trends in ORs of having no children or one child across increasing levels of prepregnant total cholesterol, TG and TG:HDL-c ratio quintiles (p trend=0.04 and <0.001 , respectively). The adjusted ORs of one lifetime pregnancy for women with BMI ≥ 25 and TG levels in the two highest quintiles were 2.1 (95% CI 0.9 to 4.8) and 3.5 (95% CI 1.6 to 7.4), and for the two highest TG:HDL-c ratio quintiles 3.1 (95% CI 1.3 to 7.4) and 4.3 (95% CI 1.9 to 10.0) compared with women in the lowest respective quintile. The risk of one lifetime pregnancy was also significantly increased for women with BMI ≥ 25 and the highest LDL and total cholesterol, as well as the lowest HDL quintiles (OR 1.8 (95% CI 0.8 to 3.8), 1.2 (95% CI 0.6 to 2.4) and 2.6 (95% CI 1.3 to 5.3), respectively). Similarly, the ORs of having no pregnancy (in women with BMI ≥ 25) were 1.7 (95% CI 1.0 to 3.0), 2.8 (95% CI 1.7 to 4.7) and 3.6 (95% CI 2.1 to 6.1) for women with the highest LDL, TG and TG:HDL-c ratio quintiles, respectively, compared with women with the lowest quintile. An increased risk of having no children was also found for the overweight and obese women

with the lowest HDL quintile (OR 1.9, 95% CI 1.2 to 3.0). Unlike in one-child mothers, the risk of having no pregnancy among overweight and obese women with higher total cholesterol levels only slightly changed from the main results. In women with prepregnant BMI < 25 , there were significant trends in the risk of having one lifetime pregnancy across increasing levels of prepregnant TG (p trend=0.04), TG:HDL-c ratio (p trend=0.04) and HDL quintiles (p trend=0.05). There were increased risks of one lifetime pregnancy in the highest TG quintile (OR 1.9, 95% CI 1.2 to 3.0) and the two highest TG:HDL-c ratio quintiles (OR 1.6, 95% CI 1.0 to 2.4; and OR 1.8, 95% CI 1.2 to 2.8, respectively), as well as the lowest HDL quintile (OR 1.7, 95% CI 1.1 to 2.6). The risks of no and one lifetime pregnancy with higher LDL and total cholesterol levels only slightly changed compared with our main results.

DISCUSSION

Prepregnant lipid levels were associated with having one lifetime pregnancy. Women with high levels of LDL, TG and TG:HDL-c ratio, as well as low HDL levels, measured years before conception, were at increased risk of having only one lifetime pregnancy. High levels of LDL and total cholesterol were associated with having no children, while in overweight and obese women this was true for all the lipids examined.

These findings provide a possible biological underpinning for a joint mechanistic pathway for reduced fertility and cardiovascular conditions.¹⁰ Our study suggests that the previously observed association between low parity and increased CVD risk may be confounded by pre-existing adverse lipid levels. This does not support the hypothesis that having additional pregnancies reduces

Table 3 OR with 95% CI for no and one lifetime pregnancy (reference: women with ≥ 2 pregnancies) by prepregnant lipid quintiles in 4322 women in the Cohort of Norway (1994–2003), analyses stratified by BMI

