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Authors

Yu, Yongfu
Arah, Onyebuchi A
Liew, Zeyan
et al.

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Maternal diabetes during pregnancy and early onset of cardiovascular disease in offspring: population based cohort study with 40 years of follow-up

Yongfu Yu,^{1,2} Onyebuchi A Arah,^{2,3} Zeyan Liew,^{4,5} Sven Cnattingius,⁶ Jørn Olsen,^{1,2} Henrik Toft Sørensen,¹ Guoyou Qin,⁷ Jiong Li^{1,8}

For numbered affiliations see end of the article.

Correspondence to: G Qin
gyqin@fudan.edu.cn
(or @YongfuYu on Twitter;
ORCID 0000-0002-1413-3117)

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ABSTRACT

OBJECTIVE

To evaluate the associations between maternal diabetes diagnosed before or during pregnancy and early onset cardiovascular disease (CVD) in offspring during their first four decades of life.

DESIGN

Population based cohort study.

SETTING

Danish national health registries.

PARTICIPANTS

All 2 432 000 liveborn children without congenital heart disease in Denmark during 1977-2016. Follow-up began at birth and continued until first time diagnosis of CVD, death, emigration, or 31 December 2016, whichever came first.

EXPOSURES FOR OBSERVATIONAL STUDIES

Pregestational diabetes, including type 1 diabetes (n=22 055) and type 2 diabetes (n=6537), and gestational diabetes (n=26 272).

MAIN OUTCOME MEASURES

The primary outcome was early onset CVD (excluding congenital heart diseases) defined by hospital diagnosis. Associations between maternal diabetes and risks of early onset CVD in offspring were studied. Cox regression was used to assess whether a maternal history of CVD or maternal diabetic complications affected these associations. Adjustments were made for calendar year, sex, singleton status, maternal factors (parity, age, smoking, education, cohabitation, residence at childbirth, history of CVD before childbirth), and paternal history of CVD before childbirth. The cumulative incidence was averaged

across all individuals, and factors were adjusted while treating deaths from causes other than CVD as competing events.

RESULTS

During up to 40 years of follow-up, 1153 offspring of mothers with diabetes and 91 311 offspring of mothers who did not have diabetes were diagnosed with CVD. Offspring of mothers with diabetes had a 29% increased overall rate of early onset CVD (hazard ratio 1.29 (95% confidence interval 1.21 to 1.37); cumulative incidence among offspring unexposed to maternal diabetes at 40 years of age 13.07% (12.92% to 13.21%), difference in cumulative incidence between exposed and unexposed offspring 4.72% (2.37% to 7.06%). The sibship design yielded results similar to those of the unpaired design based on the whole cohort. Both pregestational diabetes (1.34 (1.25 to 1.43)) and gestational diabetes (1.19 (1.07 to 1.32)) were associated with increased rates of CVD in offspring. Varied increased rates of specific early onset CVDs were also observed, particularly heart failure (1.45 (0.89 to 2.35)), hypertensive disease (1.78 (1.50 to 2.11)), deep vein thrombosis (1.82 (1.38 to 2.41)), and pulmonary embolism (1.91 (1.31 to 2.80)). Increased rates of CVD were seen in different age groups from childhood to early adulthood until age 40 years. The increased rates were more pronounced among offspring of mothers with diabetic complications (1.60 (1.25 to 2.05)). A higher incidence of early onset CVD in offspring of mothers with diabetes and comorbid CVD (1.73 (1.36 to 2.20)) was associated with the added influence of comorbid CVD but not due to the interaction between diabetes and CVD on the multiplicative scale (P value for interaction 0.94).

CONCLUSIONS

Children of mothers with diabetes, especially those mothers with a history of CVD or diabetic complications, have increased rates of early onset CVD from childhood to early adulthood. If maternal diabetes does have a causal association with increased CVD rate in offspring, the prevention, screening, and treatment of diabetes in women of childbearing age could help to reduce the risk of CVD in the next generation.

Introduction

Cardiovascular disease (CVD) remains the leading cause of mortality and morbidity worldwide.^{1 2} Incidence and mortality rates of CVD have decreased in some countries,¹⁻⁴ but the prevalence of CVD has increased in children and young adults in recent decades.^{5 6}

WHAT IS ALREADY KNOWN ON THIS TOPIC

The prevalence of cardiovascular disease (CVD) has increased in children and young adults in recent decades

Maternal diabetes before or during pregnancy is associated with increased risks of metabolic syndrome and congenital heart disease in offspring

Whether prenatal exposure to maternal diabetes affects early onset CVD in offspring during their early decades of life is not known

WHAT THIS STUDY ADDS

Maternal diabetes during pregnancy was found to be associated with an increased rate of early onset CVD among offspring across the first four decades of life, especially for the offspring of those mothers with a history of CVD or diabetic complications

Preventing, screening, and treating diabetes in women of childbearing age could be important not only for improving the health of the women but also for reducing long term risks of CVD in their offspring

The risk factors for CVD change over the course of a lifetime,⁷ and early onset CVD could have a different cause than CVD diagnosed in later adulthood. Limited but growing evidence suggests intergenerational links between maternal health during or before pregnancy and risk factors for CVD among offspring.⁸⁻¹¹ A better understanding of such a link is essential to prevent and manage CVD among children and young adults.^{8 12-14}

The prevalence of pregestational and gestational diabetes has been increasing globally.^{15 16} In pregnancies complicated by diabetes, the diabetic intrauterine environment could cause placental dysfunction and hormonal alterations, leading to disease development.¹⁵ Prenatal exposure to maternal diabetes has also been associated with congenital heart disease, obesity, and diabetes in offspring. These diseases could, in turn, lead to an increased risk of CVD in later life.^{15 17-20} The offspring of mothers with diabetes also have a higher prevalence of metabolic syndrome and other risk factors for future CVD.^{9 21 22} It is unclear, however, whether or to what extent prenatal exposure to maternal diabetes increases the risk of CVD in offspring over a lifetime.

In this Danish population based cohort study, we hypothesised that intrauterine exposure to maternal diabetes could lead to an excess risk of early onset CVD in offspring from childhood to early adulthood (up to 40 years). Using data from Danish registers, we examined associations between types of maternal diabetes and early onset of CVD in offspring, the combined effect of maternal diabetes and maternal history of CVD on the rate of early onset CVD in offspring, and whether pregestational complications of diabetes further increased the risk of CVD in offspring.

Methods

Study population

The unique personal identification number assigned to all Danish residents allows accurate linkage of individual level information from all national registries (a detailed description of registers is provided in supplementary appendix 1).²³ We conducted a population based cohort study including all live births in Denmark during 1977-2016 (n=2 475 209). Our final cohort comprised 2 432 000 births after excluding 43 209 babies with congenital heart disease identified at birth or diagnosed later. A total of 27 046 (62.6%) of these babies with congenital heart disease were diagnosed up to 1 year, 8923 (20.7%) were diagnosed between 1 and 4 years of age, 3670 (8.5%) between 5 and 9 years, 2750 (6.4%) between 10 and 19 years, and 820 (1.9%) between 20 and 40 years. Follow-up started at birth and ended at the first diagnosis of a CVD, death, emigration, or 31 December 2016, whichever came first. People who emigrated or died from a cause other than CVD during follow-up were censored at the time of emigration or death.

Exposure

Offspring born to mothers diagnosed with diabetes before childbirth were considered to have been

prenatally exposed to maternal diabetes mellitus.²³ Maternal diabetes was categorised as gestational diabetes or pregestational diabetes. Pregestational diabetes was further defined as type 1 or type 2 diabetes before childbirth (a detailed description of the methods used to identify diabetes is provided in supplementary appendix 2). If a mother was diagnosed with more than one type of diabetes during one pregnancy, possibly owing to misclassification, she was classified in the main analysis in accordance with the first type diagnosed. We also used two other approaches in sensitivity analyses: hierarchical ordering of diabetes types (type 1, type 2, and then gestational diabetes)²⁰; and restriction to offspring of mothers with only one type of diabetes. Information on the diagnosis of diabetes was retrieved from the Danish National Diabetes Register, the Danish National Patient Registry (DNPR), and the Danish National Prescription Registry (supplementary appendix 1).

Because pregestational diabetic complications could reflect diabetes severity due to poor glycaemic control, we identified mothers with such complications—namely, diabetic coma; ketoacidosis; diabetes with kidney disease; and ophthalmic, neurological, circulatory, unspecified, or multiple complications. Women with pregestational diabetic complications were classified into two groups: those with one complication and those with multiple complications.

Outcome of interest

The outcome of interest was early onset CVD in offspring, defined as the first occurrence of CVD in the DNPR or the Danish Register of Cause of Death.²³ Diagnoses of congenital heart disease were excluded. We identified the outcome by using ICD-8 and ICD-10 (international classification of diseases, 8th and 10th revisions) codes for CVD (ICD-8: 390-444.1, 444.3-458, 782.4; ICD-10: I00-I99), or surgery codes for coronary artery bypass graft surgery and percutaneous coronary interventions (supplementary table S1). We also investigated the following types of CVD: ischaemic heart disease, cerebrovascular disease, stroke, heart failure, atrial fibrillation, hypertensive disease, deep vein thrombosis, pulmonary embolism, and other types of CVD (table S1).

Covariates

Potential confounders were selected based on directed acyclic graphs (fig S1) depicting the best known relations between the variables in this study. Confounders included maternal age (<20, 20-24, 25-29, 30-34, or ≥35 years), parity (1, 2, or ≥3 children), maternal cohabitation (single or cohabitating), maternal education (0-9, 10-14, or ≥15 years), maternal residence (Copenhagen, cities with 100 000 or more inhabitants, or other), maternal smoking during pregnancy (yes or no), maternal and paternal CVD history before the birth of their child (yes or no). We also included singleton delivery, sex of offspring, and calendar period of delivery (before 1980, five year intervals during 1981-2010, and the interval 2011-16).

To deal with the problem of missing data on covariates, we used the SAS multiple imputation procedure with fully conditional specification to impute five replications (chosen to be greater than the percentage of data missing as advised by White et al²⁴ and to make the computation tractable for our large dataset with millions of observations). The covariates and multiple imputations are described in detail in supplementary appendix 3. Missing indicator method and complete case analysis were also performed for comparison.

Statistical analysis

Cox regression was used to estimate hazard ratio with 95% confidence intervals to assess the association between exposure to maternal diabetes and overall CVD/specific types of CVD in offspring, with offspring's age as the time scale. Evaluation of log-minus-log survival curves showed that the curves were roughly parallel (fig S2). Although they appeared to be converging slightly, no obvious evidence was seen to support the interaction between hazard ratio and time (P value for interaction 0.70). Thus, we find it reasonable to assume that proportional hazard assumption was not violated. Treating deaths from causes other than CVD as competing events, we performed competing risk analysis to estimate the cumulative incidence and cumulative incidence difference between exposed and unexposed offspring averaged across the distribution of the covariates, described above using the inverse probability of treatment weighting approach.²⁵⁻²⁷ We evaluated the joint effect of maternal diabetes and maternal history of CVD on early onset CVD in offspring by including the interaction term between maternal diabetes and maternal history of CVD. Furthermore, we examined the association between maternal diabetes and CVD in offspring stratified by maternal diabetic complications and the age group of the offspring (0-9, 10-14, 15-19, 20-24, and 25-40 years).

