SHORT REPORT

Progression from islet autoimmunity to clinical type 1 diabetes is influenced by genetic factors: results from the prospective TEDDY study

Andreas Beyerlein,1* Eziel Bonifacio,2,3 Kendra Vehik,4 Markus Hippich,1 Christiane Winkler,1,3 Brigitte I Frohnert,5 Andrea K Steck,5 William A Hagopian,6 Jeffrey P Krischer,4 Åke Lernmark,7 Marian J Rewers,5 Jin-Xiong She,8 Jorma Toppali,9,10 Beena Akolkar,11 Stephen S Rich,12 Anette-G Ziegler,1,3 the TEDDY Study Group

ABSTRACT

Background Progression time from islet autoimmunity to clinical type 1 diabetes is highly variable and the extent that genetic factors contribute is unknown.

Methods In 341 islet autoantibody-positive children with the human leucocyte antigen (HLA) DR3/DR4-DQ8 or the HLA DR4-DQ8/D4-DQ8 genotype from the prospective TEDDY (The Environmental Determinants of Diabetes in the Young) study, we investigated whether a genetic risk score that had previously been shown to predict islet autoimmunity is also associated with disease progression.

Results Islet autoantibody-positive children with a genetic risk score in the lowest quartile had a slower progression from single to multiple autoantibodies (p=0.018), from single autoantibodies to diabetes (p=0.004), and by trend from multiple islet autoantibodies to diabetes (p=0.06). In a Cox proportional hazards analysis, faster progression was associated with an increased genetic risk score independently of HLA genotype (HR for progression from multiple autoantibodies to type 1 diabetes, 1.27, 95% CI 1.02 to 1.58 per unit increase), an earlier age of islet autoantibody development (HR, 0.68, 95% CI 0.58 to 0.81 per year increase in age) and female sex (HR, 1.94, 95% CI 1.28 to 2.93).

Conclusions Genetic risk scores may be used to identify islet autoantibody-positive children with high-risk HLA genotypes who have a slow rate of progression to subsequent stages of autoimmunity and type 1 diabetes.

INTRODUCTION

Type 1 diabetes begins with a preclinical phase which is defined by the presence of islet autoantibodies. This preclinical phase is variable in duration, with onset of clinical diabetes occurring months to decades after the appearance of islet autoantibodies.1 Features of autoimmunity that include autoantibody titre or specificity, age and sex have been used to stratify the rate of progression to type 1 diabetes.2,3 Genes that confer susceptibility to type 1 diabetes, in particular the human leucocyte antigen (HLA) class II genes, usually exert a stronger effect on the development of autoimmunity than on disease progression,4 and although there are reports of genes that influence the progression to clinical diabetes,5,7,8 the extent to which genetic information may be used to stratify the rate of progression to clinical diabetes in islet autoantibody-positive individuals is unknown. Here, we investigated whether a previously established genetic risk score for islet autoimmunity7 is associated with progression to clinical diabetes in the longitudinal TEDDY (The Environmental Determinants of Diabetes in the Young) study.

RESEARCH DESIGN AND METHODS

TEDDY is an ongoing prospective cohort study that enrolled 8676 children with high-risk type 1 diabetes HLA genotypes between 2004 and 2010 in six clinical research centres located in the USA, Finland, Germany and Sweden.10–12 The families of children with risk of HLA genotypes were invited to participate in the follow-up study in which blood samples were obtained every 3 months for the first 4 years and biannually thereafter for the measurement of islet autoantibodies (glutamic acid decarboxylase antibody, insulin autoantibodies and insulin autoantibodies) by radio-binding assays as previously described.13,14 Samples positive for islet autoantibodies were retested at the second reference laboratory for confirmation. The outcome of islet autoantibody positivity was defined as a positive result at both reference laboratories (confirmed) and by the presence of islet autoantibodies (glutamic acid decarboxylase antibody (GADA), insulin autoantibodies-2 antibody (IAA-2A) or insulin autoantibody (IAA)) on two or more consecutive visits (persistent). The date of seroconversion to islet autoimmunity was defined as the date of drawing the first of two consecutive autoantibody-positive samples. The presence of persistent multiple islet autoantibodies was defined as the date of drawing the first sample when the second persistent and confirmed islet autoantibodies were detected. Children with positive islet autoantibodies that were due to maternal IgG transmission were not considered to be positive for that autoantibody unless the child had a negative sample before the first positive sample or the autoantibody...
The genetic score of each individual was derived from weighted values given to the HLA DR3/DR4-DQ8 or DR4-DQ8/DR4-DQ8 genotype plus a weighted value assigned to each susceptible allele of HLA class I and non-HLA SNPs (online supplementary table 1) and was applied also in this analysis. Written informed consent was obtained for all study participants from a parent or primary caretaker for genetic screening and to participate in the prospective follow-up.

