Utility of genetic and non-genetic risk factors in prediction of type 2 diabetes: Whitehall II prospective cohort study

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ABSTRACT

Objectives To assess the performance of a panel of common single nucleotide polymorphisms (genotypes) associated with type 2 diabetes in distinguishing incident cases of future type 2 diabetes (discrimination), and to examine the effect of adding genetic information to previously validated non-genetic (phenotype based) models developed to estimate the absolute risk of type 2 diabetes.

Design Workplace based prospective cohort study with three 5 yearly medical screenings.

Participants 5535 initially healthy people (mean age 49 years; 33% women), of whom 302 developed new onset type 2 diabetes over 10 years.

Outcome measures Non-genetic variables included in two established risk models—the Cambridge type 2 diabetes risk score (age, sex, drug treatment, family history of type 2 diabetes, body mass index, smoking status) and the Framingham offspring study type 2 diabetes risk score (age, sex, parental history of type 2 diabetes, body mass index, high density lipoprotein cholesterol, triglycerides, fasting glucose)—and 20 single nucleotide polymorphisms associated with susceptibility to type 2 diabetes. Cases of incident type 2 diabetes were defined on the basis of a standard oral glucose tolerance test, self report of a doctor's diagnosis, or the use of anti-diabetic drugs.

Results A genetic score based on the number of risk alleles carried (range 0-40; area under receiver operating characteristics curve 0.54, 95% confidence interval 0.50 to 0.58) and a genetic risk function in which carriage of risk alleles was weighted according to the summary odds ratios of their effect from meta-analyses of genetic studies (area under receiver operating characteristics curve 0.55, 0.51 to 0.59) did not effectively discriminate cases of diabetes. The Cambridge risk score (area under curve 0.72, 0.69 to 0.76) and the Framingham offspring risk score (area under curve 0.78, 0.75 to 0.82) led to better discrimination of cases than did genotype based tests. Adding genetic information to phenotype based risk models did not improve discrimination and provided only a small improvement in model calibration and a modest

net reclassification improvement of about 5% when added to the Cambridge risk score but not when added to the Framingham offspring risk score.

Conclusion The phenotype based risk models provided greater discrimination for type 2 diabetes than did models based on 20 common independently inherited diabetes risk alleles. The addition of genotypes to phenotype based risk models produced only minimal improvement in accuracy of risk estimation assessed by recalibration and, at best, a minor net reclassification improvement. The major translational application of the currently known common, small effect genetic variants influencing susceptibility to type 2 diabetes is likely to come from the insight they provide on causes of disease and potential therapeutic targets.

INTRODUCTION

Around one in 10 middle aged Europeans develop type 2 diabetes over a 10 year period.¹ This may rise with the threatened epidemic of type 2 diabetes, driven partly by the increase in obesity, which is projected to result in a prevalence of type 2 diabetes worldwide of 366 million by 2030 (www.who.int/diabetes/facts/world_figures/en/).² This will place a considerable burden on healthcare systems, as type 2 diabetes is a common cause of renal failure and blindness and increases the risk of cardiovascular diseases and mortality.

Estimation of people's absolute risk of coronary heart disease, based on the Framingham, PROCAM, and QRISK equations,³⁻⁵ has become established as a means of targeting preventive interventions to those at highest risk. Earlier risk functions for estimating the risk of type 2 diabetes required an oral glucose tolerance test, which is impractical for screening large numbers of people. Despite evidence that some interventions prevent or delay development of type 2 diabetes in people at high risk,⁶ the prediction of risk of new onset type 2 diabetes has not been part of routine clinical practice.