In sensitivity analyses, we evaluated risks of CVD in offspring in relation to the timing of diagnosis of maternal type 1 diabetes with respect to childbirth (pregestational diagnoses and diagnoses ≤ 2 , 2-5, and >5 years after childbirth). Corresponding risks were also evaluated in relation to the timing of diagnosis of pregestational (type 1 and type 2) diabetes. Furthermore, we restricted analyses to offspring born at term (≥ 37 completed gestational weeks).

We used a sibship design to evaluate the influence of uncontrolled confounding due to shared genetic or familial characteristics by analysing the data of sibling pairs and comparing the outcome of each sibling exposed to maternal diabetes with the outcome of their unexposed sibling.²⁸ We performed conditional Poisson regression for the sibling subcohort analysis. Similar to the stratified Cox regression,²⁹ the conditional Poisson regression included a separate stratum for each family identified by the mother's unique civil registration number, in which only sibling pairs discordant for both maternal diabetes and offspring CVD were informative and contributed

to the effect estimate. Thus, each family had its own baseline rate function reflecting the family's shared genetic or familial characteristics. The association between maternal diabetes during pregnancy and early onset CVD in offspring was analysed only among siblings. Thus, the analysis controlled for genetic or familial characteristics shared among family members. As type 1 and type 2 diabetes would not change between pregnancies, and gestational diabetes was more common in older women with higher parity, the sibship design was appropriate for gestational diabetes in a second pregnancy or later pregnancies. We further analysed the data to assess the influence of possible uncontrolled confounding. We evaluated also whether parental factors, including maternal country of origin, maternal body mass index before pregnancy, and paternal diabetes before the child's birth, affected the observed associations. In addition, we used paternal diabetes as an exposure to explore potential confounding by genetic and familial factors. We used inverse-probability-of-selection weighting to evaluate possible live birth bias due to maternal diabetes and other (uncontrolled or unmeasured) risk factors for CVD in offspring.³⁰ These risk factors could lead to fetal loss such that naive analysis of data on live births could be misleading.^{30 31}

We performed subanalyses stratified by sex of offspring, and restricted to primiparous women, and singleton offspring. Owing to ICD code changes (ICD-10 was adopted in 1994) and the availability of data on confounders (maternal smoking and maternal pregnancy body mass index became available since 1991 and 2004, respectively), we restricted subanalyses to offspring born after 1991, 1994, and 2004. All analyses were conducted using SAS 9.4 (SAS Institute, Cary, NC) and Stata 14 (StatCorp, College Station, TX).

Patient and public involvement

No patients were involved in setting the research question or the outcome measures, nor were they involved in developing plans for design or implementation of the study. No patients were asked to advise on interpretation or writing up of results. There are no plans to disseminate the results of the research to study patients or the relevant patient community.

Results

Of 2 432 000 liveborn offspring without congenital heart disease, 54 864 (2.3%) were exposed to maternal pregestational diabetes (type 1: 0.9%, type 2: 0.3%) or gestational diabetes (1.1%). The proportion of offspring born to mothers with diabetes increased over time (fig S3). A total of 93 881 offspring (3.9%) were censored at the end of follow-up owing to emigration (n=74 377) or non-cardiovascular death (n=19 504). Compared with mothers who did not have diabetes, mothers with diabetes were more likely to be older, to have had higher education, to have higher parity, to live alone, and to smoke less during pregnancy. Compared with unexposed offspring, offspring exposed to maternal

Table 1 | Associations between maternal diabetes and early onset of overall CVD and specific types of CVD in offspring

Outcome* and exposure	No of offspring with CVD	Rate per 1000 person years	Hazard ratio (95% CI), model 1	Hazard ratio (95% CI), model 2
Overall CVD				
No diabetes	91 311	2.01	1.0 (ref)	1.0 (ref)
Maternal diabetes:	1153	2.02	1.47 (1.39 to 1.56)	1.29 (1.21 to 1.37)
Pregestational diabetes	792	2.28	1.45 (1.35 to 1.56)	1.34 (1.25 to 1.43)
Pregestational diabetes, type 1	531	2.13	1.46 (1.34 to 1.59)	1.31 (1.20 to 1.43)
Pregestational diabetes, type 2	261	2.68	1.44 (1.27 to 1.62)	1.39 (1.23 to 1.57)
Gestational diabetes	361	1.60	1.51 (1.36 to 1.67)	1.19 (1.07 to 1.32)
Ischaemic heart disease				
No diabetes	3753	0.08	1.0 (ref)	1.0 (ref)
Maternal diabetes:	28	0.05	1.21 (0.83 to 1.75)	1.18 (0.82 to 1.72)
Pregestational diabetes	26	0.07	1.38 (0.94 to 2.03)	1.37 (0.93 to 2.01)
Pregestational diabetes, type 1	16	0.06	1.35 (0.83 to 2.21)	1.35 (0.83 to 2.21)
Pregestational diabetes, type 2	10	0.10	1.42 (0.76 to 2.64)	1.40 (0.75 to 2.61)
Gestational diabetes	<3	0.01	0.46 (0.11 to 1.83)	0.43 (0.11 to 1.71)
Cerebrovascular disease				
No diabetes	7113	0.15	1.0 (ref)	1.0 (ref)
Maternal diabetes:	102	0.18	1.49 (1.22 to 1.81)	1.38 (1.13 to 1.68)
Pregestational diabetes	67	0.19	1.46 (1.15 to 1.86)	1.39 (1.10 to 1.77)
Pregestational diabetes, type 1	48	0.19	1.54 (1.16 to 2.05)	1.46 (1.10 to 1.94)
Pregestational diabetes, type 2	19	0.19	1.30 (0.83 to 2.03)	1.25 (0.79 to 1.96)
Gestational diabetes	35	0.15	1.54 (1.10 to 2.14)	1.36 (0.97 to 1.90)
Stroke				
No diabetes	4678	0.10	1.0 (ref)	1.0 (ref)
Maternal diabetes:	67	0.12	1.39 (1.09 to 1.77)	1.27 (1.00 to 1.62)
Pregestational diabetes	43	0.12	1.37 (1.01 to 1.85)	1.29 (0.96 to 1.74)
Pregestational diabetes, type 1	30	0.12	1.38 (0.97 to 1.98)	1.29 (0.90 to 1.85)
Pregestational diabetes, type 2	13	0.13	1.34 (0.77 to 2.30)	1.28 (0.74 to 2.21)
Gestational diabetes	24	0.11	1.43 (0.96 to 2.13)	1.23 (0.82 to 1.85)
Heart failure				
No diabetes	1222	0.03	1.0 (ref)	1.0 (ref)
Maternal diabetes:	17	0.03	1.48 (0.92 to 2.40)	1.45 (0.89 to 2.35)
Pregestational diabetes	11	0.03	1.42 (0.78 to 2.58)	1.40 (0.77 to 2.54)
Pregestational diabetes, type 1	5	0.02	0.95 (0.40 to 2.29)	0.95 (0.39 to 2.28)
Pregestational diabetes, type 2	6	0.06	2.41 (1.08 to 5.38)	2.32 (1.04 to 5.19)
Gestational diabetes	6	0.03	1.61 (0.72 to 3.60)	1.54 (0.68 to 3.47)
Atrial fibrillation				
No diabetes	2339	0.05	1.0 (ref)	1.0 (ref)
Maternal diabetes:	17	0.03	1.17 (0.73 to 1.89)	1.12 (0.69 to 1.81)
Pregestational diabetes	16	0.05	1.36 (0.83 to 2.22)	1.33 (0.81 to 2.18)
Pregestational diabetes, type 1	11	0.04	1.51 (0.83 to 2.73)	1.48 (0.82 to 2.67)
Pregestational diabetes, type 2	5	0.05	1.12 (0.47 to 2.69)	1.10 (0.46 to 2.64)
Gestational diabetes	<3	0.00	0.36 (0.05 to 2.56)	0.31 (0.04 to 2.23)
Hypertensive disease				
No diabetes	9615	0.21	1.0 (ref)	1.0 (ref)
Maternal diabetes:	135	0.23	2.10 (1.77 to 2.49)	1.78 (1.50 to 2.11)
Pregestational diabetes	100	0.28	1.99 (1.63 to 2.42)	1.78 (1.46 to 2.17)
Pregestational diabetes, type 1	58	0.23	1.81 (1.40 to 2.35)	1.57 (1.22 to 2.04)
Pregestational diabetes, type 2	42	0.42	2.29 (1.69 to 3.10)	2.18 (1.61 to 2.95)
Gestational diabetes	35	0.15	2.50 (1.79 to 3.48)	1.77 (1.27 to 2.48)
Deep vein thrombosis				
No diabetes	4514	0.10	1.0 (ref)	1.0 (ref)
Maternal diabetes:	50	0.09	1.86 (1.41 to 2.46)	1.82 (1.38 to 2.41)
Pregestational diabetes	44	0.13	1.99 (1.48 to 2.68)	1.97 (1.47 to 2.66)
Pregestational diabetes, type 1	22	0.09	1.62 (1.07 to 2.47)	1.60 (1.05 to 2.43)
Pregestational diabetes, type 2	22	0.22	2.56 (1.68 to 3.89)	2.58 (1.70 to 3.93)
Gestational diabetes	6	0.03	1.26 (0.57 to 2.82)	1.17 (0.53 to 2.62)
Pulmonary embolism				
No diabetes	2246	0.05	1.0 (ref)	1.0 (ref)
Maternal diabetes:	27	0.05	1.85 (1.26 to 2.70)	1.91 (1.31 to 2.80)
Pregestational diabetes	21	0.06	1.82 (1.18 to 2.80)	1.86 (1.21 to 2.85)
Pregestational diabetes, type 1	7	0.03	0.97 (0.46 to 2.04)	1.01 (0.48 to 2.12)
Pregestational diabetes, type 2	14	0.14	3.22 (1.90 to 5.44)	3.22 (1.90 to 5.45)
Gestational diabetes	6	0.03	1.95 (0.87 to 4.36)	2.14 (0.96 to 4.79)
Other CVDst				
No diabetes	69 915	1.53	1.0 (ref)	1.0 (ref)
Maternal diabetes:	876	1.53	1.40 (1.31 to 1.50)	1.22 (1.14 to 1.30)
Pregestational diabetes	582	1.67	1.36 (1.26 to 1.47)	1.25 (1.15 to 1.35)
Pregestational diabetes, type 1	397	1.59	1.38 (1.26 to 1.53)	1.23 (1.12 to 1.36)

Table 1 | Continued

Outcome* and exposure	No of offspring with CVD	Rate per 1000 person years	Hazard ratio (95% CI), model 1	Hazard ratio (95% CI), model 2
Pregestational diabetes, type 2	185	1.89	1.31 (1.14 to 1.51)	1.28 (1.11 to 1.47)
Gestational diabetes	294	1.30	1.50 (1.34 to 1.68)	1.16 (1.04 to 1.30)

CVD=cardiovascular disease; ICD-8, ICD-10=international classification of disease, 8th revision, 10th revision; ref=reference; model 1=offspring's age as time scale; model 2=offspring's age as time scale, and controlled for calendar year, sex, singleton status, parity, maternal smoking, maternal education, maternal cohabitation, maternal residence at birth, maternal history of CVD before childbirth, paternal history of CVD before birth of the child, and maternal age (restricted cubic spline with five knots at five evenly spaced centiles).

