Effect of intrauterine diabetes exposure on the incidence of end-stage renal disease in young adults with type 2 diabetes

Running title: Intrauterine exposure to diabetes

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Objective. We examined the effect of intrauterine diabetes exposure (IDE) on the incidence of diabetic end-stage renal disease (ESRD) in Pima Indians with type 2 diabetes.

Research Design and Methods. Individuals were followed from their first diabetic examination until December 2006, death, ESRD, or age of 45 years.

Results. Among the 1,850 diabetic participants, 102 had IDE. ESRD developed in 57, five of whom had IDE. Cumulative incidence of ESRD by age 45 was 19.3% in participants with IDE and 5.1% in those without; age-sex-adjusted incidence rate ratio was 4.12, (95% CI=1.54-11.02). After additional adjustment for age at diabetes onset, ESRD incidence was similar in the two groups (incidence rate ratio=1.38, 95% CI=0.45-4.24).

Conclusions. IDE increases the age-sex-adjusted incidence of ESRD 4-fold in young adults with type 2 diabetes, mediated primarily by the earlier onset of type 2 diabetes in those with IDE.

The intrauterine environment plays a central role in fetal development and an adverse environment may enhance the risk of chronic diseases in the offspring [1-5]. We examined the effect of intrauterine diabetes exposure (IDE) on incidence of diabetic end-stage renal disease (ESRD) in Pima Indians with type 2 diabetes.

RESEARCH DESIGN AND METHODS

Between 1965 and 2007, Pima Indians from the Gila River Indian Community participated in a longitudinal diabetes study. Each member of the community ≥5 years old was invited to have a research examination every 2 years that included a 75-gram oral glucose tolerance test (OGTT). In addition, many pregnant women also had a 75-gram OGTT in the third trimester. Diabetes was defined by a 2-hour post-load plasma glucose concentration ≥11.1 mmol/l (200 mg/dl), or by a clinical diagnosis between examinations. IDE was considered present if the mother was diagnosed with diabetes before the child's birth.

Diabetic ESRD was defined as the initiation of renal replacement therapy or death from diabetic nephropathy, and was ascertained independently of the research examinations. Cause of ESRD in those receiving renal replacement therapy was determined from clinical records. Death was attributed to diabetic nephropathy if ICD-9 code 250.4 was specified as the underlying cause [6]. The study population included diabetic subjects aged 5 to 45 years old who attended research examinations and resided in the community. The longitudinal study was approved by the review board of the National Institute of Diabetes and Digestive and Kidney Diseases. Each participant gave informed consent or assent.

Statistical analysis. Clinical features at baseline were compared by IDE with analysis of covariance for continuous variables or Mantel-Haenszel Chi-square for categorical variables stratified by sex. Variables with non-normal distributions were analyzed with the Kruskal–Wallis test.

Incidence of ESRD was computed and events and person-years (pyrs) of follow-up were stratified in a time-dependent fashion by decades of age. Individuals were followed from their first diabetic examination after January 1, 1965 until December 31, 2006, death, onset of ESRD, or age of 45 years, whichever came first. Individuals above 45 years of age were not included, because few
people whose IDE could be determined were followed past 45 years of age. Unadjusted cumulative incidence of ESRD in diabetic subjects, stratified by IDE, was estimated by the Kaplan-Meier product-limit method either as a function of age or duration of diabetes. When calculated by age, subjects entered at the diagnosis of diabetes, and the Kaplan-Meier method was modified for left-truncated data [7]. Differences in cumulative incidence were assessed by the log-rank test. Age-sex-adjusted incidence rates of ESRD were standardized to the 1985 Pima Indian population and rate ratios were computed from these age-sex-adjusted rates. Incidence rate ratios adjusted for age, sex, and age at onset of diabetes were computed by Mantel-Haenszel stratification. Proportion of incident ESRD due to IDE in the diabetic population was computed as the difference in age-sex-adjusted incidence of ESRD in the total diabetic population and in the diabetic population without IDE divided by the age-sex-adjusted incidence in the total diabetic population.

RESULTS

Among the 1,850 diabetic subjects, 102 (5.5%) were offspring of diabetic mothers. Median follow-up from the first diabetic examination was 7.1 years (interquartile range 3.2 to 12.5 years). Those exposed to diabetes in utero were significantly younger at baseline than the unexposed (17.5 vs. 34.2 years; p <0.0001), owing to the earlier onset of diabetes characteristic of this exposure, and had lower BMI, blood pressure, and serum cholesterol concentration. At the end of follow-up, median age of the unexposed participants remained significantly greater than in those with IDE (26.7 vs. 45.0 years; p<0.0001), but the median duration of diabetes was similar (7.6 vs. 8.8 years; p=0.22).

ESRD developed in 57 participants, five of whom were exposed to diabetes in utero. Cumulative incidence of ESRD by age was higher in participants with IDE than in those without (p=0.001), but was equivalent in the two groups when examined by diabetes duration (p=0.79; Figure). Unadjusted incidence was 5.61 cases/1,000 pyrs in participants with IDE and 3.52 cases/1,000 pyrs in those without. Age-sex-adjusted incidence in those with IDE was 4.12 (95% CI=1.54-11.02) times that of the unexposed. After additional adjustment for age at diabetes onset, ESRD incidence was similar in the two groups (incidence rate ratio=1.38, 95% CI=0.45-4.24). The proportion of diabetic ESRD attributable to IDE in the diabetic population under 45 years old was 18.9%.

CONCLUSIONS

Exposure to IDE was associated with a 4-fold increase in the incidence of ESRD in young adults with type 2 diabetes when adjusted for age and sex. This effect was explained largely by their earlier age at onset of diabetes. To our knowledge, this study is the first to link IDE to an earlier onset of ESRD.

Recent improvements in diabetes management has reduced the overall incidence of diabetic ESRD in some populations [8, 9]. Less effective therapeutic management of youth and young adults with type 2 diabetes [10], however, may limit these beneficial trends in younger people [8]. In the present study, estimates of the population attributable risk of ESRD ascribed to IDE suggest that interventions which delay diabetes onset in women until after the childbearing years could ultimately eliminate 18.9% of the ESRD that would occur before the age of 45 years in the children of these women.

IDE exposure increased nearly 4-fold in Pima Indians between 1967 and 1996 and the prevalence of diabetes in youth related to this exposure doubled during the same period [11]. Nevertheless, because type 2 diabetes in
youth is a recent phenomenon, even in the Pima Indians, the number of cases of ESRD attributable to IDE in the present study was small. The effect of this exposure, however, was large and trends with respect to diabetes and its complications in this population have been a reliable harbinger for trends in other populations. Implementation of appropriate behavioral interventions to prevent or delay diabetes in women of childbearing age may be an effective long-term strategy to reduce the increasing incidence of diabetic ESRD in young adults.

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**Disclaimer:** The findings and conclusions in this paper have not been formally disseminated by the Centers for Disease Control and Prevention and should not be construed to represent any agency determination or policy.

**Figure Legend**

**Figure.** Cumulative incidence of diabetic ESRD by age (left) and by duration of diabetes (right) according to exposure to diabetes in utero (—— Exposed, ----- Unexposed). The five cases of ESRD in the exposed group result in five steps in the cumulative incidence plot by age. Only four cases are shown in the cumulative incidence plot by duration, because the fifth case occurred at a duration beyond the range of the figure.

**REFERENCES**