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Lamin A/C Polymorphisms, Type 2 Diabetes, and the Metabolic Syndrome:

Case-Control and Quantitative Trait Studies

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Abstract

Mutations in the *LMNA* gene, encoding the nuclear envelope protein lamin A/C, are responsible for a number of distinct disease entities including Dunnigan-type familial partial lipodystrophy. Dunnigan-type lipodystrophy is characterized by loss of subcutaneous adipose tissue, insulin resistance, dyslipidemia, and type 2 diabetes and shares many of the features of the metabolic syndrome. Furthermore, several genome-wide linkage scans for type 2 diabetes have found evidence of linkage at chromosome 1q21.2, the region that harbors the *LMNA* gene. Therefore, *LMNA* is a biological and positional candidate for type 2 diabetes susceptibility. Previous studies have reported association between a common *LMNA* variant (1908C>T; rs4641) and adverse metabolic traits in ethnically diverse populations from Asia and North America. In the present study, we characterized the common variation across the *LMNA* gene (including rs4641) and tested for association with type 2 diabetes in two large case-control studies ($n = 2,052$) and with features of the metabolic syndrome in a separate cohort study ($n = 1,572$). Despite our study being sufficiently powered to detect effects similar and even smaller in magnitude than those previously reported, none of the *LMNA* single nucleotide polymorphisms were statistically significantly associated with type 2 diabetes or the metabolic syndrome. Thus, it appears unlikely that variation at *LMNA* substantially increases the risk of type 2 diabetes or related traits in U.K. Europeans.

Deleterious mutations in the *LMNA* gene, encoding the nuclear envelope protein lamin A/C, are responsible for a number of distinct disease entities known as “laminopathies” including Dunnigan-type familial partial lipodystrophy (FPLD; OMIM 151660) (1-5). Dunnigan-type FPLD is characterized by loss or absence of adipose tissue and severe metabolic derangements, including insulin resistance, dyslipidemia, and type 2 diabetes. Recent evidence has demonstrated that common variants in genes previously implicated in monogenic disorders of glucose imbalance increase the risk of common type 2 diabetes (6). These observations support the examination of common variants in *LMNA* for their effect

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on type 2 diabetes and metabolic syndrome risk. Furthermore, genome-wide linkage scans for type 2 diabetes have found evidence for linkage at chromosome 1q21-q24 (7-9), the region that harbors the *LMNA* gene. Therefore, *LMNA* is a biological and positional candidate for type 2 diabetes susceptibility.

The most frequently studied variant at the *LMNA* gene in relation to type 2 diabetes- and metabolic syndrome-related traits is the silent C>T substitution at nucleotide 1908 (1908C>T; rs4641). This variant affects the third base at codon 566 (His566His) in exon 10, just before the alternative splicing site that gives rise to the two distinct proteins, lamin A and C (10). Several studies have reported on the association between the 1908T allele and type 2 diabetes-related traits in ethnically diverse populations (11-17).

In the present study, which is the most comprehensive of its kind on the *LMNA* gene, we characterized the common variation at the *LMNA* gene (including 1908C>T) and tested for association with risk of type 2 diabetes and related phenotypes in three large population-based cohorts.

RESEARCH DESIGN AND METHODS

This study included two type 2 diabetes case-control studies of middle-aged U.K. Euroid: the type 2 diabetes case-control European Prospective Investigation into Cancer (EPIC) Study and the Cambridgeshire Case-Control (CCC) Study. Genetic associations with the features of the metabolic syndrome were assessed in a separate cohort of middle-aged U.K. Euroids (the Medical Research Council [MRC] Ely Study). Participants from these studies reside within a region that has traditionally had low admixture rates. Because of its genetic homogeneity, it is likely to be free from population stratification, which may be an important source of confounding in some cohorts. Descriptive characteristics are given in online appendix Table 1 (available at <http://dx.doi.org/10.2337/db06-1055>).