*Ischaemic heart disease (ICD-8 codes: 410-414; ICD-10 codes: I20-I25), cerebrovascular disease (ICD-8 codes: 430-438; ICD-10 codes: I60-I69), stroke (ICD-8 codes: 430-436; ICD-10 codes: I61-I64), heart failure (ICD-8 codes: 427.0, 427.1, 782.4; ICD-10 codes: I110, I130, I132, I50), atrial fibrillation (ICD-8 codes: 427.93, 427.94; ICD-10 code: I48), hypertensive disease (ICD-8 codes: 400-404; ICD-10 codes: I10-I15), deep vein thrombosis (ICD-8 codes: 451.00; ICD-10 codes: I80.1-I80.3), pulmonary embolism (ICD-8 codes: 450.99; ICD-10 codes: I26), other types of CVD (the remainder of the codes included under CVD overall (ICD-8 codes: 390-444.1, 444.3-458, 782.4; ICD-10 codes: I00-I99)).

†The most important diagnostic entities contributing to this category: cardiomyopathy, cardiac arrest, paroxysmal tachycardia, complications and ill-defined descriptions of heart disease, diseases of capillaries, varicose veins of other sites, non-specific lymphadenitis, and hypotension.

diabetes were more likely to have a parental history of CVD and to have a higher rate of developing diabetes, obesity, hypertension, hypercholesterolaemia, and chronic kidney diseases (tables S2-3).

During up to 40 years of follow-up, 1153 offspring of mothers with diabetes and 91 311 offspring of mothers who did not have diabetes were diagnosed with CVD. Exposed offspring had a higher rate of overall CVDs than unexposed offspring (hazard ratio 1.29 (95% confidence interval 1.21 to 1.37); cumulative incidence among unexposed offspring at 40 years of age (13.07% (12.92% to 13.21%)), cumulative incidence difference between the exposed and the unexposed offspring (4.72% (2.37% to 7.06%); table 1, fig 1, and fig S4). An increased rate of early onset CVD was seen in offspring exposed to pregestational diabetes (1.34 (1.25 to 1.43)) or gestational diabetes (1.19 (1.07 to 1.32)). Early onset of CVD in offspring was of similar magnitude in maternal type 1 and type 2 diabetes. The rates for most specific types of CVD were also increased in offspring exposed to maternal diabetes, with higher rates seen for heart failure (1.45 (0.89 to 2.35)), hypertensive disease (1.78 (1.50 to 2.11)), deep vein thrombosis (1.82 (1.38 to 2.41)), and pulmonary embolism (1.91 (1.31 to 2.80); table 1). We generally observed increased hazard ratios for offspring exposed to maternal diabetes in each age group in childhood (before 20 years of age) and early adulthood (from 20 to 40 years of age), regardless of the type of maternal diabetes (fig 2).

We found a higher incidence of CVD in offspring of mothers with both diabetes and comorbid CVD (1.73 (1.36 to 2.20)) than in offspring of mothers with diabetes only (1.29 (1.21 to 1.37)). This increased incidence was due to the added influence of comorbid CVD but not their interaction on the multiplicative scale (P value for interaction 0.94; table 2). Offspring of mothers with pregestational diabetes and diabetic complications had a higher incidence of CVD (1.60 (1.25 to 2.05)) than the offspring of mothers with pregestational diabetes but no diabetic complications (1.31 (1.16 to 1.48); table 3), representing a 22% increased rate (1.22 (0.92 to 1.62)) due to diabetic complications. Our study population for this analysis was young (up to 21 years of age with a median age of 11), and only a limited number of CVD events (n=62) in offspring of mothers with diabetes with complications were available for this analysis.

Sensitivity analysis yielded similar findings to those of the main analyses (tables S4-6), including use of hierarchical ordering of diabetes types (type 1, type 2, and gestational diabetes), and restriction to offspring of mothers diagnosed with only one diabetes type during their pregnancy; or offspring born at term. The results from sibship design in the sibling cohort (1.26 (1.18 to 1.35)) were similar to those of the main analysis in the unmatched whole population cohort. Moreover, for the timing of the diagnosis of maternal type 1 diabetes, the association between maternal type 1 diabetes and CVD was strongest when mothers were diagnosed before childbirth (1.35 (1.23 to 1.48)). The association was attenuated with greater time to diagnosis after birth (table S7). The pattern was similar when type 1 and type 2 diabetes were combined. The results were essentially unchanged when we controlled for maternal country of origin, maternal body mass index before pregnancy, and paternal diabetes before the birth of the child, either separately or together in the model (table S8). In addition, paternal diabetes remained weakly associated with early onset CVD in offspring (table S9).

In our evaluation of live birth bias, the inverse-probability-weighting approach yielded results similar to those of the primary analyses, showing that maternal diabetes increased the rate of early onset CVD in offspring (table S10). The analysis stratified by sex of the offspring showed similar results. Results from separate analyses restricted to singletons, offspring of primiparous women, individuals with complete data, and offspring born after 1991, 1994, or 2004, and using the missing indicator method, were similar to those obtained in the primary analyses (table S11).

Discussion

In this large population based cohort study, we found that maternal diabetes before or during pregnancy was associated with increased rates of early onset CVD in general, and with most specific types of early onset CVD in offspring, persisting from childhood through early adulthood. The strongest associations were found among offspring of mothers with diabetic complications or with diabetes and a history of CVD.

Comparisons with other studies

During pregnancies complicated by diabetes, large amounts of maternal glucose freely cross the placenta, which could lead to increased secretion of fetal

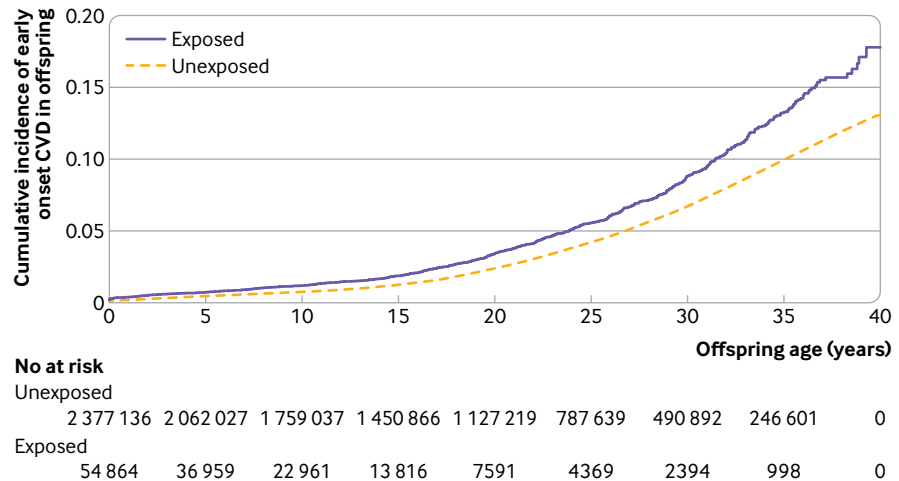


Fig 1 | Adjusted cumulative incidence of early cardiovascular disease (CVD) onset among offspring exposed versus unexposed to maternal diabetes. The adjusted cumulative incidence was averaged across the distribution of the covariates—calendar year, sex, singleton status, parity, age, smoking, education, cohabitation, residence at childbirth, history of CVD before childbirth, and paternal history of CVD before birth of the child—using the inverse probability of treatment weighting approach

insulin.¹⁵ This increase would result in a state of hyperinsulinaemia and hyperglycaemia in the fetal circulation.¹⁵ Exposure to hyperinsulinaemia and hyperglycaemia in utero could have longlasting effects on fetal vascular gene expression and result in changes in vascular function, thereby contributing to higher CVD risks in offspring.^{15 32} The results of animal studies^{33 34} indicated abnormalities in vascular reactivity and increased risks of hypertension and cardiovascular dysfunction in offspring of diabetic rats. Several human studies have shown that the diabetic intrauterine environment could have a programming effect on fetal vascular dysfunction, leading to a poor CVD risk profile after birth.^{15 32} This profile would include increased concentrations of markers of endothelial dysfunction, macrosomia, increased aortic intima-media thickness, and arterial stiffening.^{15 32} Poor glycaemic control in pregnant women could result in more unhealthy fetuses with more adverse birth outcomes,^{15 35-37} which are risk factors for future CVD.³⁸ In addition, offspring of mothers with diabetes tend to be at increased risk of exposure to many classic cardiovascular risk factors, such as hypercholesterolaemia, hypertension, obesity, diabetes, chronic kidney disease, and metabolic syndrome.^{9 10 15 32 37 39-41} Moreover, factors such as genetic susceptibility, family environment variables, or lifestyle characteristics might also contribute to the increased risk of CVD in children born to mother with diabetes.^{15 37}

Empirical evidence is lacking for an association of maternal diabetes during pregnancy with the overall risk of early onset CVD in offspring, with the exception of an increased risk of congenital heart diseases.^{18 42} One study reported that the risk of congenital heart disease was higher among offspring exposed to pregestational diabetes than among those exposed to gestational diabetes.¹⁸ A Swedish study found

that pregestational and gestational diabetes were associated with congenital heart defects.⁴² In our large study, we found increased rates of early onset CVD in offspring prenatally exposed to three types of maternal diabetes (pregestational type 1 and type 2, and gestational diabetes). The variation in the magnitudes of the estimated effect size might be due to the different mechanisms underlying different types of diabetes.^{15 43} The similarity in disease risk due to maternal type 1 versus type 2 diabetes for early onset CVD and congenital heart disease, respectively, suggests a shared pathological process between both types of diabetes in fetal heart development.^{18 43} The higher CVD rate in offspring of those with pregestational diabetes, compared with offspring of mothers with gestational diabetes, suggests that a high glucose level in early pregnancy has a major role in CVD development in offspring.^{18 44 45} The difference in associations found between maternal diabetes and specific types of CVD could stem from different underlying mechanisms for CVD.^{46 47}

Notably, an almost twofold risk of higher early onset of CVD was seen in offspring of mothers with diabetes and a history of CVD, compared with the offspring of mothers with no such history. The greater effect of coexisting maternal CVD and maternal diabetes on CVD risk in offspring requires additional research to examine the burden of multimorbidity during pregnancy.