Women with prepregnant BMI<25										Women with prepregnant BMI≥25									
Lipid quintiles* in mmol/L	Nulliparous					One-child mothers OR (95% CI)	P for trend	Nulliparous OR (95% CI)					One-child mothers OR (95% CI)	P for trend					
	n	n ^a	n ^b	N	OR (95% CI)			n	n ^a	n ^b	N								
LDL cholesterol†						0.84									0.44				
<2.42	377	219	69	665	1.0 (reference)		72	41	11	124	1.0 (reference)		1.0 (reference)						
2.43–2.84	343	240	64	647	1.1 (0.9 to 1.5)		90	68	20	178	1.1 (0.6 to 2.1)		1.4 (0.6 to 3.3)						
2.85–3.24	341	216	67	624	1.0 (0.7 to 1.3)		113	109	23	245	1.5 (0.8 to 2.6)		1.4 (0.6 to 3.3)						
3.25–3.76	315	198	69	582	1.0 (0.8 to 1.4)		111	152	39	302	1.7 (1.0 to 3.0)		1.8 (0.8 to 4.0)						
>3.77	230	187	64	481	1.2 (0.9 to 1.6)		158	233	57	448	1.7 (1.0 to 3.0)		1.8 (0.8 to 3.8)						
Total cholesterol						0.13									0.04				
<4.19	359	198	75	632	1.0 (reference)		73	60	19	152	1.0 (reference)		1.0 (reference)						
4.20–4.61	341	231	57	629	1.4 (1.1 to 1.9)		115	73	17	205	0.6 (0.3 to 1.0)		0.7 (0.3 to 1.5)						
4.62–5.0	332	191	64	587	1.0 (0.7 to 1.3)		102	114	27	243	1.3 (0.8 to 2.2)		1.2 (0.6 to 2.5)						
5.1–5.63	327	244	72	643	1.3 (0.9 to 1.7)		114	167	37	318	1.5 (0.9 to 2.5)		1.2 (0.6 to 2.4)						
>5.64	248	196	66	510	1.3 (0.9 to 1.7)		143	197	53	393	1.2 (0.7 to 2.0)		1.2 (0.6 to 2.4)						
TG						0.04									<0.001				
<0.66	363	311	69	743	1.0 (reference)		66	59	12	137	1.0 (reference)		1.0 (reference)						
0.67–0.86	364	248	82	694	0.8 (0.6 to 1.0)		83	102	18	203	1.7 (1.0 to 3.0)		1.2 (0.5 to 2.8)						
0.87–1.09	339	208	65	612	0.9 (0.7 to 1.2)		115	85	23	223	1.6 (0.9 to 2.8)		1.7 (0.7 to 3.8)						
1.10–1.45	319	156	58	533	0.8 (0.6 to 1.1)		133	146	30	309	2.5 (1.5 to 4.4)		2.1 (0.9 to 4.8)						
>1.46	222	137	59	418	1.0 (0.7 to 1.4)		149	219	70	438	2.8 (1.7 to 4.7)		3.5 (1.6 to 7.4)						
HDL cholesterol						0.05									0.17				
<1.20	198	70	47	315	0.9 (0.7 to 1.2)		127	191	50	368	1.9 (1.2 to 3.0)		2.6 (1.3 to 5.3)						
1.21–1.40	189	138	43	370	0.9 (0.7 to 1.1)		81	122	31	234	1.7 (1.0 to 2.8)		2.3 (1.1 to 4.9)						
1.41–1.60	475	287	93	855	0.8 (0.6 to 1.1)		158	143	37	338	1.4 (0.9 to 2.2)		1.8 (0.9 to 3.6)						
1.61–1.84	358	272	74	704	0.7 (0.5 to 1.0)		85	82	18	185	1.4 (0.8 to 2.4)		1.8 (0.8 to 4.0)						
>1.85	387	293	77	757	1.0 (reference)		96	73	17	186	1.0 (reference)		1.0 (reference)						
TG:HDL-c ratio						0.04									<0.001				
<0.39	374	321	70	765	1.0 (reference)		67	42	10	119	1.0 (reference)		1.0 (reference)						
0.40–0.54	365	245	75	685	0.8 (0.6 to 1.0)		85	99	17	201	2.2 (1.2 to 4.0)		1.9 (0.7 to 4.9)						
0.55–0.73	365	213	71	649	0.8 (0.6 to 1.1)		101	91	22	214	2.4 (1.3 to 4.3)		2.2 (0.9 to 5.5)						
0.74–1.04	281	172	64	517	0.9 (0.7 to 1.2)		138	131	33	302	3.3 (1.9 to 5.9)		3.1 (1.3 to 7.4)						
>1.05	222	109	53	384	0.8 (0.6 to 1.0)		155	248	71	474	3.6 (2.1 to 6.1)		4.3 (1.9 to 10.0)						

The estimates were obtained by multinomial logistic regression, stratified by BMI and adjusted for age at examination, educational level, smoking, time since last meal and oral contraceptive use.

Number of women: with ≥ 2 children (n, reference group), nulliparous women (n^a), one-child mothers (n^b) and total number of women within the category (N).

*Quintiles calculated on a total sample prior to pregnancy.