The EPIC Study is nested within the EPIC Norfolk Study, a population-based cohort study of Euroid men and women aged 40–78 years. Both the case-control and full cohort (18,19) study have been previously described in detail. Briefly, the case-control study consists of 417 incident type 2 diabetic case subjects and two sets of 417 control subjects, each matched in terms of age, sex, general practice, and recruitment date, with one set additionally matched for BMI. A case was defined by a physician's diagnosis of type 2 diabetes, with no insulin prescribed within the 1st year following diagnosis, and/or A1C >7% at the health check. Control subjects were randomly selected from the EPIC-Norfolk cohort from among those without diabetes, cancer, stroke, or myocardial infarction at baseline and who had not developed diabetes by the time of selection. Potential control subjects with measured A1C levels >6% were excluded.

The CCC Study consists of 552 patients with type 2 diabetes aged 45–76 years, randomly sampled from a population-based diabetes register, and 552 control subjects recruited from the same population and individually matched for age, sex, and geographical location. The study design and methods have been described in detail elsewhere (20). Briefly, case subjects were defined by onset of diabetes after the age of 30 years, without insulin treatment in the 1st year following diagnosis. Diabetes was excluded in control subjects by medical record search and by an A1C measurement <6%. In total, 2,052 individuals (41.5% with type 2 diabetes) from both the EPIC Study and CCC Study were genotyped for the 13 *LMNA* variants.

The MRC Ely Study commenced in 1990 as a prospective population-based cohort study of the etiology and pathogenesis of type 2 diabetes and related metabolic disorders. Part of the study design and methods have been described in detail elsewhere (21,22). At baseline

(phase 1), 1,122 individuals (aged 40–67 years) were randomly selected from Ely (a town in East Anglia, U.K.). Of these, 937 were followed-up 4.5 years later, and an additional 183 young adults (aged 30–40 years) were recruited for the same clinical examinations (phase 2). Phase 3 was carried out 10 years after the start of the study. Of all phase 1 and 2 participants, 936 attended phase 3 examinations and 714 were newly recruited from the same geographical area. The current study examines the individuals of phase 3 ($n = 1,572$ men and women aged 35–79 years) for whom genotypic and phenotypic data (metabolic traits) were available. Of all participants, 30.2% had the metabolic syndrome as previously defined (based on obesity, dyslipidemia, elevated blood pressure, glucose intolerance, and insulin resistance) (23). Ethical permission for the three studies was granted by their respective local research ethics committee, and study participants provided written informed consent.

Single nucleotide polymorphism selection

Single nucleotide polymorphism (SNP) selection was undertaken before the release of the International HapMap (24) and was based on literature reports and on the dbSNP database. The selection was performed in a stepwise manner to ensure even-spaced coverage of the gene, including 10 kb up- and downstream of the gene. First, we selected all common SNPs, as well as all polymorphic missense variants, that were reported in the literature. Subsequently, we selected all common SNPs (minor allele frequency [MAF] $\geq 5\%$) and missense variants that were catalogued in the dbSNP database. Furthermore, SNPs from the dbSNP database for which no allele frequency was reported but for which validation from multiple sources was available were also included. Finally, to ensure coverage of at least one SNP every 3 kb, we also included SNPs from the dbSNP database for which no allele frequency was available. Eventually, 20 SNPs were genotyped, of which 13 SNPs that had a MAF $>1\%$ were included in the analyses (Fig. 1).

The International HapMap Phase 2 (24) reports 14 common SNPs (MAF ≥ 0.05) for the Caucasian population (CEU), which can be tagged by four tag SNPs. Of the 13 SNPs we genotyped, 3 (SNP1, SNP2, and SNP9) were in common with HapMap SNPs. These 3 SNPs capture the genetic variation of 8 (57%) of the 14 HapMap SNPs (r^2 threshold at 0.80). The six HapMap SNPs that were not captured by SNP1, SNP2, and SNP9 are in perfect linkage disequilibrium (LD) ($r^2 = 1$) and can be tagged by one SNP. The remaining 10 SNPs we genotyped were not typed in the HapMap project, such that we cannot accurately assess the coverage of the genetic variation in *LMNA*. However, on the basis of their physical location on the gene, we can expect that at least four of our SNPs may represent the remaining six HapMap SNPs, as these four SNPs are located within the six-SNP block that shows perfect LD. Therefore, we can conclude that our 13 SNPs adequately represent the common genetic variation in the *LMNA* gene (online appendix Fig. 1 and Table 2). All genotypes conformed to expectations under Hardy-Weinberg equilibrium ($P > 0.01$).