Higher insulin resistance is also associated with an increased risk of pregestational diabetic complications.⁴⁸ Therefore, diabetic complications due to poor glycaemic control might reflect the severity of pregestational diabetes. Our observation suggests that children of mothers with diabetic complications have a higher risk of very early onset CVD. This result is consistent with earlier studies that found a higher risk of genital anomalies and congenital heart diseases in

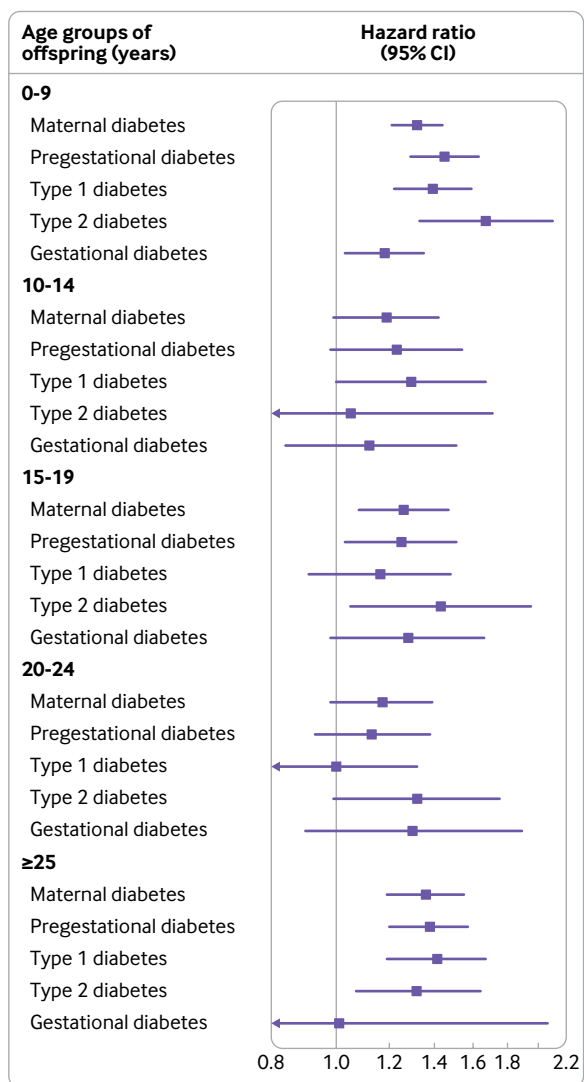


Fig 2 | Associations between maternal diabetes and early onset of cardiovascular disease (CVD) in offspring by offspring age and diabetes type. Maternal diabetes included gestational diabetes and pregestational diabetes. Pregestational diabetes included type 1 and type 2 diabetes. Associations were controlled for calendar year, sex, singleton status, parity, maternal smoking, maternal education, maternal cohabitation, maternal residence at birth, maternal history of CVD before childbirth, paternal history of CVD before birth of the child, and maternal age (restricted cubic spline with five knots at five evenly spaced centiles). Arrows indicate a lower limit of 0.8

offspring of mothers with diabetic complications.^{18 20} Since the diagnostic code for diabetic complications was available only in ICD-10 (since 1994), our study population was relatively too young for enough CVD events to have occurred for this analysis. Further follow-up and investigation in other study settings would be valuable to confirm these findings.

Strengths and limitations of this study

The main strength of our study is the high quality, prospectively collected data covering all Danish residents with nearly complete follow-up, thus minimising the possibility of recall or selection bias. The large sample size and long follow-up of up to 40 years

provide an opportunity to investigate specific types of CVD and to evaluate the long term consequences of maternal diabetes for offspring of different ages. In addition, we used a sibship design to evaluate the influence of uncontrolled confounding due to shared familial (genetic or environmental) characteristics. Such (unmeasured) characteristics are often difficult to control in a conventional cohort study. We could also adjust for a wide range of covariates, such as maternal sociodemographic and lifestyle variables, and history of CVD. Previous studies have reported that maternal diabetes could increase the risks of preterm birth, pre-eclampsia, and macrosomia^{15 35-37}; therefore, these factors could lie in the pathway from maternal diabetes during pregnancy to early onset CVD in offspring. Thus, we considered these variables to be primarily potential mediators, and did not adjust for these factors in the main analysis, as reported in the main text. Nevertheless, for explorative purposes, in an additional analysis, we did adjust for maternal pre-eclampsia, preterm birth, and large for gestational age, and found that only slightly attenuated associations were seen, compared with the main analysis (table S11).

Several limitations must also be noted. Firstly, although we adjusted for a wide range of confounders, we could not rule out the possibility of residual confounding by some uncontrolled genetic or familial environment or lifestyle characteristics, such as maternal alcohol consumption, glycaemic control, folic acid supplementation, prenatal care, physical inactivity, psychological stress, offspring smoking, and genetic variants, in fetal susceptibility to the diabetic intrauterine environment. Residual confounding of unmeasured factors may partially account for the association between maternal diabetes and early onset CVD in offspring. However, our sibship design yielded results similar to those of the unpaired design based on the whole cohort. We have to acknowledge that, although the sibship design has a number of methodological strengths, it also has several drawbacks.⁴⁹⁻⁵¹ For example, compared with an unpaired design (which was also used in our study), the sibship design may lead to bias in estimating exposure effects if carryover or spillover effects are present.^{50 51} In addition, a sibship design cannot adequately control for unmeasured confounders that change across pregnancies. Moreover, the sibship design restricts the analysis to women who have had at least two pregnancies, leading to loss of generalisability to the target population. Thus, the sibship design could not fully rule out the possibility of uncontrolled confounding, possibly requiring bias analysis.⁴⁹⁻⁵² The strongest associations were for offspring born to mothers with a diagnosis of diabetes before childbirth. This association was attenuated when maternal diabetes was diagnosed after childbirth. These findings suggest that non-genetic effects are important determinants of early onset CVD in offspring, and that the observed associations are not entirely attributable to confounding by genetics and familial environment.

Table 2 | Joint effect of maternal diabetes and maternal CVD history before childbirth on early CVD onset in offspring

Exposure	No of offspring with CVD	Rate per 1000 person years	Hazard ratio (95% CI), model 1	Hazard ratio (95% CI), model 2
No maternal diabetes and CVD	89 372	2.00	1.0 (ref)	1.0 (ref)
Maternal diabetes only	1086	1.99	1.45 (1.37 to 1.54)	1.29 (1.21 to 1.37)
Maternal CVD only	1939	2.23	1.41 (1.35 to 1.48)	1.33 (1.27 to 1.39)
Joint effect of maternal diabetes and CVD	67	2.46	2.10 (1.65 to 2.67)	1.73 (1.36 to 2.20)
Maternal diabetes×CVD	—	—	1.02 (0.80 to 1.31)	1.01 (0.79 to 1.30)
P value for interaction (maternal diabetes×CVD)	—	—	0.86	0.94

CVD=cardiovascular disease; model 1=offspring's age as time scale; model 2=offspring's age as time scale, and controlled for calendar year, sex, singleton status, parity, maternal smoking, maternal education, maternal cohabitation, maternal residence at birth, maternal history of CVD before childbirth, paternal history of CVD before birth of the child, and maternal age (restricted cubic spline with five knots at five evenly spaced centiles).

In addition, the considerably greater influence of maternal diabetes, compared with paternal diabetes, on the risk of CVD in offspring, further suggests that our findings are unlikely to be attributable entirely to uncontrolled confounding.

Secondly, there is a possibility of live birth selection bias, which we assessed using information available on stillbirths. Our sensitivity analyses provided reassurance that restricting our sample to live births was unlikely to affect the observed associations appreciably.

Thirdly, potential misclassification bias might remain, because the same code (ICD-8:250) was used to record type 1 and type 2 diabetes before 1986.¹⁸ However, the ascertainment and verification of diabetes in Denmark are considered highly reliable, implying that misclassification of maternal diabetes overall is unlikely.⁵³ A validation study that examined cardiovascular diagnoses in the DNPR in 2010-2012 indicated that cardiovascular diagnoses were of overall high quality and adequate for epidemiological research.⁵⁴ Although we could not rule out misclassification bias, we think that it would probably be non-differential, would tend to attenuate our estimates, and would not change our conclusion substantially.

Finally, our study could not examine the risk of CVD offspring in late adulthood because the longest follow-up period so far is 40 years. The maximum follow-up period for CVD events in offspring of mothers with diabetic complications was only 21 years, which might

prevent us from obtaining stable risk estimates for some subgroup analyses. However, this is the best available evidence and we expect the associations to be even stronger when older offspring can be studied. Future studies with longer follow-up are well warranted.

Conclusion and policy implications

The diabetic intrauterine environment could have a programming effect on the development of CVD in offspring. Our study provides evidence that children of mothers with diabetes, especially those with a history of CVD or with diabetic complications, had increased rates of early onset CVD throughout the early decades of life. These findings highlight the importance of effective strategies for screening and preventing diabetes in women of childbearing age. We need to monitor CVD risks in offspring of mothers with diabetes and investigate possible life course interventions that could reduce the occurrence of CVD. A history of CVD or diabetic complications in women with diabetes should be taken into account in designing public health strategies that target offspring at increased risk of early onset CVD. Future research should examine the degree of glycaemic control during pregnancy that would minimise the risk of CVD in offspring throughout their life.

AUTHOR AFFILIATIONS

¹Department of Clinical Epidemiology, Aarhus University Hospital, Aarhus, Denmark

²Department of Epidemiology, Fielding School of Public Health, University of California, Los Angeles (UCLA), CA, USA

Table 3 | Associations between maternal pregestational diabetes with or without diabetic complications and early onset of CVD in offspring born during 1996-2016*

Exposure	No of offspring with CVD	Rate per 1000 person years	Hazard ratio (95% CI), model 1	Hazard ratio (95% CI), model 2
No pregestational diabetes	16 251	1.22	1.0 (ref)	1.0 (ref)
Pregestational diabetes without diabetic complications	266	1.64	1.38 (1.23 to 1.56)	1.31 (1.16 to 1.48)
Pregestational diabetes with diabetic complications	62	1.99	1.67 (1.30 to 2.14)	1.60 (1.25 to 2.05)
1 complication	24	1.90	1.59 (1.07 to 2.37)	1.55 (1.04 to 2.32)
≥2 complications	38	2.05	1.73 (1.25 to 2.37)	1.63 (1.18 to 2.24)
Pregestational diabetes with diabetic complication v without diabetic complication	—	—	1.21 (0.91 to 1.59)	1.22 (0.92 to 1.62)

CVD, cardiovascular disease; HR, hazard ratio; ICD-10=international classification of disease, 10th revision; model 1=offspring's age as time scale; model 2=offspring's age as time scale, and controlled for calendar year, sex, singleton status, parity, maternal smoking, maternal education, maternal cohabitation, maternal residence at birth, maternal history of CVD before childbirth, paternal history of CVD before birth of the child, and maternal age (restricted cubic spline with five knots at five evenly spaced centiles).

*Pregestational diabetic complications: ICD-10 codes (E10.0–E10.8, E11.0–E11.8, and H36.0). Women with pregestational diabetic complications were classified as having one or multiple complications (ICD-10: E10.7, E11.7; ≥2 complications as provided above)

³Department of Statistics, College of Letters and Science, University of California, Los Angeles (UCLA), CA, USA

⁴Department of Environmental Health Sciences, Yale School of Public Health, New Haven, CT, USA

⁵Yale Center for Perinatal, Pediatric, and Environmental Epidemiology, Yale School of Public Health, New Haven, CT, USA

⁶Division of Epidemiology, Department of Medicine Solna 17176, Karolinska University Hospital, Karolinska Institute, Stockholm, Sweden

⁷Department of Biostatistics, School of Public Health, and The Key Laboratory of Public Health Safety of Ministry of Education, Fudan University, Shanghai 200032, China

⁸Ministry of Education—Shanghai Key Laboratory of Children's Environmental Health, Xinhua Hospital, Shanghai Jiao Tong University School of Medicine, Shanghai, China

Contributors: GQ (gyqin@fudan.edu.cn) and JL (jl@clin.au.dk) are corresponding authors and senior authors who contributed equally to this study. YY, GQ, and JL have full access to all the data in this study and take full responsibility as guarantors for the integrity of the data and the accuracy of the data analysis. YY, OAA, GQ, and JL conceived and designed the study. YY, OAA, and GQ undertook the statistical analysis. YY drafted the manuscript. All authors provided critical input to the analyses, interpreted the data, and revised the manuscript critically. The corresponding author confirms that all listed authors meet authorship criteria and that no others meeting the criteria have been omitted.