†Missing data within lipids on 16 cases of LDL, 2 cases of TG and 2 cases of TG:HDL-c ratio.

BMI, body mass index; LDL, low-density lipoprotein; TG, triglyceride; TG:HDL, TG to high-density lipoprotein ratio.

CVD risk.³ Rather, unfavourable lipid profiles may be related to both subfertility and later CVD.

There is a lack of studies evaluating the relation between preconception lipid levels and fertility in women. The LIFE (Longitudinal Investigation of Fertility and the Environment) study found concentrations of free cholesterol to be associated with fecundity in both sexes.⁹ In contrast to our study, TGs and total cholesterol were not found to be significant in individual and couple-based adjusted models (as well as two other measured lipid components: phospholipids and total lipids); however, the authors used a different study design and lipid measurement methods. In accordance with our findings is the Framingham Heart Study, which detected a trend towards TG elevation and lower HDL serum levels among women with self-reported infertility (as not achieving pregnancy for ≥ 1 year).¹⁵ The presence of HDL cholesterol and ApoB in follicular fluid from human oocytes suggests that these lipids play a direct role in reproduction.^{7 8 16} Previous animal studies have reported an association between dyslipidaemia and infertility.¹⁷ Posed explanations have been that abnormalities in HDL metabolism including change in structure, concentration or function compromise female fertility.^{7 8 16} It has been suggested that genetic polymorphisms that alter function in proteins engaged in cholesterol metabolism may affect human fertility.^{18 19} One of the possible molecular mechanisms could be through a mediating role of HDL on paraoxonase 1 activity. Paraoxonase is an HDL-associated enzyme that inhibits LDL oxidation, and thus protects cells from oxidative stress.²⁰ Its stability and binding affinity are strongly influenced by changes in shape and size of HDL particles.²¹ These changes may lead to decreased antioxidative capacity and consecutively oxidative stress. Oxidative stress is associated with adverse cardiovascular and fertility outcomes, including atherosclerosis, PCOS, pre-eclampsia, endometriosis and infertility.^{19 22} A recent study in women of reproductive age with upper normal ranges of thyroid-stimulating hormone has suggested a direct link between unfavourable lipid profile and increased oxidative membrane damage.²³

Recent insights suggest TG:HDL-c ratio to be a reliable marker of insulin resistance and atherogenicity,²⁴ highlighting its ability to identify insulin resistance in apparently healthy individuals.²⁵ Observed higher levels of TG:HDL-c ratio in our study are indicative of possible pre-existing metabolic risk factors among women with one lifetime pregnancy, as well as a subpopulation of nulliparous women (overweight, obese and with reported partners—as a proxy for exposed to pregnancy). This is also consistent with increasing rates of infertility in both sexes among population with metabolic syndrome.⁹ The higher proportion of diabetes in this group of women further supports this notion. In agreement, the Japan Nurses' Health Study reported a significant increase in the risk of diabetes in young nulliparous women (<45 years of age) with ovarian infertility.²⁶ Accordingly, the Framingham Heart Study found infertile premenopausal women to have increased odds of diabetes and obesity.¹⁵

Given the accompanying metabolic irregularities among major causes of female infertility,^{15 27} substantially higher proportion of IVF treatment among one-child mothers indirectly supports metabolic implications. The latter finding remains after exclusion of women older than 34 years at the time of first delivery.

Dyslipidaemia is associated with PCOS.^{28–30} However, we only identified three women with PCOS in our study sample. Thus, the presence of subclinical forms or under-reporting may be present.

In accordance with the literature,^{27 31} the risk of having no and only one child showed strong effects in overweight and obese women (BMI ≥ 25) in stratified analyses (table 3). Nevertheless, the higher risk of having only one child remained in normal-weight women (BMI < 25) with the lowest HDL quintile and the highest TG and TG:HDL-c ratio quintiles. These findings mirror observations from the literature of metabolic irregularities among normal-weight women as an independent risk factor for future fertility impairments.^{32 33} The LIFE study reported both female and male lipid concentrations to affect fecundity, irrespective of their BMI.⁹