Here, we analysed the data of all 341 children who had developed islet autoantibodies during follow-up and for whom the genetic risk score could be determined. We calculated the Kaplan-Meier curves for progression from (1) any autoantibodies to multiple autoantibodies, (2) any autoantibodies to type 1 diabetes onset and (3) multiple autoantibodies to type 1 diabetes onset in children stratified by quartiles of the genetic risk score (lower quartile: <13.47; upper quartile: >14.88; the two middle quartiles were combined into one group, ie, 13.47–14.88). In order to determine the potential contribution of HLA and non-HLA SNPs and explore potential confounding by other, basically unmodifiable factors, we applied Cox proportional hazards regression analysis for the three progression times with the genetic risk score as the main predictor variable, and with HLA genotype (DR3/DR4-DQ8 or DR4-DQ8/DR4-DQ8), sex, age at onset of the previous event (eg, of any autoantibodies in the model of progression from any autoantibodies to type 1 diabetes) and country of ascertainment as confounder variables. Specifically, the HR of each outcome variable was determined as log(HR) = β_GRS × genetic risk score + β_HLA × HLA genotype (reference: HLA DR4-DQ8/DR4-DQ8) + β_sex × sex (reference: boys) + β_age × age at onset (continuous variable) + β_COUNTRY × country (reference: USA). In these models, we used the genetic risk score without inclusion of the HLA class II genotypes so that the contributions of HLA class II genotype and the remainder of the genes in the risk score could be determined separately. All analyses were performed using R version 3.3.3 (R Foundation for Statistical Computing, Vienna, Austria). Significance was defined by a two-sided significance level of 0.05.

RESULTS
There were 341 children who developed islet autoantibodies at a median age of 2.7 (IQR, 1.5–5.0) years, of whom 141 (41.3%) were female, and 250 had the HLA DR3/DR4-DQ8 and 91 the HLA DR4-DQ8/DR4-DQ8 genotype. The subjects were followed to a median age of 7.9 (IQR, 6.2–9.5) years. During this follow-up period, 214 children (62.8%) developed multiple autoantibodies at a median age of 2.8 (IQR, 1.8–5.1) years, and 107 (31.4%) of the children developed clinical type 1 diabetes at a median age of 5.0 (IQR, 3.0–7.1) years, with 96 children (28.2%) developing both multiple autoantibodies and type 1 diabetes. The median genetic risk score was 14.23 (IQR, 13.47–14.88) in all children and was higher in the children who developed clinical type 1 diabetes (median, 14.36; IQR, 13.73–15.03) as compared with the children who remained single islet autoantibody-positive at last visit (median, 14.01; IQR, 13.15–14.56; p=0.007 from Mann-Whitney U test). The children with a genetic risk score in the lowest quartile progressed more slowly from single to multiple islet autoantibodies (p=0.018), from single autoantibodies to diabetes (p=0.004), and by trend from multiple islet autoantibodies to type 1 diabetes (p=0.06; figure 1) than the children with genetic risk scores in the upper three quartiles. In a Cox proportional hazards analysis, an increased genetic risk score calculated without HLA genotype and an earlier age of islet autoantibody development were consistently associated with a faster progression to subsequent stages of autoimmunity and type 1 diabetes. Girls progressed faster from multiple autoantibodies to type 1 diabetes than boys. The HLA DR3/4-DQ8 genotype and country of ascertainment were not associated with the rate of progression at any stage after the appearance of islet autoantibodies (table 1).