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Data sharing: No additional data available.

The lead authors (the manuscript's guarantors) affirm that the manuscript is an honest, accurate, and transparent account of the study being reported; that no important aspects of the study have been omitted; and that any discrepancies from the study as planned have been explained.

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Web appendix: Supplementary material

Supplementary Material

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Figure S4. Adjusted cumulative incidence of early CVD onset among offspring, exposed to maternal diabetes versus unexposed

Appendix S1. Detailed description of registers used in this study

The Danish National Health Service provides universal tax-supported healthcare, guaranteeing unfettered access to general practitioners and hospitals, and partial reimbursement for prescribed drug treatment. Since the 1960s, when the population registration systems started, all live births and new residents in Denmark have been assigned a unique individual personal identification number (Central Personal Register number, CPR). The CPR is encrypted by a unique code, which can be used for linkage between different national registers and ensures that national data protection, privacy, and confidentiality requirements are fulfilled.

The Danish Civil Registration System (CRS)

The Danish Civil Registration System (CRS)¹ was established on April 2, 1968, and registers all people living in Denmark. The CRS includes the unique individual personal identification number, sex, date and place of birth, current address, marital status, spouse, parents, children, and vital statistics.

The Danish Medical Birth Registry (MBR)

The Danish Medical Birth Registry (MBR)² was established in 1968, and has been computerized since 1973. The MBR includes birth characteristics, such as gestational age, birth weight, Apgar score at 5 minutes, parity, date of birth, sex, singleton or not, and maternal smoking during pregnancy.

The Danish National Patient Register (DNPR)

The Danish National Patient Register (DNPR)³ was established in 1977. The DNPR contains hospital discharge diagnoses from 1977, and outpatient and emergency diagnoses are included from 1995 and onwards. We used the Danish National Patient Registry covering all Danish hospitals to identify the study population, comorbidities within the study population, and non-fatal endpoints. Each hospital discharge or outpatient visit is recorded in the registry with one primary diagnosis and potentially several secondary

diagnoses, each classified according to the ICD-8 codes (*International Classification of Disease* codes, 8th revision) and ICD-10 codes when implemented.

The Danish Register of Causes of Death

The Danish Register of Causes of Death⁴ has been computerized since 1970. Danish law mandates registration of the date and cause of death for all Danish citizens who die in Denmark. The ICD-8 was used to classify the causes of death during 1970-1993 and the ICD-10 was used from 1994 onward.

The Danish National Prescription Registry

The Danish National Prescription Registry⁵ contains information on all prescriptions dispensed at community pharmacies in Denmark since 1995, including date of dispensing and drug class according to the Anatomical Therapeutic Chemical [ATC] classification system.

The Danish National Diabetes Register

The Danish National Diabetes Register^{6,7} was established based on data from existing Danish health registers, including from the Danish National Registry of Patients³, the National Health Insurance Service Register⁸, the Danish National Prescription Registry⁵, and the Civil Registration System¹. The Danish National Diabetes Register contains information on individuals diagnosed with diabetes by general and specialist practitioners in Denmark. Individuals are classified as having diabetes if they are registered with at least one of the following criteria: (1) a diagnosis of diabetes (ICD-8 codes: 249, 250; ICD-10 codes: E10-E11, H36.0, O24 excluding O24.4 and O24.9); (2) receipt of chiropody for diabetic patients; (3) two redeemed prescriptions for insulin (ATC code: A10A); (4) two redeemed prescriptions for oral antidiabetics (ATC code: A10B); or (5) two blood glucose measurements per year in five consecutive years.

The Danish Integrated Database for Longitudinal Labour Market Research

The Danish Integrated Database for Longitudinal Labour Market Research⁹ was established in 1981, and contains information on personal labour market affiliation and education.

Appendix S2. Detailed description of methods used to identify diabetes

Information on diabetes diagnoses was retrieved from the Danish National Diabetes Register (1995-2016),⁷ the Danish National Patient Registry (1977-2016),¹⁰ and the Danish National Prescription Registry (1994-2016)⁵ using *International Classification of Disease* codes (ICD-8 codes during 1970-1993 and ICD-10 codes since 1994) and *Anatomical Therapeutic Chemical* (ATC) classification codes. Diabetes was categorized as gestational diabetes (ICD-8 codes: 634.74, Y6449; ICD-10 codes: O24.4, O24.9) or pregestational diabetes. Pregestational diabetes was ascertained using the following criteria: (1) an diagnosis of diabetes (ICD-8 codes: 249, 250; ICD-10 codes: E10-E11, H36.0, O24 excluding O24.4 and O24.9); (2) receipt of chiropody for diabetic patients; (3) two redeemed prescriptions for insulin (ATC code: A10A); or (4) two redeemed prescriptions for oral antidiabetics (ATC code: A10B). Blood glucose measurements were not used due to their poor validity.⁶ Pregestational diabetes was further classified as type 1 diabetes (ICD-8 code: 249; ICD-10 codes: E10, O24.0; ATC code: A10A) or type 2 diabetes (ICD-8 code: 250; ICD-10 codes: E11, O24.1; ATC code: A10B). Pregestational diabetes was defined as the diagnosis of type 1 or type 2 diabetes before giving birth rather than before conception, because preexisting but unrecognized diabetes was likely to be identified and registered during pregnancy. Because diabetes was recorded using a single code (250) during 1977-1986, we used two approaches to distinguish between type 1 and type 2 diabetes during that period: 1) a specific code for type 1 or type 2 diabetes applied later in time; or 2) age of diabetes onset [cut-off for type 1: <30 years and cut-off for type 2: ≥ 30 years].^{11,12} If a mother was diagnosed with multiple types of diabetes during one pregnancy, she was classified according to the first diagnosed type. Once a mother was diagnosed with a specific code for type 1 or type 2 diabetes, she retained this status for any subsequent pregnancy. A mother with gestational diabetes diagnosed in one pregnancy was not assumed to have gestational diabetes or preexisting diabetes in the future pregnancies unless a further diagnosis was made.

Appendix S3. Detailed description of covariates and multiple imputation

Information on maternal and birth characteristics was retrieved from the Danish Medical Birth Register,² and socioeconomic factors were retrieved from the Danish Integrated Database for Longitudinal Labour Market Research and the Danish Civil Registry System.¹³ Information on parental CVD history was obtained from the Danish National Patient Registry.¹⁰ These included maternal age (<20, 20-24, 25-29, 30-34, or ≥ 35 years), parity (1, 2, or ≥ 3 children), maternal cohabitation (single or cohabitating), maternal education (0-9, 10-14, or ≥ 15 years), maternal residence (Copenhagen, cities with $\geq 100,000$ inhabitants, or other), maternal smoking during pregnancy (yes or no), maternal and paternal CVD history before the birth of their child (yes or no). We also included singleton delivery (yes or no), sex of offspring (male or female), and calendar period of delivery (before 1980, or 5-year intervals during 1981-2010 or 2011-2016). Maternal age was included as a continuous covariate in the regression models using restricted cubic splines rather than categorization.

The frequency of missing data in our main analysis was relatively low: 0.04% for sex, 2.27% for maternal education, 0.14% for maternal cohabitation, and 0.30% for paternal history of CVD. 97.41% offspring have complete information for these covariates. The frequency of missing data for maternal smoking (data available since 1991) was 4.31%. Regarding the other variables used in this study, the frequency of missing data were 0.30% for maternal country of origin, 0.30% for paternal history of diabetes before childbirth, and 5.71% for maternal prepregnancy BMI (data available since 2004).

We imputed missing variables using the fully conditional specification (FCS) method in which logistic regression was used for categorical variables and predictive mean matching for continuous variables. The imputation model included the following variables: maternal diabetes during pregnancy (yes or no), offspring CVD (yes or no), maternal age (continuous variables), parity (1, 2, or ≥ 3 children), maternal cohabitation (single or cohabitating), maternal education (0-9, 10-14, or ≥ 15 years), maternal residence (Copenhagen, cities with $\geq 100,000$ inhabitants, or other), maternal smoking during pregnancy (yes or no), maternal and paternal CVD history before the birth of their child (yes or no), maternal country of origin

(Nordic countries, others), paternal history of diabetes before childbirth (yes or no), singleton delivery (yes or no), sex of offspring (male or female), and calendar period of delivery (before 1980, or 5-year intervals during 1981-2010 or 2011-2016). As the data on maternal prepregnancy BMI was available since 2004, the imputation model for maternal prepregnancy BMI was only restricted to offspring born since 2004. The distributions of observed and imputed variables were similar.

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Table S1. Definition of cardiovascular disease^a

	ICD-8	ICD-10	Procedure/Surgery Codes
Overall cardiovascular disease (CVD)	390-444.1, 444.3-458, 782.4	I00-I99	30350, 30354, 30240, KFNG, KFNF 30009, 30019, 30029, 30039, 30049, 30059, 30069, 30079, 30089, 30099, 30109, 30119, 30120, 30129, 30139, 30149, 30159, 30169, 30179, 30189, 30199, 30200, KFNA-KFNE, KFNH20
Ischemic heart disease	410-414	I20-I25	
Cerebrovascular disease	430-438	I60-I69	
Stroke	430-436	I61-I64	
Heart failure	427.0,427.1,782.4	I110, I130, I132, I50	
Atrial fibrillation	427.93, 427.94	I48	
Hypertensive disease	400-404	I10-I15	
Deep vein thrombosis	451.00	I80.1-I80.3	
Pulmonary embolism	450.99	I26	
Coronary artery bypass graft (CABG)			30009, 30019, 30029, 30039, 30049, 30059, 30069, 30079, 30089, 30099, 30109, 30119, 30120, 30129, 30139, 30149, 30159, 30169, 30179, 30189, 30199, 30200, KFNA-KFNE, KFNH20
Percutaneous coronary intervention (PCI)			30350, 30354, 30240, KFNG, KFNF
Other CVDs	Remainder of codes (390-444.1, 444.3-458, 782.4)	Remainder of codes (I00-I99)	Without the surgery codes above for PCI and CABG

^a The information on cardiovascular disease is from the Danish National Patient Registry or the Danish Register of Cause of Death.