Compared with women with two or more pregnancies, total cholesterol levels above clinically recommended range were associated with risk of having no children, irrespective of BMI. The LIFE study reported a higher percentage of women with a history of irregular menstrual cycles in the highest quartile of free cholesterol.⁹ The Japan Nurses' Health Study found women with ovarian infertility to be at high risk of hypercholesterolaemia.²⁶ In our study, total cholesterol levels were not associated with the risk of having one lifetime pregnancy, except among overweight and obese women. This could suggest that total cholesterol levels play varied roles in different subfecundity or infertility subtypes. In addition, nulliparous women in our study were older at examination and had higher BMI. Both age and obesity are associated with systemic oxidative stress.^{19 22} It is possible that in such physiological environment, clinically abnormal levels of certain lipids might activate additional pathological processes that adversely affect reproductive function.²⁸

In our study, women with one lifetime pregnancy had poorer lifestyle factors (BMI, smoking), were older and less educated. The lower mean education among one-child mothers is in agreement with a Nordic demographic study,³⁴ which shows that later onset of childbearing is related to a lower number of children finally born in women with low education. Given that educational level and occupation are key indicators of socioeconomic status,³⁵ observed lower parity among women with low education could also reflect unfavourable socioeconomic position as a limiting factor to further pregnancies. However, a study exploring age at first birth, parity and postreproductive mortality suggests that late childbearing in itself may be a signal of pre-existing poor health of a woman.³⁶

The observed risk differences between nulliparous women and one-child mothers in our main results

(figure 2, table 2) could be explained by the heterogeneity of causes for childlessness among nulliparous women in this cohort. The risk may, therefore, be diluted by low-risk groups of women who are voluntarily childless³⁷ or have not been exposed to pregnancy (ever). Given the lack of information on women's reproductive planning in our data, we tried to address this in a subanalysis including only women with reported partner (ever) as a proxy for being exposed to pregnancy. Here we found that the results for nulliparous women were similar to our main results for one-child mothers. Women with reported partner had higher risk of having no children (compared with partnered women with ≥ 2 births) if their TG and TG:HDL-c ratio levels were in the highest quintiles and HDL in the lowest quintile (OR 1.9, 95% CI 0.9 to 4.2; OR 2.0, 95% CI 1.0 to 4.1; and OR 1.6, 95% CI 0.7 to 3.6, respectively). These findings support the role of serum lipids in lifetime nulliparity among women with partners.

Strengths and limitations

Our subset of women was from a large population-based health study with prepregnant health data. Linked data from the MBRN provided complete registration of total reproduction. The prospective design minimised the potential for bias. A weakness is that blood sampling was performed in a non-fasting state. Studies show that TG levels are sensitive to recent food intake, while cholesterol levels seem to be less affected.³⁸ We addressed this by adjusting our analyses for time since last meal and the main results were unchanged, suggesting that non-fasting lipids are not likely to introduce a systematic bias. Non-fasting lipids have successfully been used in lipid and CVD research.^{9 39 40}

The study lacked data on apolipoprotein E genotype, CRP (C-reactive protein) and thyroid tests/thyroid antibodies, factors that all are found to affect female fertility.^{23 41} No assessments of duration or temporal proximity of OC use, dietary intake or stress were available; therefore, unmeasured confounding cannot be ruled out. Smoking adversely influences female fertility,⁴² with most of its effects attributed to HDL cholesterol decrease.⁴³ We accounted for this in our analyses; however, smoking status of participants was only available at enrolment. The ethnic homogeneity of the included women may reduce the generalisability of our findings.

Unfavourable prepregnant lipid levels were associated with having no and one lifetime pregnancy. Women's metabolic homeostasis is important for reproduction and also has cardiometabolic implications.^{32 44} Pre-existing poor lipid and metabolic profiles could represent one of the possible linkages between previously observed reduced fertility and later CVD.

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Contributors All authors had full access to the data and are responsible for the integrity of the data. AP, RS, LAD and N-HM designed the study. AP and N-HM conducted the analyses. N-HM created the figure and AP created the tables and the flow chart. AP drafted the manuscript. N-HM, RS and LAD reviewed the preliminary analyses and initial draft and provided critical comments.

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Data sharing statement This data set contains personal data and cannot be made public due to confidentiality requirements according to Norwegian legislations. Researchers who are interested in analysing data from CONOR or the MBRN may apply to the appropriate organisations after having obtained all necessary approvals according to Norwegian legislations.

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