Two risk scores were published in 2007 and 2008. They were based on demographic, anthropometric,

and biochemical measures but did not require information from an oral glucose tolerance test. The Cambridge type 2 diabetes risk score is a pragmatic algorithm incorporating the routinely assessed variables age, sex, drug treatment, family history of type 2 diabetes, body mass index, and smoking status.⁷ In validating this risk score in the EPIC-Norfolk study, Rahman et al reported that the model performed well in distinguishing cases of later type 2 diabetes (discrimination), with an area under the receiver operating characteristics curve of 0.75 (where 1 indicates perfect discrimination and 0.5 indicates no discrimination).⁷ The Framingham offspring type 2 diabetes risk score also makes use of routinely collected clinical data and biochemical measures including age, sex, parental history of type 2 diabetes, body mass index, high density lipoprotein cholesterol, triglycerides, and fasting glucose. This model also provided good discrimination of incident cases of diabetes, with an area under the receiver operating characteristics curve of 0.85.⁸ Notably, the addition of other non-routinely collected measures of glucose metabolism such as the homeostasis model assessment, fasting insulin, or two hour oral glucose tolerance test did not improve the risk prediction.⁸

Both risk algorithms include information on family history of type 2 diabetes. However, family history reflects shared environment as well as genotype, and whereas biochemical and non-biochemical features associated with type 2 diabetes alter with age, genetic predisposition is determined at conception and fixed through life. Identifying genotypes that alter susceptibility to type 2 diabetes, and using them on their own or in combination with existing phenotype based risk algorithms, might therefore improve prediction of risk, particularly if genotypes identify differences in the activity of pathways unmeasured in non-genetic risk models.

Large case-control studies using whole genome arrays of single nucleotide polymorphisms (genome-wide association studies) have identified multiple common risk alleles for type 2 diabetes, many of which reside in or near genes not previously implicated in the pathogenesis of diabetes,⁹⁻¹² in addition to confirming loci identified by previous candidate gene studies.¹³⁻¹⁴ Around 20 single nucleotide polymorphisms associated with type 2 diabetes have been identified, for which the findings from the discovery study have been independently replicated in additional datasets and the effect sizes evaluated precisely by meta-analysis (as detailed in web table A).¹⁵⁻¹⁸ For all loci identified to date, effect sizes have been modest, with odds ratios for risk of type 2 diabetes ranging from 1.37 for single nucleotide polymorphisms in the gene *TCF7L2* to 1.09 for those in the gene *ADAMTS9*,¹⁸ raising the question of whether information on genotype would be useful for prediction of risk. Moreover, risk alleles have been discovered mainly by using case-control studies, which, although efficient for the discovery of disease associated single nucleotide polymorphisms, are suboptimal for evaluating their performance as predictors of risk. This is optimally assessed

by genotyping the genetic markers in prospective studies of initially healthy people in whom cases of incident disease accrue over time. However, the study designs, the number of disease associated single nucleotide polymorphisms evaluated, the metrics used to evaluate predictive performance, and the methods used to compare the performance of genotype with non-genetic risk factors have been variable. Despite this, genetic tests for the prediction of type 2 diabetes have already become available in the commercial sector (for example, www.23andme.com and www.decodedme.com/).

In this study, we first compared the ability of two phenotype based risk algorithms, the Cambridge risk score and the Framingham offspring risk score,^{7,8} with genotype based risk models for the discrimination of incident cases of type 2 diabetes in the prospective Whitehall II study of civil servants in the United Kingdom. We then evaluated the effect of addition of genotypes to the two non-genetic risk algorithms on the ability to discriminate between people who developed type 2 diabetes and those who remained free of the disease. We further assessed the effect of adding genotype to the calibration of the risk models (which involves assessing the closeness of the observed event rate to the predicted risk in different risk categories) as well as reclassification (which assesses the extent to which addition of genetic information shifts people to categories of predicted risk that better reflect their eventual outcome).¹⁹

METHODS

Study population

Between 1985 and 1988, all civil servants aged between 35 and 55 years in 20 departments in central London were invited to a medical examination at their workplace.^{20,21} With 73% participation, the cohort included 10 308 participants at entry to the study. At phase 3 in 1991-3, all participants known to be alive and in the country were invited to the screening clinic, which included an oral glucose tolerance test; 6058 men and 2758 women (85.5% of the original sample) attended. For this analysis, phase 3 (when an oral glucose tolerance test was done for the first time) serves as the baseline. Biochemical screening was repeated at phase 5 (1997-9) and phase 7 (2003-4). Additional questionnaire-only phases assessed diabetes status at phase 4 (1995-6) and phase 6 (2001). Of the baseline participants, 6156 participated in phase 7 screening, at which DNA was collected.