Table S2. Baseline characteristics of offspring exposed *in utero* to maternal diabetes, in Denmark, 1977-2016

Variable ^a	Exposure to type 1 diabetes (n=22 055)	Exposure to type 2 diabetes (n=6537)	Exposure to gestational diabetes (n=26 272)	Unexposed (n=2 377 136)
Singleton				
No	948 (4)	365 (6)	1554 (6)	75 294 (3)
Yes	21 107 (96)	6172 (94)	24 718 (94)	2 301 842 (97)
Sex				
Male	11 349 (51)	3370 (52)	13 715 (52)	1 219 022 (51)
Female	10 705 (49)	3163 (48)	12 553 (48)	1 157 052 (49)
Missing	1 (0)	4 (0)	4 (0)	1062 (0)
Maternal parity				
1	9222 (42)	2304 (35)	10 519 (40)	1 069 854 (45)
2	8421 (38)	2374 (36)	9320 (35)	882 200 (37)
≥3	4412 (20)	1859 (28)	6433 (24)	425 082 (18)
Maternal age at childbirth, years				
<20	188 (1)	58 (1)	156 (1)	59 566 (3)
20-24	2342 (11)	630 (10)	1 893 (7)	434 191 (18)
25-29	7020 (32)	1484 (23)	6754 (26)	869 159 (37)
30-34	7682 (35)	2097 (32)	9429 (36)	702 488 (30)
35+	4823 (22)	2268 (35)	8040 (31)	311 732 (13)
Maternal smoking during pregnancy ^b				
No	15 924 (80)	3 701 (76)	20 902 (81)	1 223 564 (77)
Yes	3194 (16)	903 (19)	3853 (15)	302 428 (19)
Missing	746 (4)	261 (5)	982 (4)	68 888 (4)
Maternal education at childbirth, years				
0-9	4178 (19)	2048 (31)	5697 (22)	637 947 (27)
10-14	9194 (42)	2599 (40)	11 601 (44)	1 015 220 (43)
15+	8356 (38)	1708 (26)	7613 (29)	670 697 (28)
Missing	327 (1)	182 (3)	1361 (5)	53 272 (2)
Maternal cohabitation at childbirth				
No	10 079 (46)	2573 (39)	10 935 (42)	1 054 960 (44)
Yes	11 968 (54)	3964 (61)	15 317 (58)	1 318 900 (55)
Missing	8 (<1)	0 (0)	20 (<1)	3276 (<1)
Maternal residence at childbirth				
Copenhagen	3061 (14)	901 (14)	2413 (9)	267 353 (11)
Big cities ≥100 000 inhabitants	2430 (11)	1012 (15)	3514 (13)	303 220 (13)
Others	16 564 (75)	4624 (71)	20 345 (77)	1 806 563 (76)
Maternal CVD history before childbirth				
No	20 648 (94)	6025 (92)	24 847 (95)	2 311 824 (97)
Yes	1407 (6)	512 (8)	1425 (5)	65 312 (3)
Paternal CVD history before birth of the child				
No	20 997 (95)	6 217 (95)	24 827 (94)	2 297 308 (97)
Yes	1043 (5)	303 (5)	1429 (5)	72 482 (3)
Missing	15 (<1)	17 (<1)	16 (<1)	7346 (<1)

^a Expressed as frequency (percentage). Percentages have been rounded and may not total to 100.

^b Maternal smoking during pregnancy was available from 1991 to 2016.

Table S3. Incidence rates of cardiovascular risk factors among offspring according to the presence of maternal diabetes^a

Exposure	Cardiovascular risk factors among offspring									
	Diabetes		Chronic kidney disease		Hypertension		Hypercholesterol emia		Obesity	
	No. of cases	Rate per 1000 person-years	No. of cases	Rate per 1000 person-years	No. of cases	Rate per 1000 person-years	No. of cases	Rate per 1000 person-years	No. of cases	Rate per 1000 person-years
No diabetes	30 653	0.67	3522	0.08	8612	0.19	2395	0.05	69 827	1.53
Maternal diabetes	727	1.27	58	0.10	123	0.21	33	0.06	1138	1.99
Pregestational diabetes	606	1.75	40	0.11	90	0.26	27	0.08	719	2.07
Type 1	401	1.61	30	0.12	53	0.21	20	0.08	420	1.68
Type 2	205	2.10	10	0.10	37	0.37	7	0.07	299	3.08
Gestational diabetes	121	0.53	18	0.08	33	0.15	6	0.03	419	1.86

^a Chronic kidney disease: (ICD-8 codes: 580.00, 582.00-584.99, 593.20; ICD-10 codes: N02.0-02.8, N03.0-03.9, N05.0-6.8, N07.0-8.8, N14.0-14.4, N18.0-18.9, I12.0); hypertension: (ICD-8 codes:400-404, 426; ICD-10 codes:I10-I15, I27); hypercholesterolemia: (ICD-8 codes:27200; ICD-10 codes:E780); obesity: (ICD-8 codes:277; ICD-10 codes:E66).

Table S4. Associations between maternal diabetes and early CVD onset in offspring, by hierarchical ordering of diabetes types (type 1, type 2, and gestational diabetes)

Exposure	No. of CVD cases	Rate per 1000 person-years	HR (95% CI) Model 1	HR (95% CI) Model 2
No diabetes	91 311	2.01	1.0 (ref)	1.0 (ref)
Maternal diabetes	1153	2.02	1.47 (1.39 to 1.56)	1.29 (1.21 to 1.37)
Pregestational diabetes	820	2.29	1.45 (1.35 to 1.56)	1.34 (1.25 to 1.43)
Type 1	589	2.11	1.46 (1.34 to 1.59)	1.31 (1.20 to 1.43)
Type 2	231	2.92	1.44 (1.27 to 1.62)	1.39 (1.23 to 1.57)
Gestational diabetes	333	1.55	1.51 (1.36 to 1.67)	1.19 (1.07 to 1.32)

Abbreviations: CVD, cardiovascular disease; HR, hazard ratio.

Model 1: Offspring' s age as time scale.

Model 2: Offspring' s age as time scale. Controlled for calendar year, sex, singleton status, parity, maternal smoking, maternal education, maternal cohabitation, maternal residence at birth, maternal history of CVD before childbirth, paternal history of CVD before birth of the child, and maternal age (restricted cubic spline with five knots at five evenly spaced).

Table S5. Associations between maternal diabetes and early CVD onset after excluding offspring of mothers diagnosed with multiple types of diabetes in one pregnancy

Exposure	No. of CVD cases	Rate per 1000 person-years	HR (95% CI) Model 1	HR (95% CI) Model 2
No diabetes	91 311	2.01	1.0 (ref)	1.0 (ref)
Maternal diabetes	1005	2.02	1.44 (1.35 to 1.53)	1.27 (1.19 to 1.35)
Pregestational diabetes	672	2.37	1.43 (1.32 to 1.54)	1.34 (1.24 to 1.44)
Type 1	462	2.14	1.43 (1.30 to 1.56)	1.29 (1.18 to 1.42)
Type 2	210	3.08	1.43 (1.25 to 1.64)	1.45 (1.26 to 1.66)
Gestational diabetes	333	1.55	1.47 (1.32 to 1.63)	1.15 (1.04 to 1.29)

Abbreviations: CVD, cardiovascular disease; HR, hazard ratio.

Model 1: Offspring' s age as time scale.

Model 2: Offspring' s age as time scale. Controlled for calendar year, sex, singleton status, parity, maternal smoking, maternal education, maternal cohabitation, maternal residence at birth, maternal history of CVD before childbirth, paternal history of CVD before birth of the child, and maternal age (restricted cubic spline with five knots at five evenly spaced).

Table S6. Associations between maternal diabetes and early CVD onset in term-born offspring^a

Exposure	No. of CVD cases	Rate per 1000 person-years	HR (95% CI) Model 1	HR (95% CI) Model 2
No diabetes	73 360	1.89	1.0 (ref)	1.0 (ref)
Maternal diabetes	869	1.83	1.41 (1.32 to 1.50)	1.23 (1.15 to 1.32)
Pregestational diabetes	557	2.03	1.36 (1.25 to 1.48)	1.26 (1.16 to 1.37)
Type 1	372	1.88	1.39 (1.25 to 1.54)	1.24 (1.12 to 1.37)
Type 2	185	2.40	1.32 (1.14 to 1.52)	1.30 (1.12 to 1.50)
Gestational diabetes	312	1.56	1.49 (1.33 to 1.67)	1.19 (1.07 to 1.33)

Abbreviations: CVD, cardiovascular disease; HR, hazard ratio.

Model 1: Offspring' s age as time scale.

Model 2: Offspring' s age as time scale. Controlled for calendar year, sex, singleton status, parity, maternal smoking, maternal education, maternal cohabitation, maternal residence at birth, maternal history of CVD before childbirth, paternal history of CVD before birth of the child, and maternal age (restricted cubic spline with five knots at five evenly spaced).

^a Term-born offspring: live-births with a gestational age \geq 37 weeks.

Table S7. Associations between maternal diabetes (type 1 and type 2) and early CVD onset in offspring, according to the timing of the maternal diabetes diagnosis

Timing of maternal diabetes diagnosis	No. of CVD cases	Rate per 1000 person-years	HR (95% CI) Model 1	HR (95% CI) Model 2
Maternal type 1 diabetes diagnosis				
No diabetes diagnosed	87 826	1.99	1.0 (ref)	1.0 (ref)
Diagnosed before birth	462	2.10	1.53 (1.40 to 1.68)	1.35 (1.23 to 1.48)
Diagnosed ≤ 2 years after birth	82	1.70	1.36 (1.10 to 1.69)	1.19 (0.96 to 1.48)
Diagnosed 2-5 years after birth	203	1.92	1.42 (1.23 to 1.63)	1.28 (1.11 to 1.46)
Diagnosed > 5 years after birth	3269	2.45	1.06 (1.03 to 1.10)	1.10 (1.06 to 1.13)
Maternal type 1 and type 2 diabetes diagnosis				
No diabetes diagnosed	82 464	1.96	1.0 (ref)	1.0 (ref)
Diagnosed before birth	766	2.28	1.45 (1.35 to 1.55)	1.34 (1.25 to 1.44)
Diagnosed ≤ 2 years after birth	152	2.08	1.35 (1.15 to 1.58)	1.27 (1.09 to 1.49)
Diagnosed 2-5 years after birth	277	1.97	1.34 (1.19 to 1.51)	1.24 (1.10 to 1.39)
Diagnosed > 5 years after birth	8444	2.65	1.11 (1.08 to 1.13)	1.14 (1.12 to 1.17)

Abbreviations: CVD, cardiovascular disease; HR, hazard ratio.

Model 1: Offspring' s age as time scale.

Model 2: Offspring' s age as time scale. Controlled for calendar year, sex, singleton status, parity, maternal smoking, maternal education, maternal cohabitation, maternal residence at birth, maternal history of CVD before childbirth, paternal history of CVD before birth of the child, and maternal age (restricted cubic spline with five knots at five evenly spaced).

