FOXN3 hyperglycemic risk allele and insulin sensitivity in humans

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ABSTRACT

Objective The rs8004664 variation within the FOXN3 gene is significantly and independently associated with fasting blood glucose in humans. We have previously shown that the hyperglycemia risk allele (A) increases FOXN3 expression in primary human hepatocytes; over-expression of human FOXN3 in zebrafish liver increases fasting blood glucose; and heterozygous deletion of the zebrafish ortholog foxn3 decreases fasting blood glucose. Paralleling these model organism findings, we found that rs8004664 A/A homozygotes had blunted glucagon suppression during an oral glucose tolerance test. Here, we test associations between insulin sensitivity and the rs8004664 variation.

Research design and methods 92 participants (49±13 years, body mass index: 32±6 kg/m², 28 with and 64 without type 2 diabetes mellitus) were genotyped at rs8004664. Insulin sensitivity was measured by the euglycemic-hyperinsulinemic clamp technique.

Results The “A” allele frequency was 59%; the protective (G) allele frequency was 41% (A/A: n=29; G/G: n=12; A/G: n=50). Clamp-measured glucose disposal rate (GDR) was not different by genotype (F=0.046, p=0.96) or by “A” allele carrier (p=0.36). Female G/G homozygotes had better insulin sensitivity compared to female “A” allele carriers (GDR; G/G: 9.9±3.0 vs A/A+G/G: 7.1±3.0 mg/kg fat-free mass+17.7/min; p=0.04). Insulin sensitivity was not different by genotype or by “A” allele carriers.

Conclusion The rs8004664 variation within the FOXN3 gene may modulate insulin sensitivity in women.

INTRODUCTION

In a large cohort of non-diabetic subjects, the single nucleotide variation rs8004664 within the first intron of the FOXN3 gene in humans was found to be significantly and independently associated with fasting blood glucose.1 The molecular mechanism for how this variation increases the fasting blood glucose set-point remains elusive; nevertheless, we previously showed that the fasting hyperglycemia allele at rs804464 increases FOXN3 expression in primary human hepatocytes.2 We modeled this increased liver FOXN3 expression by over-expressing the human FOXN3 cDNA in zebrafish livers and observed an increase in fasting blood glucose without any additional dietary challenge. Since FOXN3 is a transcriptional repressor,3 we performed whole transcriptome analyses in livers over-expressing FOXN3: the MYC ortholog mycb transcript, which encodes a driver of liver glucose utilization during fasting,4 was strongly down-regulated. We showed that FOXN3 directly represses MYC expression.5 This indicates that liver FOXN3 increases fasting blood glucose by repressing a driver of liver glucose utilization, providing more glucose for export from the liver.6

In follow-up investigations, we found that glucagon injection into mice rapidly decreases liver FOXN3 protein, indicating hormonal regulation of FOXN3. When we prepared a viable loss-of-function mutation in the orthologous foxn3 gene, we observed decreased fasting blood glucose, blood glucagon, and alpha cell mass.7 Concordantly, over-expression of human FOXN3 in zebrafish liver increased alpha cell mass in the zebrafish endocrine pancreas. In this second study, we also explored the effect of the rs8004664 variation within the foxn3 gene in livers over-expressing FOXN3: the MYC ortholog mycb transcript, which encodes a driver of liver glucose utilization during fasting,4 was strongly down-regulated. We showed that FOXN3 directly represses MYC expression.5 This indicates that liver FOXN3 increases fasting blood glucose by repressing a driver of liver glucose utilization, providing more glucose for export from the liver.6

Significance of this study

What is already known about this subject?
► The role of FOXN3 in modulating insulin sensitivity was not known.
► We genotyped a cohort of subjects who underwent high-dose euglycemic-hyperinsulinemic clamp for the rs8004664.
► The fasting hyperglycemia variant of rs8004664 was associated with increased glucose disposal in female subjects, but not in men or the combined cohort.

What are the new Findings?
► FOXN3 may modulate insulin sensitivity in a sexually dimorphic manner.
variation on oral glucose tolerance in a large cohort of human subjects: rs8004644 hyperglycemia risk allele carriers show diminished suppression of glucagon over the oral glucose tolerance test (as reflected by decreased area below baseline), but show no differences in fasting glucagon.5

Our working model for how FOXN3 regulates fasting glucose does not exclude a potential role for insulin sensitivity, and therefore glucose disposal rate (GDR) during a glucose clamp. Here, we tested in a cohort of adults with and without type 2 diabetes whether the rs8004644 variation modulates insulin-mediated glucose uptake by examining associations between rs800464 variants and insulin sensitivity measured by the gold-standard euglycemic-hyperinsulinemic clamp technique.6

METHODS

Study population
In this single group cross-sectional design, 92 participants who previously underwent a euglycemic-hyperinsulinemic clamp6 were genotyped at the rs8004644 variant. Participants were initially part of a larger prospective study called “The Pennington Center Longitudinal Study” designed to assess the effects of obesity and lifestyle factors on the chronic disease development, including type 2 diabetes mellitus. The current study utilized a subset of participants (with available DNA) from the original cohort, consisting of 92 white adults. Participants were categorized as having type 2 diabetes by self-report (“yes” response to having diabetes) or by fasting plasma glucose ≥126 mg/dL, on study visit.