Clinical characteristics

We measured age, sex, body mass index, waist circumference, blood pressure, lipids, C reactive protein, fibrinogen, glucose, smoking, and coronary heart disease and diabetes status at baseline. Weight was measured in underwear to the nearest 0.1 kg on Soehnle electronic scales. We measured height in bare feet to the nearest 1 mm by using a stadiometer with the participant standing erect with head in the Frankfort plane. We calculated body mass index as weight (kilograms)/

height (metres) squared. We measured waist circumference, taken as the smallest circumference at or below the costal margin, with participants unclothed in the standing position by using a fibreglass tape measure at 600 g tension. We measured systolic blood pressure and diastolic blood pressure twice in the sitting position after five minutes' rest with the Hawksley random zero sphygmomanometer. We took the average of the two readings to be the measured blood pressure. We took venous blood in the fasting state or at least five hours after a light, fat free breakfast, before a two hour 75 g oral glucose tolerance test was done. Serum for lipid analyses was refrigerated at -4°C and assayed within 72 hours. We used a Cobas Fara centrifugal analyser (Roche Diagnostics System, Nutley, NJ) to measure cholesterol and triglyceride concentrations. We measured high density lipoprotein cholesterol by precipitating non-high density lipoprotein cholesterol with dextran sulfate-magnesium chloride with the use of a centrifuge and measuring cholesterol in the supernatant fluid. We used the Friedewald formula to calculate low density lipoprotein cholesterol concentration. We measured C reactive protein in serum stored at -70°C with a high sensitivity immunonephelometric assay in a BN ProSpec nephelometer (Dade Behring, Milton Keynes). We measured fibrinogen by an automated Clauss assay in an MDA-180 coagulator (Organon Teknika, Cambridge), using the manufacturer's reagents and the international fibrinogen standard. We measured blood glucose by the glucose oxidase method on a YSI Model 23A glucose analyser,^{22,23} and serum insulin with an in-house human insulin radioimmunoassay.²⁴ We defined prevalent coronary heart disease as meeting MONICA criteria,²⁵ giving positive responses to questions about chest pain,²⁶ physician's diagnosis, evidence from medical records, or positive electrocardiographic findings.

Diabetes, HbA_{1c}, and homoeostasis model assessment

We determined diabetes status on the basis of a 75 g oral glucose tolerance test (phases 3, 5, and 7), use of anti-diabetic drugs, or self report of a doctor's diagnosis (phases 3 to 7). We defined diabetes by a two hour glucose of at least 11.1 mmol/l or fasting glucose of at least 7 mmol/l.²⁷ Glycated haemoglobin (HbA_{1c}) was measured at phase 7 in EDTA whole blood on a calibrated HPLC system with automated haemolysis before injection. HbA₁ is resolved as a separate peak, which does not interfere with quantification of HbA_{1c}.²⁸ We calculated homoeostasis model assessment as (fasting glucose (mmol/l) $\times$ fasting insulin (mU/l)/22.5).²⁹ Non-fasting participants (fasting less than five hours) were assigned a missing value (n=435, 9.1%).

Genotyping

We extracted DNA from blood samples by using magnetic bead technology (Medical Solutions, Nottingham). Medical Solutions used SNPLex (Applied Biosciences) to determine the following single nucleotide polymorphisms, chosen because their association with diabetes risk has been replicated and the effect size

has been estimated with precision from large scale meta-analysis (web table A): *ADAMTS9* rs4607103, *CALPN10* rs3792267, *CDC123* (*CAMK1D*) rs12779790, *CDKN2B* rs10811661, *FTO* rs1421085 (which is complete LD with rs9930506³⁰), *HHEX* rs1111875, *HNF1A* (*TCF2*) rs1800574, *IGF2BP2* rs440296, *JAZF1* rs864745, *KCNJ11* rs521, *NOTCH2* rs10923931, *PPARG* rs1801282, *SLC30A8* rs13266634, *TCF2* (*HNF1B*) rs4430796, *TCF7L2* rs12255372, *THADA* rs7578597, and *TSPAN8* (*LGR5*) rs7961581. Single nucleotide polymorphisms in *BCL11A* rs10490072, *CDKAL1* rs17036101, *TCF7L2* rs7901695, and *VEGFA* rs9472138 were genotyped by using the 50K IBC CVD chip (web table B).³¹ All single nucleotide polymorphisms were in Hardy-Weinberg equilibrium (χ^2 , $P>0.05$).