Genetic analyses

Samples were arrayed on 96-well plates with three replicates and one water control per plate. For the case-control populations, case and control samples were randomly distributed across each 96-well plate, with approximately the same number of cases and controls per plate. Genotyping of samples was performed in 384-well plates at the Wellcome Trust Sanger Institute, Cambridge, using an adaptation of the homogenous MassExtend protocol supplied by Sequenom for the MassArray system (25). Call rates $\geq 95\%$ were observed for all SNPs, except SNP 11 that had an 86% call rate. Concordance rates between duplicate samples were $\geq 98\%$.

Statistical analysis

Analyses were performed using Stata SE 8.2 for Windows (StataCorp, College Station, TX) and SPSS 12.0 for Windows (SPSS, Chicago, IL). Genotype frequencies were tested for Hardy-Weinberg equilibrium using the χ^2 test. LD was assessed using the expectation-maximization algorithm. We calculated the effect of *LMNA* variants on discrete outcomes (type 2 diabetes and metabolic syndrome) using logistic regression. Before combining the two type 2 diabetes case-control cohorts, we tested for heterogeneity between the populations. For one SNP (rs536857), evidence ($P = 0.04$) of heterogeneity was observed, which was adjusted for in a random effects model. To test association between *LMNA* variants and continuous metabolic traits, we used generalized linear models adjusted for age, sex, BMI, and study. We first tested for an additive effect of the SNP on the risk of type 2 diabetes and the metabolic traits. If these results showed evidence for a dominant or recessive inheritance, the respective models were tested as well. Because the number of rare homozygotes was small ($n = 1-5$) for SNP3 to SNP11, a dominant model was also tested. We tested for sex-specific effects by including genotype-by-sex interaction terms in the models. Since no evidence for genotype-by-sex interaction was found ($P > 0.24$), we do not present stratified results. Haplotype blocks were defined using Gabriel's method (26), as implemented in Haploview (27). Haplotype frequencies were inferred using the expectation-maximization algorithm as implemented in the HelixTree Genetic Analyses Software (version 4.4.1; Golden Helix, Bozeman, MT). Haplotypes prevalent at $>5\%$ were retained for further analysis. A χ^2 test was used to compare haplotype frequencies of case and control samples. Haplotype trend regression analysis was performed to test for association between the haplotypes and continuous metabolic traits. Power calculations were performed using Quanto version 1.1.1 (<http://hydra.usc.edu/gxe>).

RESULTS

Figure 2A shows the extent of LD (r^2) between SNPs. In the EPIC Study and CCC Study, none of the SNPs were associated with diabetes risk after adjustment for age, sex, BMI, and study (Fig. 2B). In the MRC Ely Study, none of the SNPs were associated with the metabolic syndrome after adjustment for age and sex (Fig. 2C). Results for the unadjusted models were similar (data not shown).

We tested for association between the 13 SNPs and continuous metabolic traits (i.e., plasma HDL, LDL, and total cholesterol and BMI) in a combined nondiabetic population comprising the MRC Ely Study, as well as control subjects from both the EPIC and CCC Studies. We also examined the associations with fasting glucose and insulin, which was only available in the MRC Ely Study. None of the *LMNA* variants were significantly associated with any of the continuous metabolic traits, adjusted for age, sex, BMI, and study (online appendix Table 3). However, carriers of the 1908T allele tended to have lower triglyceride ($P = 0.06$) and total cholesterol ($P = 0.05$) levels than C1908C homozygotes, and SNP1 T-allele carriers tended to have lower glucose levels ($P = 0.05$).

None of the haplotypes were significantly associated with risk of type 2 diabetes or the metabolic syndrome. Also, for the continuous metabolic traits, haplotypes did not provide any additional information beyond the individual marker associations (online appendix Tables 6 and 7).

Both case-control and cohort studies had sufficient power to detect similar and even smaller effect sizes than those previously reported; i.e., for the previously studied 1908C>T (rs4641) variant, our study was powered at 95% to detect an odds ratio (OR) 1.35 for type 2 diabetes per copy of the minor allele, at a significance of 1%. For SNPs 1, 2, 12, and 13, we had sufficient power ($>80\%$) to detect ORs >1.25 in the case-control study and differences of

>2% in the cohort study at a significance of 5%. SNP3 to SNP11 had a lower MAF and are therefore slightly less powered; i.e., we had 69–83% power to detect ORs >1.4 in the case-control study, and differences as small as 3% could be detected in the cohort study at a significance of 5% (online appendix Tables 4 and 5).