Anthropometrics/body composition
As previously described,6 metabolic body weight and height were measured, and body mass index was calculated from these respective values. Per cent body fat was determined from dual-energy X-ray absorptiometry (DXA, Hologic QDR 4500A; Hologic, Bedford, MA, USA). Fat mass and fat-free mass (FFM) were calculated from these respective values. Per cent body fat was determined from dual-energy X-ray absorptiometry (DXA, Hologic QDR 4500A; Hologic, Bedford, MA, USA). Fat mass and fat-free mass (FFM) were calculated from these respective values.

Insulin sensitivity
Peripheral insulin sensitivity was measured using the gold-standard euglycemic-hyperinsulinemic clamp technique, as previously described.5 Briefly, plasma glucose was clamped between 90 and 100 mg/dL during continuous insulin infusion (120 mU/m²/min). GDRs presented herein are normalized for metabolic size of the participant (FFM +17.7) as well as average insulin concentrations during steady state of the clamp procedure.7 Average insulin concentrations measured during steady state of the clamp for the entire study cohort was 227.1±68 µU/mL. Fasting glucose, insulin, lipids and triglycerides were measured from a fasting blood draw before the clamp procedure.

RESULTS
Participant characteristics are shown in table 1. No baseline differences were observed. Of the 92 participants, 29 were homozygous for the fasting hyperglycemia risk

| Characteristic | Genotype | Genotype |
|---------------|----------|----------|
| Age (y)       | 50±13    | 46±15    | 0.32   |
| Sex (M/F)     | 36/43    | 7/6      | NA     |
| BMI (kg/m²)   | 33±6     | 32±7     | 0.91   |
| Body weight (kg) | 93±19     | 93±22     | 0.99   |
| FFM (kg)      | 58.9±11.7| 59.6±14.0| 0.85   |
| Fasting glucose (mg/dL) | 102±17 | 102±12 | 0.97 |
| Fasting insulin (µU/mL) | 15±7 | 17±21 | 0.56 |
| HOMA-IR       | 3.9±2.1  | 4.6±6.2  | 0.40   |
| DXA (% Fat)   | 36±9     | 35±8     | 0.83   |
| Cholesterol (mg/dL) | 203±42 | 200±44 | 0.79 |
| HDL (mg/dL)   | 50±12    | 53±17    | 0.49   |
| LDL (mg/dL)   | 120±36   | 117±43   | 0.80   |
| Triglycerides (mg/dL) | 165±115 | 148±118 | 0.61 |
| Diabetes status (Y/N) | 25/54 | 3/10 | NA |
| Glucose disposal (mg/kg FFM +17.7/min) | 6.7±2.9 | 7.5±3.1 | 0.36 |

BMI, body mass index; FFM, fat-free mass; HDL, high-density lipoprotein; LDL, low-density lipoprotein.

Genotyping
DNA was extracted from buffy coat of blood samples (Qiagen PureGene Blood Core Kit #158389). Genotyping of the rs8004644 variant was completed using a TaqMan SNP Genotyping Assay (ThermoFisher ID C__29386020_10) as previously described.5

Statistical analysis
Data are presented as mean±SD, unless otherwise noted. SPSS (IBM V. 25) was used for statistical analysis. Participants were grouped by genotype (G|G, A|G, and A|A), and differences in metabolic outcomes were tested using one-way analysis of variance. Secondarily, participants were grouped by hyperglycemic risk-carrying allele versus protective allele homozygotes (A|G+A|G combined vs G|G), and differences were tested using unpaired Student’s t-test. Significance was accepted at p=0.05. To explore the role of type 2 diabetes status on the relationship between genetic variants and metabolic outcomes, participants were categorized by diabetes status and a similar analysis was performed among these two groups (those with and without type 2 diabetes). Furthermore, the relationship between genetic variants and metabolic outcomes was also performed separately among women and men.
allele (A|A), 13 were homozygous for the protective allele (G|G), and 50 participants were heterozygous (A|G). The high-risk allele (A) frequency was 59% and the protective allele frequency (G) was 41%. No differences were observed in GDR when grouped by genotype (F = 0.046, p = 0.96) or when grouped by carrier of hyperglycemic risk allele versus homozygotes for the protective allele “G” (p = 0.36; table 1).