CONCLUSIONS
This study suggests that an islet autoimmunity genetic risk score is predictive of the rate of progression to clinical onset of type 1 diabetes in islet autoantibody-positive children with the HLA DR3/DR4-DQ8 or the HLA DR4-DQ8/DR4-DQ8 genotype.

Figure 1 Cumulative risks of (A) development of multiple islet autoantibodies after first appearance of any autoantibodies, (B) development of type 1 diabetes after first appearance of any autoantibodies and (C) development of type 1 diabetes after first appearance of multiple autoantibodies, in children with the HLA DR3/DR4-DQ8 or the HLA DR4-DQ8/DR4-DQ8 genotype. P values were calculated using log-rank tests. The groups were defined by quartiles of the genetic risk score (green: lower quartile; blue: two medium quartiles; orange: upper quartile). HLA, human leucocyte antigen.
Importantly, this risk score was predictive also when weighting for the HLA class II genes was not included, suggesting the impact of genetic variants on progress is independent of either HLA DR3/4-DQ8 or DR4-DQ8/DR4-DQ8 genotypes in these TEDDY participants. Previous studies have indicated that protective HLA genotypes are associated with a slower progression to clinical diabetes in islet autoantibody-positive individuals, but there are little or no differences in the progression rate between the high-risk HLA class II genotypes. Our findings are consistent with previous reports of associations between the rate of progression from preclinical to clinical type 1 diabetes and individual type 1 diabetes susceptibility genes. Of practical relevance, a low genetic risk score may be used to identify a subset of islet autoantibody-positive children with slower progression to clinical type 1 diabetes, and therefore be an exclusion criterion for the sake of interpretability; however, exact age (ie, not rounded) was used in the regression models.

Author affiliations

1Institute of Diabetes Research, Helmholtz Zentrum München, and Forschgruppe Diabetes, Technical University of Munich, at Klinikum rechts der Isar, Munich-Neuherberg, Germany
2DFG Center for Regenerative Therapies Dresden, Faculty of Medicine, Technische Universität Dresden, Dresden, Germany
3Forschgruppe Diabetes eV at Helmholtz Zentrum München, Munich-Neuherberg, Germany
4Health Informatics Institute, Morsani College of Medicine, University of South Florida, Tampa, Florida, USA
5Barbara Davis Center for Childhood Diabetes, University of Colorado, Aurora, Colorado, USA
6Pacific Northwest Diabetes Research Institute, Seattle, Washington, USA
7Department of Clinical Sciences, Lund University/CRC, Skåne University Hospital, Malmö, Sweden
8Center for Biotechnology and Genomic Medicine, Medical College of Georgia, Augusta University, Augusta, Georgia, USA
9Institute of Biomedicine, Research Centre for Integrative Physiology and Pharmacology, Turku University Hospital, Turku, Finland
10Department of Physiology, University of Turku, Turku, Finland
11National Institute of Diabetes and Digestive and Kidney Diseases, Bethesda, Maryland, USA
12Center for Public Health Genomics, University of Virginia, Charlottesville, Virginia, USA

Contributors AB (guarantor) analysed the data and wrote the first and final draft of the manuscript together with EB and A-GZ. KV, MH, CW, BIF, AKS, WAH, JPK, AL, MIR, J-IX, IT, BA and SSR contributed to the interpretation of the results, reviewed the manuscript and contributed to subsequent drafts.

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Competing interests A patent has been applied for (LU100334) with the title ‘Method the risk to develop type 1 diabetes’ by Helmholtz Zentrum München Deutsches Forschungszentrum für Gesundheit und Umwelt. EB, A-GZ and CW are among the inventors. The patent includes the genetic score that is examined in the manuscript.

Patient consent Not required.

Ethics approval The study was approved by local institutional review boards and is monitored by an external advisory board established by the US National Institutes of Health.

Provenance and peer review Not commissioned; externally peer reviewed.

Data sharing statement The program code and the data can be provided upon reasonable request.

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REFERENCES