DISCUSSION

In the present study, the largest of this kind on the *LMNA* gene, we were unable to detect association between variants in the *LMNA* gene and risk of type 2 diabetes or the metabolic syndrome, despite our study being sufficiently powered. We observed tentative associations between 1908C>T and lipid levels. However, the direction of these trends contrast with those reported in other studies, and we conclude that our observations are likely to be false positive due to the number of statistical comparisons undertaken. Given the appropriate design of this study and the fact that no other associations were detected strongly suggests that, in this population, *LMNA* variants are unlikely to play a major role in the development of type 2 diabetes or metabolic syndrome. However, small effects (i.e., OR <1.25 or <3% for quantitative traits) cannot be excluded.

The *LMNA* gene encodes two nuclear proteins, lamin A and C, which are essential structural components of most differentiated mammalian cells. Rare mutations in exon 8 of *LMNA* cause the autosomal-dominant Dunnigan-type FPLD (2-5). At birth, individuals with FPLD have normally distributed adipose tissue but experience a progressive loss of subcutaneous fat from the extremities, gluteals, and torso following puberty. In later life, individuals with FPLD often experience severe metabolic derangements, including insulin resistance, dyslipidemia, heart disease, and type 2 diabetes. Mutations elsewhere in the *LMNA* gene are associated with several additional autosomal-dominant diseases (1).

Common variants in the *LMNA* gene, the 1908C>T variant in particular, have been associated with the risk of type 2 diabetes and with related metabolic phenotypes in several ethnically distinct populations (11-17). Hegele and colleagues (16,17) first reported that in two genetically distinct aboriginal populations, Oji-Cree and Inuit, the T1908T homozygotes had significantly higher measures of obesity-related traits than C-allele carriers. In a case-control study of Japanese men, Murase et al. (15) reported an association of borderline statistical significance ($P=0.08$) between the 1908C>T variant and type 2 diabetes, with the 1908T allele being more frequent (44%) in case than control subjects (32%). In addition, T-allele carriers of both case and control subjects had significantly higher fasting insulin, cholesterol, and triglyceride levels than C/C homozygotes. In the case group from the same study, Liang et al. (14) reported a modest ($P=0.06$) increased risk of diabetic nephropathy in 1908T allele carriers compared with 1908C allele homozygotes. In 908 Old Order Amish (13), six *LMNA* SNPs were identified and tested for association with type 2 diabetes and the metabolic syndrome. Only the 1908C>T variant was modestly ($P<0.03$) associated with the metabolic syndrome and triglyceride and HDL cholesterol levels but not with diabetes. In a small cohort of Pima Indians, weak evidence of linkage (logarithm of odds 1.73) for subcutaneous abdominal adipocyte size was localized to the *LMNA* region of chromosome 1 (1q21-q23), and association analyses showed that adipocytes of C1908C homozygotes were larger than those of T1908T homozygotes (12). As enlarged subcutaneous abdominal adipocyte size is an independent predictor of type 2 diabetes in Pima Indians (28), these data suggest that C1908C homozygotes may be at increased risk of diabetes, which contradicts the findings by Steinle et al. (13). In a subsequent study of 1,338 Pima Indians, no evidence of association was observed between the 1908C>T variant and type 2 diabetes or any related quantitative traits (11). Taken together, although the *LMNA* gene appears to be a strong functional and positional candidate for type 2 diabetes and related phenotypes, the associations reported thus far have been weak and inconsistent.

In the present study, participants were exclusively European, living in the same geographical area in eastern England. Although our current and previous findings suggest that this is an ethnically homogeneous population, we cannot completely exclude the existence of underlying genetic heterogeneity. It should also be highlighted that the current study was aimed at examining the effect of common genetic variation in the *LMNA* gene, and we cannot exclude the possibility that rare SNPs (MAF <0.05) may affect the risk of type 2 diabetes or related metabolic traits.