Among those with type 2 diabetes (n=28), no significant differences were observed in GDR by genotype group (F = 0.852, p = 0.44) or by carrier of hyperglycemic risk allele (p = 0.54). Similarly, among those without type 2 diabetes (n=64), no significant differences were observed in GDR grouped by genotype (F = 0.149, p = 0.86) or by carrier of hyperglycemic risk allele (p = 0.61). Insulin concentrations were not different by genotype group (INS, G|G: 261±149.7 vs A|A+AG: 221.5±41.6 µU/mL; p = 0.05) although trending. This is driven by a single outlier within the GG group. Re-analysis after outlier removal shows similar means between groups (INS, G|G: 225.8±69.9 vs A|A+AG: 221.5±41.6 µU/mL; p = 0.87). Insulin concentrations were not different between males and females (INS, males: 235.6±83.5 vs female: 220.3±42.8 µU/mL; p = 0.26).

Female homozygotes for the protective allele “G” had significantly higher GDR (indicative of increased insulin sensitivity) compared to female carriers of the hyperglycemic-risk allele “A” (G|G: 9.9±3.0 vs A|A+AG: 7.1±3.0 mg/kgFFM/min; p = 0.04). Homeostatic model assessment of insulin resistance (HOMA-IR) was however not different between groups (p = 0.33). However, re-analysis of data after removal of an outlier with excessive hyperinsulinemia revealed a significant difference (G|G: 1.6±0.8 vs A|A+AG: 3.8±1.9; p = 0.02). When grouped by genotype, no differences were observed in GDR (F = 2.310, p = 0.11) or HOMA-IR (F = 0.507, p = 0.61). Among men, no significant differences were observed in GDR grouped by genotype group (F = 0.592, p = 0.56) or by high-risk carrier (p = 0.43).

**DISCUSSION**

This is the first study to examine the links between FOXN3 genetic variants and insulin sensitivity measured by the euglycemic-hyperinsulinemic clamp technique in humans. GDR (insulin sensitivity) was not statistically different by genotype (G|G, A|G, and A|A), or when grouped by hyperglycemic risk-carrying allele (G vs A|A+AG). Among women, protective allele homozygotes (G|G) had significantly higher GDR (indicative of better insulin sensitivity) compared to carriers of the hyperglycemic high-risk allele, suggesting a potential role for sex in modulating the relationship between genotype and glucose metabolism. Indeed, we observed a greater increase in fasting blood glucose when FOXN3 was over-expressed in female zebrafish liver.2 The findings herein are supportive but should be interpreted cautiously due the limited sample size (G|G: n=6; A|A+AG: n=43).

While the rs8004664 fasting hyperglycemia variation within the FOXN3 gene does not appear to impact glucose disposal, the encoded protein’s role in the development of type 2 diabetes merits further investigation. In particular, FOXN3’s role in responding to and modulating glucagon function will be the subject of future studies. It is clear from a large body of preclinical and early clinical work that total genetic or pharmacological blockade of hepatic glucagon action is not a viable therapeutic approach. While blood glucose is lowered, hepatic transaminases, circulating triglycerides, and low-density lipoprotein cholesterol are elevated; and alpha cell hyperplasia occurs when glucagon receptor-blocking antibodies are administered.8 Thus, a more thorough understanding of the effects and effectors of glucagon action could lead to the development of more selective, glucagon-leveraging therapies for type 2 diabetes mellitus.9 10

In summary, we provide novel data on the relationship between the FOXN3 rs8004664 variant and insulin sensitivity measured by the euglycemic-hyperinsulinemic clamp technique in humans. We did not find differences among genotype, suggesting that other physiological factors may be responsible for modulating fasting blood glucose. To further define the liver FOXN3-glucagon axis, future studies should explore links between this variant and other physiological modulators of glycemia including hepatic insulin sensitivity, glucagon secretion, and insulin secretion. Critically, FOXN3 is expressed in several tissues, often by more than one cell type in a tissue.11 Thus, a combination of human physiological characterization of the effect of the rs8004664 variation and a systematic effort to modulate FOXN3 expression in different cell types in model organisms is warranted.

**Contributors** MLE analyzed and interpreted data, and drafted manuscript. SK conceptualized study, collected and interpreted data, and edited the manuscript. ER conceptualized study, collected and interpreted data, and edited the manuscript. AS conceptualized study, collected and interpreted data, and edited the manuscript. ER is the guarantor of the study, having full access to all study data; he had the final responsibility for data integrity, accuracy of data analysis, and decision to submit for publication. All authors have approved the final version of this manuscript.

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**Competing interests** None declared.

**Ethics approval** The study was performed in accordance with the principles contained within the Declaration of Helsinki. The protocol was approved by the Institutional Review Board of the Pennington Biomedical Research Center. All participants provided informed, written consent prior to study participation.

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**Data availability statement** Data are available upon reasonable request.

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