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Smoking during pregnancy and diabetes mellitus in a British longitudinal birth cohort
Scott M Montgomery, Anders Ekbom

Exposures in utero may increase the risk of type 2 diabetes. We tested the hypothesis that maternal smoking during pregnancy increases both the risk of early onset type 2 diabetes and non-diabetic obesity in offspring.

Methods and results
We used data from the British National Child Development Study (NCDS), based on the Perinatal Mortality Survey (PMS) of about 17,000 births from 3 to 9 March 1958. The first sweep of the study in 1965 had 15,996 responses. The cohort remained generally representative at age 33 years (n = 11,559). Missing data reduced the proportion in social class V from 6.4% in sweep 1 to 5.3%. Ethics committee approval was obtained for research involving medical examinations, and cohort members signed consent forms at age 33 years allowing access to medical records.

Medical examinations and record reviews by local authority medical officers were conducted at ages 7 and 16 years. Children with incomplete or equivocal information on diabetes or with a recorded onset before the age of 16 were not included in the main analysis as they are unlikely to have type 2 diabetes. A personal interview at age 33 years asked about diabetes. Those with only gestational diabetes were also excluded: 15 men and 13 women with an onset of diabetes. We tested the hypothesis that maternal smoking during pregnancy increases both the risk of early onset type 2 diabetes and non-diabetic obesity in offspring.

At birth midwives recorded information on the child’s sex, birth weight, mother’s age, her age on leaving full-time education, family social class, and smoking during pregnancy (after the 4th month) divided into non-smokers, medium (1–9 cigarettes/day) and heavy (>10), and variable (a balance of medium and heavy). Details of maternal smoking were again recorded in 1974, as non-smoking and <1, 1–5, 6–10, 11–20, 21–30, or >30 cigarettes/day.

Cohort members’ own smoking behaviour was recorded during an interview at age 16 and they were classified as non-smokers or as smoking <1, 1–9, 10–19, 20–29, >29 cigarettes/week. Interviewers measured height in centimetres and weight in kilograms using stadiometers and electronic balances at age 33. Multiple logistic regression analysis was used for two outcomes: diabetes and body mass index (BMI) of over 30, independent of diabetes. Where obesity was the outcome, those with diabetes were excluded and we adjusted for sex, own smoking at age 16, and all the maternal factors at birth.

Some 10% (n = 602) were obese (BMI > 30) at age 33. After we excluded the diabetic cohort members the adjusted odds ratios (and 95% confidence intervals) for obesity associated with maternal smoking during pregnancy are 1.34 (1.07 to 1.69), 1.35 (0.95 to 1.92), and 1.38 (1.06 to 1.79), with a statistically significant trend (P = 0.003) for medium, variable, and heavy smokers, respectively (table). Non-diabetic cohort members who smoked at age 16 did not have an increased risk of obesity.

Comment
The association of diabetes with maternal smoking during pregnancy (independent of finer-grain measures of mothers’ smoking in 1974, own smoking at age 16, and other potential confounding factors) suggests that it is a true risk factor for early adult onset diabetes.

Cigarette smoking as a young adult was also independently associated with an increased risk of subsequent diabetes.

Maternal smoking in pregnancy, own smoking at age 16 years, and risk of diabetes among young adults. Odds ratio with 95% confidence intervals using logistic regression with a diagnosis of diabetes after age 16 as the dependent variable

<table>
<thead>
<tr>
<th>Smoking during pregnancy</th>
<th>People without diabetes No (%)</th>
<th>People with diabetes No (%)</th>
<th>Unadjusted OR (95% CI) P</th>
<th>Adjusted* OR (95% CI) P</th>
<th>Adjusted† OR (95% CI) P</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mother smoking while pregnant (after 4th month)</td>
<td>Non-smoker 3430 (69.8) 12 (2.9) 1.00 1.00 1.00</td>
<td>Medium smoker 698 (14.0) 3 (10.7) 1.24 (0.35 to 4.42) 0.735 1.11 (0.31 to 4.04) 0.872 1.01 (0.23 to 4.53) 0.990</td>
<td>Varies between 277 (5.6) 4 (14.3) 4.13 (1.32 to 12.88) 0.015 4.13 (1.27 to 13.40) 0.018 3.55 (0.88 to 14.38) 0.076</td>
<td>Heavy smoker 521 (10.6) 9 (32.1) 4.94 (2.07 to 11.77) &lt;0.001 4.55 (1.82 to 11.38) 0.001 4.02 (1.14 to 14.14) 0.030</td>
<td></td>
</tr>
<tr>
<td>Own smoking at age 16 years (cigarettes/week)</td>
<td>Non-smoker 3251 (66.1) 13 (46.4) 1.00 1.00 1.00</td>
<td>1–5 147 (3.0) 3 (35.6) 1.20 (0.22 to 13.09) 0.610 2.09 (0.25 to 17.23) 0.492 2.07 (0.25 to 17.19) 0.502</td>
<td>1–9 510 (10.4) 3 (64.4) 1.47 (0.42 to 5.18) 0.548 1.76 (0.48 to 6.46) 0.394 1.92 (0.52 to 7.10) 0.332</td>
<td>10–19 176 (3.6) 2 (17.6) 2.84 (0.64 to 12.69) 0.171 2.41 (0.51 to 11.41) 0.268 2.48 (0.52 to 11.97) 0.257</td>
<td>20–29 189 (3.8) 3 (36.7) 1.32 (0.17 to 10.17) 0.788 1.42 (0.17 to 11.43) 0.741 1.61 (0.20 to 12.96) 0.653</td>
</tr>
</tbody>
</table>