The control subjects of the case-control studies were not tested for type 2 diabetes, which may introduce misclassification in the control sample and, as a consequence, bias the observation toward the null. We have attempted to diminish this misclassification using A1C. At an A1C level of 6%, previous reports have suggested sensitivity to be 100% and specificity 91% (29). Thus, by using this cutoff, we would exclude all the cases of undiagnosed diabetes in the control subjects but would only be excluding <10% of the potential control subjects who in fact had normal glucose tolerance.

We conclude that although common genetic variation at *LMNA* may underlie adverse metabolic phenotypes in Asian and North American populations, our results in large population-based cohorts indicate that it is unlikely that common variation in the *LMNA* gene increases the susceptibility to type 2 diabetes or features of the metabolic syndrome in U.K. Europeans.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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Glossary

CCC	Cambridgeshire Case-Control
EPIC	European Prospective Investigation into Cancer
FPLD	familial partial lipodystrophy
LD	linkage disequilibrium
MAF	minor allele frequency
MRC	Medical Research Council
SNP	single nucleotide polymorphism

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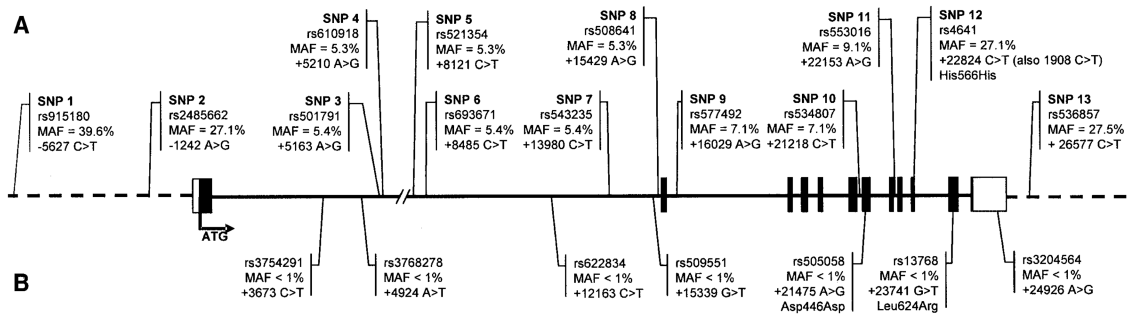


FIG. 1. Position (relative to the ATG methionine initiation codon) and MAF of the *LMNA* SNPs selected for genotyping. *A*: Polymorphic (MAF ≥ 1%) SNPs included in the analyses. *B*: SNPs with a MAF < 1%, not analyzed.

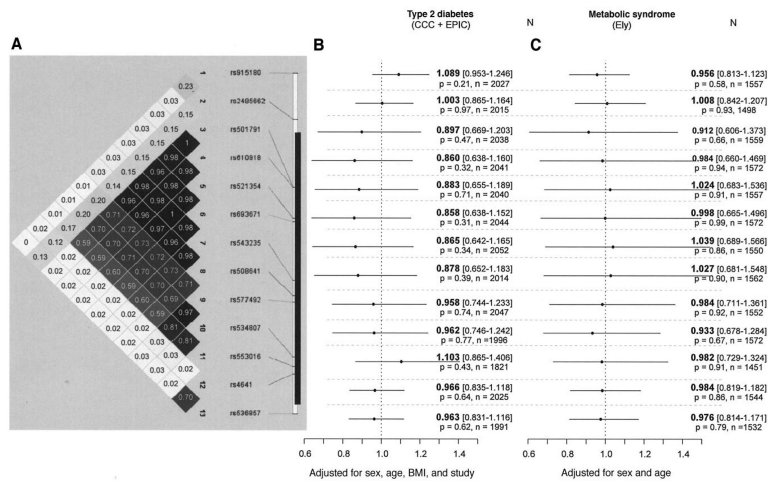


FIG. 2.
A: Pairwise LD estimates (r^2 values) assessed in control subjects for the 13 polymorphic *LMNA* SNPs included in the study. (The black solid line represents the gene region.) **B:** Adjusted OR of *LMNA* variants for type 2 diabetes incidence for the additive model. **C:** Adjusted OR of *LMNA* variants for metabolic syndrome incidence for the additive model. Values in **B** and **C** represent ORs (increased risk per additional minor allele) and 95% CIs (OR [95 CI]), *P* values for the additive model, and number of participants with full genotype and phenotype data.