Table S8. Associations between maternal diabetes and early CVD onset in offspring, with additional adjustment for parental factors^a

Additional covariate	Exposure	HR (95% CI)^b
Maternal country of origin	No diabetes	1.0 (ref)
	Maternal diabetes	1.29 (1.22 to 1.37)
	Pregestational diabetes	1.34 (1.25 to 1.43)
	Type 1	1.31 (1.20 to 1.43)
	Type 2	1.39 (1.24 to 1.58)
	Gestational diabetes	1.20 (1.08 to 1.33)
Paternal diabetes before birth of the child	No diabetes	1.0 (ref)
	Maternal diabetes	1.29 (1.21 to 1.36)
	Pregestational diabetes	1.34 (1.24 to 1.43)
	Type 1	1.31 (1.20 to 1.43)
	Type 2	1.39 (1.23 to 1.57)
	Gestational diabetes	1.19 (1.07 to 1.32)
Maternal pre-pregnancy BMI^c	No diabetes	1.0 (ref)
	Maternal diabetes	1.26 (1.13 to 1.40)
	Pregestational diabetes	1.35 (1.17 to 1.57)
	Type 1	1.31 (1.11 to 1.56)
	Type 2	1.52 (1.11 to 2.08)
	Gestational diabetes	1.17 (1.00 to 1.37)
All three variables above	No diabetes	1.0 (ref)
	Maternal diabetes	1.26 (1.13 to 1.41)
	Pregestational diabetes	1.36 (1.17 to 1.57)
	Type 1	1.31 (1.11 to 1.55)
	Type 2	1.53 (1.11 to 2.09)
	Gestational diabetes	1.18 (1.01 to 1.38)

Abbreviations: CVD, cardiovascular disease; HR, hazard ratio.

^a As data on pre-pregnancy BMI was available only from 2004 to 2016, an analysis with additional adjustment for maternal pre-pregnancy BMI was restricted to offspring born during 2004-2016. Other analyses included offspring born during 1977-2016.

^b Offspring' s age as time scale. Controlled for calendar year, sex, singleton status, parity, maternal smoking, maternal education, maternal cohabitation, maternal residence at birth, maternal history of CVD before childbirth, paternal history of CVD before birth of the child, and maternal age (restricted cubic spline with five knots at five evenly spaced).

^c Pre-pregnancy BMI was available from 2004 to 2016 and was included in the model using restricted cubic spline with five knots at five evenly spaced).

Table S9. Associations between paternal diabetes and early CVD onset in offspring

Exposure	No. of CVD cases	Rate per 1000 person-years	HR (95% CI) Model 1	HR (95% CI) Model 2
No paternal diabetes	92 025	2.01	1.0 (ref)	1.0 (ref)
Paternal diabetes	439	1.86	1.21 (1.10 to 1.32)	1.10 (1.00 to 1.20)
Type 1	321	1.95	1.20 (1.08 to 1.34)	1.12 (1.01 to 1.25)
Type 2	118	1.67	1.22 (1.02 to 1.46)	1.03 (0.86 to 1.23)

Abbreviations: CVD, cardiovascular disease; HR, hazard ratio.

Model 1: Offspring' s age as time scale.

Model 2: Offspring' s age as time scale. Controlled for calendar year, sex, singleton status, parity, maternal smoking, maternal education, maternal cohabitation, maternal residence at birth, maternal history of CVD before childbirth, paternal history of CVD before birth of the child, and maternal age (restricted cubic spline with five knots at five evenly spaced).

Table S10. Sensitivity analysis of live-birth bias using inverse probability weighting

Exposure	No. of CVD cases	Rate per 1000 person-years	HR (95% CI) Model 1	HR (95% CI) Model 2
No diabetes	91 311	2.01	1.0 (ref)	1.0 (ref)
Maternal diabetes	1153	2.02	1.47 (1.39 to 1.56)	1.29 (1.21 to 1.38)
Pregestational diabetes	792	2.28	1.45 (1.35 to 1.56)	1.35 (1.25 to 1.45)
Type 1	531	2.13	1.46 (1.34 to 1.59)	1.32 (1.21 to 1.45)
Type 2	261	2.68	1.44 (1.27 to 1.62)	1.41 (1.23 to 1.62)
Gestational diabetes	361	1.60	1.51 (1.36 to 1.67)	1.18 (1.05 to 1.32)

Abbreviations: CVD, cardiovascular disease; HR, hazard ratio.

Model 1: Offspring' s age as time scale.

Model 2: Offspring' s age as time scale. Controlled for calendar year, sex, singleton status, parity, maternal smoking, maternal education, maternal cohabitation, maternal residence at birth, maternal history of CVD before childbirth, paternal history of CVD before birth of the child, and maternal age (restricted cubic spline with five knots at five evenly spaced).

Table S11. Subanalyses of the association between maternal diabetes and early CVD onset in offspring

Sub-analysis	Exposure	No. Of CVD cases	Rate per 1000 person-years	HR (95% CI)
Male offspring	No diabetes	43 631	1.87	1.0 (ref)
	Maternal diabetes	595	2.02	1.27 (1.17 to 1.38)
	Pregestational diabetes	395	2.23	1.32 (1.20 to 1.46)
	Type 1	266	2.09	1.29 (1.15 to 1.46)
	Type 2	129	2.57	1.39 (1.17 to 1.65)
	Gestational diabetes	200	1.71	1.18 (1.03 to 1.36)
Female offspring	No diabetes	47 680	2.15	1.0 (ref)
	Maternal diabetes	558	2.01	1.30 (1.20 to 1.42)
	Pregestational diabetes	397	2.34	1.35 (1.22 to 1.49)
	Type 1	265	2.16	1.33 (1.17 to 1.50)
	Type 2	132	2.81	1.40 (1.18 to 1.66)
	Gestational diabetes	161	1.49	1.19 (1.02 to 1.39)
Singleton offspring	No diabetes	89 037	2.01	1.0 (ref)
	Maternal diabetes	1112	2.04	1.30 (1.23 to 1.38)
	Pregestational diabetes	770	2.32	1.35 (1.26 to 1.45)
	Type 1	517	2.16	1.33 (1.22 to 1.45)
	Type 2	253	2.71	1.40 (1.24 to 1.59)
	Gestational diabetes	342	1.61	1.20 (1.08 to 1.33)
Primiparous women	No diabetes	40 980	2.00	1.0 (ref)
	Maternal diabetes	461	2.08	1.31 (1.19 to 1.44)
	Pregestational diabetes	313	2.33	1.35 (1.21 to 1.51)
	Type 1	222	2.15	1.29 (1.13 to 1.48)
	Type 2	91	2.91	1.53 (1.24 to 1.87)
	Gestational diabetes	148	1.70	1.22 (1.04 to 1.44)
Offspring born after 1991	No diabetes	28 523	1.37	1.0 (ref)
	Maternal diabetes	730	1.66	1.28 (1.19 to 1.38)
	Pregestational diabetes	405	1.77	1.36 (1.23 to 1.50)
	Type 1	322	1.77	1.38 (1.24 to 1.54)
	Type 2	83	1.76	1.30 (1.05 to 1.61)
	Gestational diabetes	325	1.54	1.19 (1.07 to 1.33)
Offspring born after 1994	No diabetes	20 600	1.27	1.0 (ref)
	Maternal diabetes	660	1.61	1.27 (1.18 to 1.38)
	Pregestational diabetes	360	1.72	1.36 (1.23 to 1.51)
	Type 1	290	1.71	1.36 (1.21 to 1.53)
	Type 2	70	1.76	1.37 (1.08 to 1.73)
	Gestational diabetes	300	1.50	1.18 (1.06 to 1.33)
Offspring born after 2004	No diabetes	6118	1.24	1.0 (ref)
	Maternal diabetes	347	1.64	1.26 (1.13 to 1.41)
	Pregestational diabetes	178	1.77	1.36 (1.17 to 1.58)
	Type 1	139	1.71	1.31 (1.11 to 1.56)
	Type 2	39	2.01	1.52 (1.11 to 2.09)
	Gestational diabetes	169	1.53	1.17 (1.01 to 1.37)
Complete case analysis	No diabetes	26 607	1.36	1.0 (ref)

	Maternal diabetes	669	1.64	1.27 (1.17 to 1.37)
	Pregestational diabetes	372	1.74	1.34 (1.21 to 1.49)
	Type 1	298	1.74	1.36 (1.21 to 1.53)
	Type 2	74	1.72	1.27 (1.01 to 1.60)
	Gestational diabetes	297	1.52	1.19 (1.06 to 1.33)
Missing indicator method	No diabetes	91 311	2.01	1.0 (ref)
	Maternal diabetes	1153	2.02	1.29 (1.21 to 1.36)
	Pregestational diabetes	792	2.28	1.33 (1.24 to 1.43)
	Type 1	531	2.13	1.31 (1.20 to 1.43)
	Type 2	261	2.68	1.39 (1.23 to 1.57)
	Gestational diabetes	361	1.60	1.19 (1.07 to 1.32)
Additional adjustment for LGA^a, preterm birth and pre-eclampsia	No diabetes	91 311	2.01	1.0 (ref)
	Maternal diabetes	1153	2.02	1.24 (1.16 to 1.31)
	Pregestational diabetes	792	2.28	1.27 (1.18 to 1.36)
	Type 1	531	2.13	1.24 (1.14 to 1.36)
	Type 2	261	2.68	1.33 (1.18 to 1.50)
	Gestational diabetes	361	1.60	1.16 (1.05 to 1.29)

Abbreviations: CVD, cardiovascular disease; HR, hazard ratio.

Offspring' s age as time scale. Controlled for calendar year, sex, singleton status, parity, maternal smoking, maternal education, maternal cohabitation, maternal residence at birth, maternal history of CVD before childbirth, paternal history of CVD before birth of the child, and maternal age (restricted cubic spline with five knots at five evenly spaced).

^a LGA: large for gestational age (birthweight below the 10th, between 10th and 90th, above the 90th percentile for infants of the same gestational age, sex, and birth year).