Statistical analysis

We used two sample *t* tests to compare mean baseline values of continuous variables in people who developed diabetes and those who did not. Where appropriate, we log transformed variables and present geometric means and approximate standard deviations. We used the χ^2 test to compare categorical variables. We assessed the association of each genotype with risk of diabetes by logistic regression analysis and summarised the data by odds ratios and 95% confidence intervals. We used published regression coefficients to calculate the Cambridge type 2 diabetes risk score and Framingham offspring study type 2 diabetes risk score for each participant.^{7,8} In addition, we calculated two genetic scores. In the first, we assigned each person a score based simply on the number of risk alleles carried. Thus for *CDKAL1*, *CDC123*/*CAMK1D*, *FTO*, *HNF1A*, *IGFBP2*, *KCNJ11*, *NOTCH2*, *TCF2*, *TCF7L2*, *TSPAN8*/*LRG5*, and *VEGFA*, we coded genotypes "0" for common allele homozygotes,¹¹ "1" for heterozygotes, and "2" for rare allele homozygotes,²² and for *ADAMTS9*, *BCL11A*, *CALPN10*, *CDKN2A/2B*, *HHEX*, *JAZF1*, *PPARG*, *SLC30A8*, and *THADA*, coding was "2" for common allele homozygotes and "0" for rare allele homozygotes,¹¹ as the rare allele is reported to be protective (see web table B). In the second score, we calculated a genetic risk function by using weights derived from the risk coefficient for each gene based on odds ratios reported in previous meta-analyses (web table A).^{15 16 18 32} Risk estimates for each allele were available for 18 genes, and we multiplied these coefficients by 0, 1, or 2 according to the number of risk alleles carried by each person. Where effect estimates were reported for carriage of either one or two copies of each risk allele as a single group (*CALPN10* and *HNF1A*), we multiplied risk coefficients by a score of 0 or 1. We assumed genetic and clinical variables to be independent and added the weighted genetic score to each of the risk algorithms to provide a combined phenotypic and genetic score.

We assessed discrimination with the detection rate, which is equivalent to sensitivity and defined as the

proportion of all cases detected for a pre-specified false positive rate, as well as the area under the receiver operating characteristics curve. We assessed the calibration of the Cambridge risk score and Framingham offspring risk score in the estimation of the absolute risk of type 2 diabetes by comparing the difference between observed and expected event rates in different categories of risk with the Hosmer-Lemeshow test, with Akaike's information criterion and the likelihood ratio test as global measures of model fit.³³ We used the net reclassification improvement measure to assess the extent to which adding the genetic variables reassigned people to risk categories that better reflected their final outcome.³⁴

RESULTS

Web table C shows the baseline characteristics of the study participants. We excluded 103 people with prevalent type 2 diabetes at baseline from subsequent analysis. Of the 5135 participants followed for a median of 11.7 years to phase 7, with complete phenotype and genotype measures, 302 had developed type 2 diabetes by phase 7.

Discrimination using phenotype based risk scores

The odds ratio for developing diabetes was 8.4 (95% confidence interval 5.1 to 13.9) for people in the top fifth of the Cambridge risk score distribution compared with those in the bottom fifth, and the area under the receiver operating characteristics curve was 0.72 (95% confidence interval 0.69 to 0.76) at phase 7 (table 1). The odds ratio for type 2 diabetes in participants in the top fifth of the Framingham offspring risk score distribution compared with those in the bottom fifth was 18.2 (9.8 to 33.8), with a significantly greater area under the receiver operating characteristics curve of 0.78 (0.75 to 0.82) ($P=0.01$) (table 1 and fig 1). Detection rates (sensitivity) for a 5% false positive rate were 19.7 (95% confidence interval 14.1 to 25.3) for the Cambridge risk score (cut-off point 0.34) and 30.6 (24.1 to 37.1) for the Framingham offspring risk score (cut-off point 0.08). The respective detection rates for a 10% false positive rate were 34.2 (27.5 to 40.9; cut-off point 0.26) and 43.0 (36.0 to 50.0; cut-off point 0.05) (table 1).