*Adjusted for maternal smoking during pregnancy, sex, mother’s age at birth of cohort member, age mother left school, family social class at birth, birth weight, own smoking at age 16 years, and BMI at age 33 years.
†Adjusted for all of the above and maternal smoking in 1974.
In utero exposures due to smoking during pregnancy may increase the risk of both diabetes and obesity through programming, resulting in lifelong metabolic dysregulation, possibly due to fetal malnutrition or toxicity. The odds ratios for obesity without type 2 diabetes are more modest than those for diabetes and the scope for confounding may be greater. Smoking during pregnancy may represent another important determinant of metabolic dysregulation and type 2 diabetes in offspring. Smoking during pregnancy should always be strongly discouraged.

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Drug points

Ticlopidine associated with acute arthritis

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Ticlopidine is an antiplatelet thienopyridine drug that works by non-competitive antagonism of the ADP receptor. It is used widely to prevent thrombosis after coronary stent placement and has been shown to be at least as effective as aspirin in preventing events in patients with cerebrovascular disease.\(^1\) Its most common side effects include diarrhoea, nausea, vomiting, and skin rash.\(^3\) It also has serious—but rare—side effects such as neutropenia, thrombotic thrombocytopenic purpura, and bone marrow aplasia.\(^4\) We report a case of acute arthritis associated with ticlopidine.

A 65 year old woman was admitted to hospital because of chest pain. She was known to be taking insulin for diabetes and had developed non-Q wave myocardial infarction. She was also hyperlipidaemic. Cardiac catheterisation showed two vessel coronary artery disease, for which she had successful angioplasty with stent placement. Sanofi-Synthelabo, the manufacturer of ticlopidine, has not reported any case of polyarthritis associated with ticlopidine implantation. Ticlopidine, has not reported any case of polyarthritis associated with ticlopidine intake. Treatment with ticlopidine was discontinued, and one week later her rash resolved completely but her arthritis persisted. After treatment with a non-steroidal anti-inflammatory drug (diclofenac 75 mg intramuscularly twice daily) for 10 days, her arthritis gradually resolved. Two weeks later her joints were completely normal and her erythrocyte sedimentation rate decreased to 32 mm/h. On her last evaluation, six months after the onset of arthritis, she had had no recurrence of her joint pain, and her erythrocyte sedimentation rate had dropped to 18 mm/h. The clinical features and laboratory findings of arthritis in this case suggest a drug induced hypersensitivity (leucocytoclastic) vasculitis. Before March 2001, one case of polyarthritis and three cases of arthralgia associated with ticlopidine had been reported to the Committee on Safety of Medicines in the United Kingdom. Two case reports of arthritis associated with clopidogrel have also been published.\(^4\) Clopidogrel is a thienopyridine drug with a similar chemical structure to ticlopidine and is commonly used in patients undergoing coronary stent implantation. Sanofi-Synthelabo, the manufacturer of ticlopidine, has not reported any case of polyarthritis associated with taking the drug. We suggest that thienopyridine derivatives be considered as a potential cause of acute arthritis.

The presumptive diagnosis was a symmetrical polyarthritis associated with ticlopidine intake. Treatment with ticlopidine was discontinued, and one week later her rash resolved completely but her arthritis persisted. After treatment with a non-steroidal anti-inflammatory drug (diclofenac 75 mg intramuscularly twice daily) for 10 days, her arthritis gradually resolved. Two weeks later her joints were completely normal and her erythrocyte sedimentation rate decreased to 32 mm/h. On her last evaluation, six months after the onset of arthritis, she had had no recurrence of her joint pain, and her erythrocyte sedimentation rate had dropped to 18 mm/h. The clinical features and laboratory findings of arthritis in this case suggest a drug induced hypersensitivity (leucocytoclastic) vasculitis. Before March 2001, one case of polyarthritis and three cases of arthralgia associated with ticlopidine had been reported to the Committee on Safety of Medicines in the United Kingdom. Two case reports of arthritis associated with clopidogrel have also been published.\(^4\) Clopidogrel is a thienopyridine drug with a similar chemical structure to ticlopidine and is commonly used in patients undergoing coronary stent implantation. Sanofi-Synthelabo, the manufacturer of ticlopidine, has not reported any case of polyarthritis associated with taking the drug. We suggest that thienopyridine derivatives be considered as a potential cause of acute arthritis.

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