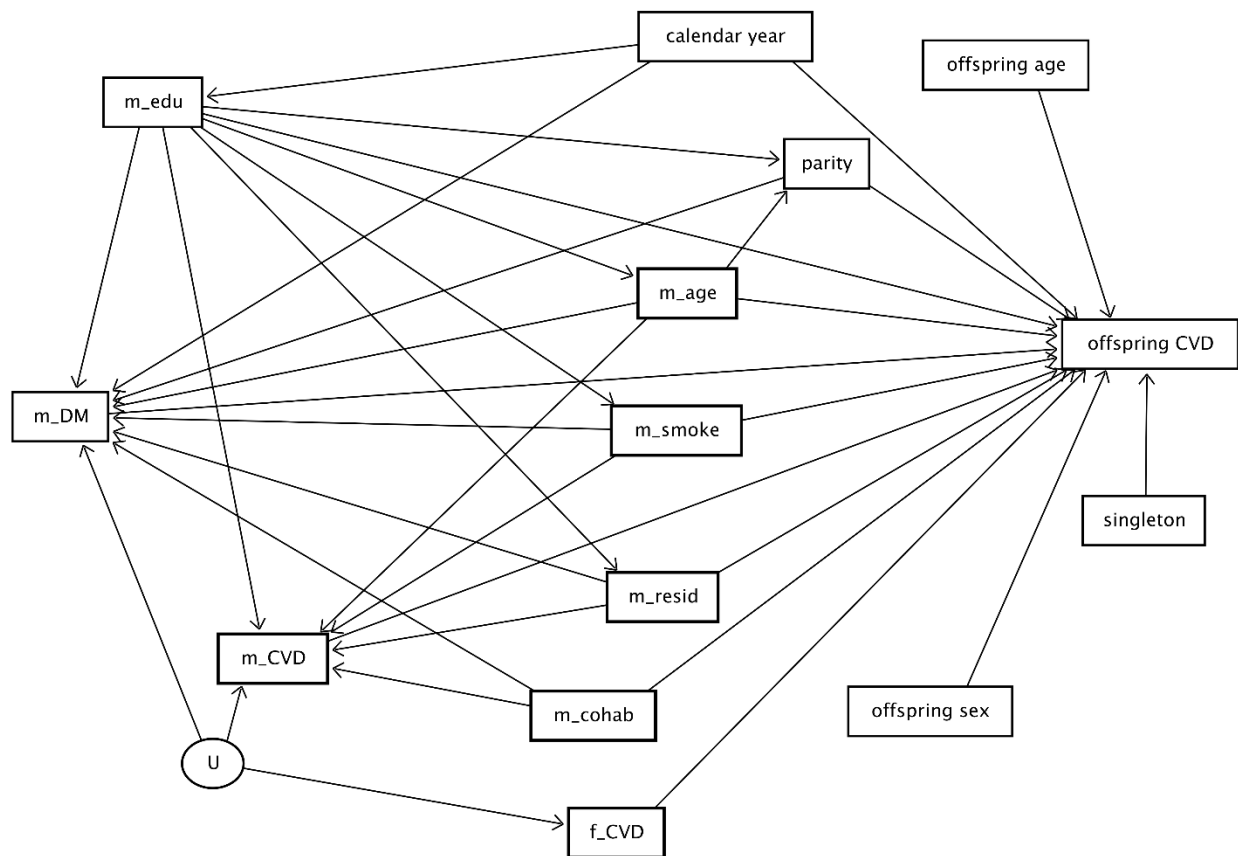


Figure S1. Causal diagram showing selection of covariates for confounding control^a

^a m_DM: maternal diabetes, m_age: maternal maternal age, m_smoke: maternal smoking, m_edu: maternal education, m_cohab: maternal cohabitation, m_resid: maternal residence at birth, m_CVD: maternal history of CVD before childbirth, f_CVD: paternal history of CVD before birth of the child, U: unmeasured variable

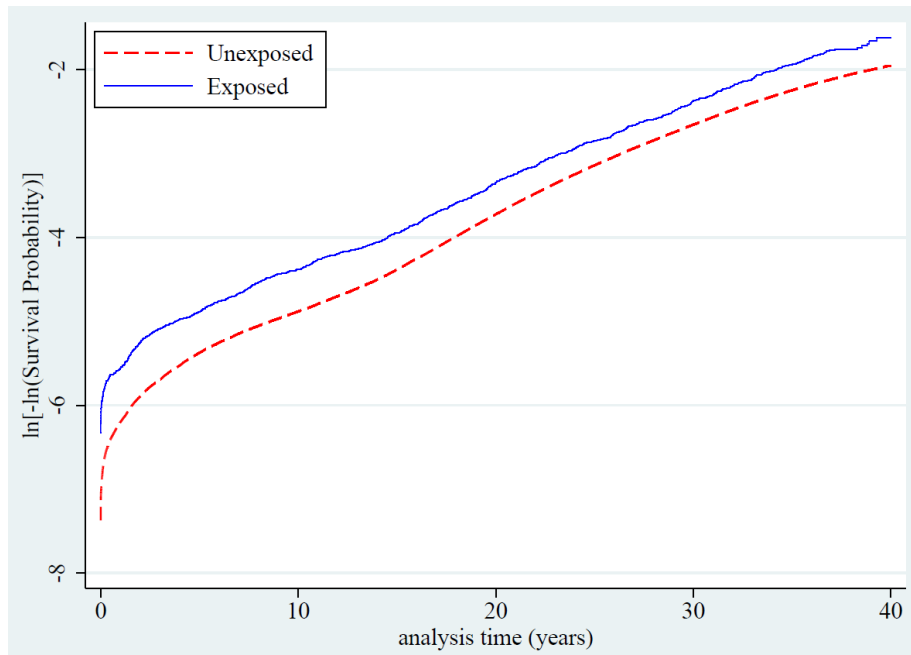


Figure S2. The log-minus-log survival curve

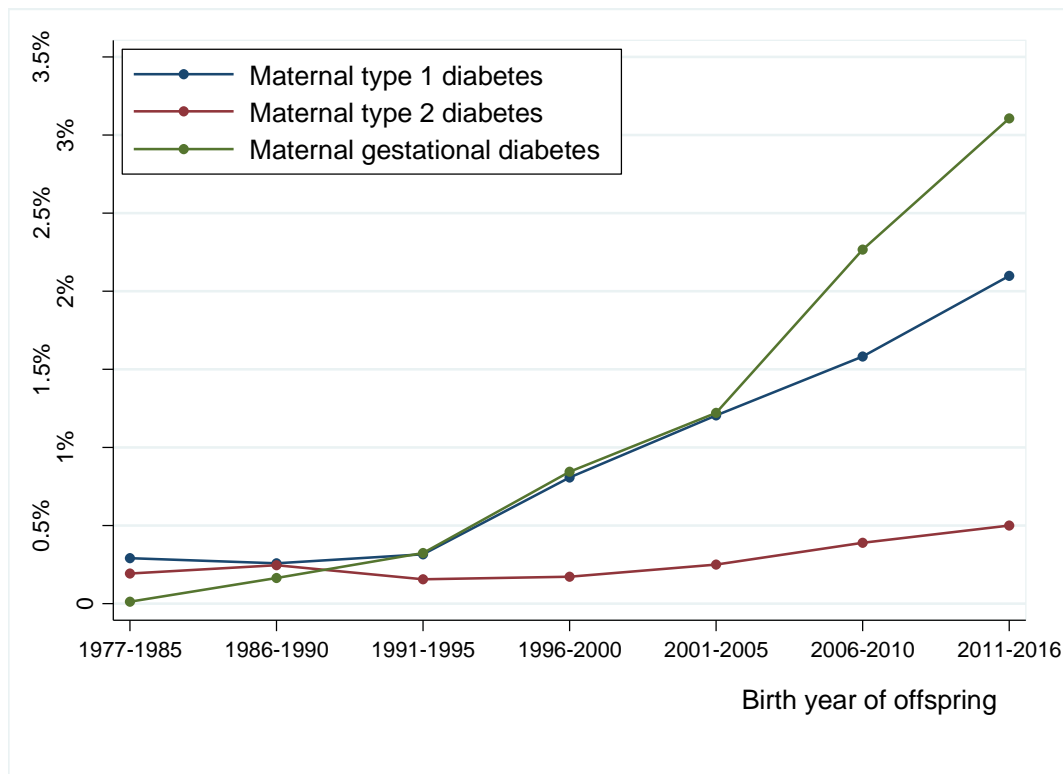
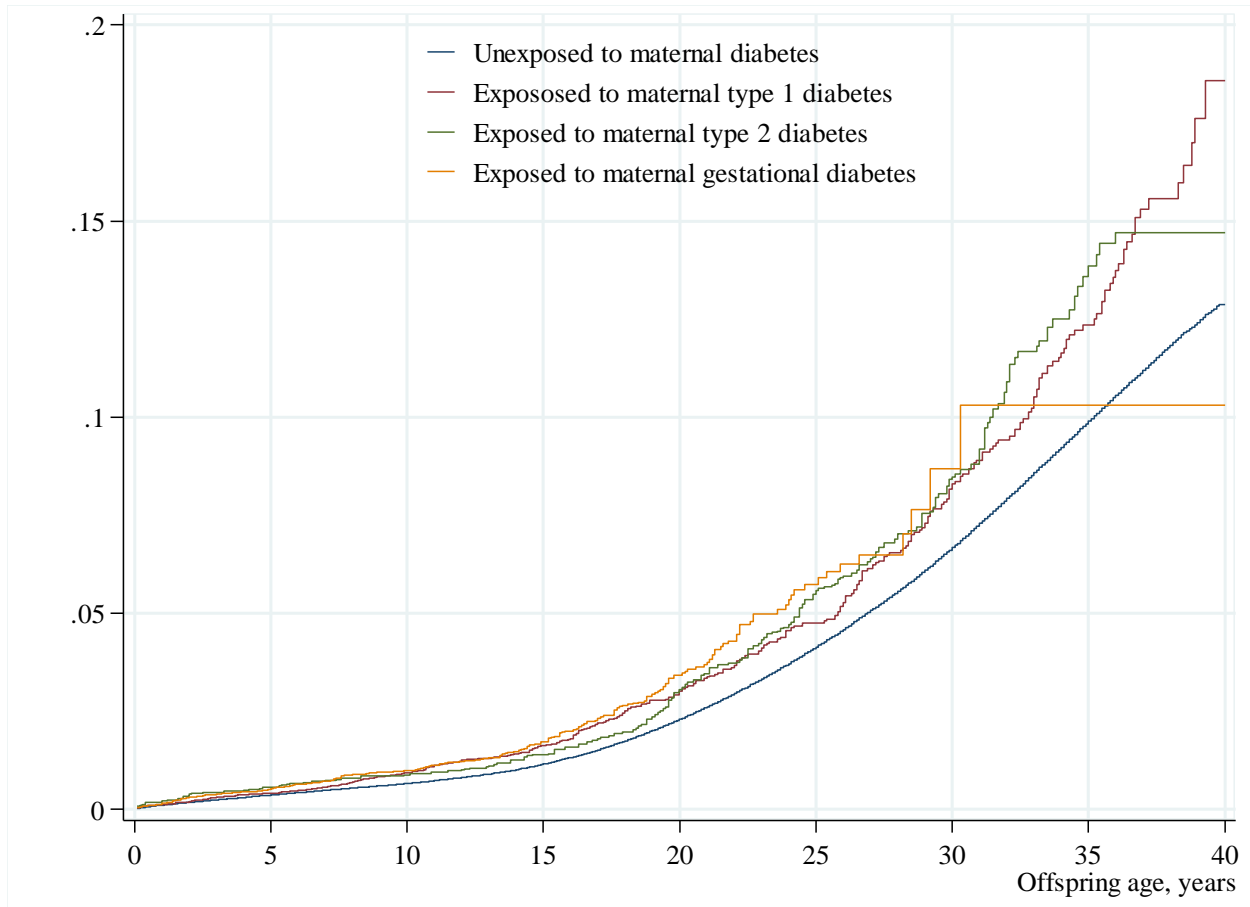


Figure S3. The proportion of offspring born to mothers with diabetes, by birth years and types of maternal diabetes



Number at risk									
Unexposed	2 377 136	2 062 027	1 759 037	1 450 866	1 127 219	787 639	490 892	246 601	0
Type 1	22 055	15 437	10 295	6323	3408	2160	1408	628	0
Type 2	6537	4924	3616	2769	2144	1606	927	360	0
GDM	26 272	16 598	9050	4724	2039	603	59	10	0

Figure S4. Adjusted cumulative incidence of early CVD onset among offspring, exposed to maternal diabetes versus unexposed^a

^a Adjusted cumulative incidence averaged across the distribution of the covariates (calendar year, sex, singleton status, parity, age, smoking, education, cohabitation, residence at childbirth, history of CVD before childbirth, and paternal history of CVD before childbirth) using the inverse probability of treatment weighting approach. aternal diabetes.