Discrimination using genotype based risk scores

Web table B shows details of the 20 single nucleotide polymorphisms typed, together with risk estimates for type 2 diabetes from published meta-analyses (web table A). All genotypes were in Hardy-Weinberg equilibrium. In line with previous reports, *FTO* rs1421085 was associated with body mass index ($P<0.001$).³⁰ *CDC123/CAMK10* rs10811661 ($P=0.001$), *KCNJ11* rs5219 ($P<0.01$), and *TCF7L2* rs7901695 ($P<0.002$) were associated with two hour glucose after an oral glucose tolerance test. *CALPN10* rs3792267 was associated with significant differences in concentrations of total cholesterol and low density lipoprotein cholesterol ($P=0.01$). We found no other associations between genotypes and phenotypes included in either

of the non-genetic risk models at phase 3 or phase 7 (web tables D-J). The point estimates of the effect sizes were consistent with previous meta-analyses involving many thousands of cases of type 2 diabetes (web table A). However, as would be expected in a prospective study with fewer cases of diabetes and in agreement with the previous report from a prospective study of similar size,³⁵ most associations were not significant at $P<0.05$ in this dataset.

Figure 2 shows the distribution of the gene count score in people who developed type 2 diabetes and those who remained free of diabetes; the web figure shows the equivalent data for genetic risk function. Fourteen per cent of people with type 2 diabetes were in the top fifth of the gene count score compared with 13.9% of those without type 2 diabetes (web table K). The odds ratio for type 2 diabetes for participants in the top fifth of the score was 1.6 (0.9 to 2.5) compared with those in the bottom fifth. Twenty-five per cent of people with type 2 diabetes were in the top fifth of the genetic risk function, compared with 20% of those without diabetes, giving an odds ratio for type 2 diabetes of 2.3 (1.5 to 3.8) (web table K). The mean gene count score was 21.1 (SD 2.6) in people with type 2 diabetes and 20.5 (2.7) in those without type 2 diabetes (web table K). The simple gene count and the genetic risk function gave very similar discrimination, with areas under the receiver operating characteristics curve of 0.54 (0.50 to 0.58) (fig 1) and 0.55 (0.51 to 0.59) (table 1). Table 1 shows the detection rates at 5% and 10% false positive rates.

Effect of adding genetic information to discrimination and calibration of risk models

The addition of genetic information to either of the phenotype based risk models (using the simple gene

Fig 1 | Receiver operating characteristics curves for gene count score alone (area under curve 0.54, 95% CI 0.50 to 0.58), Framingham offspring risk score (area under curve 0.78, 0.75 to 0.82), and gene count score incorporated into Framingham offspring risk score (area under curve 0.81, 0.75 to 0.82)

Table 1 Odds ratios and area under receiver operating characteristics curve for Cambridge type 2 diabetes risk score, Framingham offspring type 2 diabetes risk score, and genetic risk scores alone and in combination in Whitehall II study

Risk model	Odds ratio* (95% CI)	Area under receiver operating characteristics curve (95% CI)	Detection rate for 5% false positive rate	Detection rate for 10% false positive rate
Cambridge risk score	8.4 (5.1 to 13.9)	0.72 (0.69 to 0.76)	19.7 (14.1 to 25.3)	34.2 (27.5 to 40.9)
Framingham offspring risk score	18.2 (9.8 to 33.8)	0.78 (0.75 to 0.82)	30.6 (24.1 to 37.1)	43.0 (36.0 to 50.0)
Gene count score	1.7 (0.9 to 2.5)	0.54 (0.50 to 0.58)	6.5 (3.5 to 10.8)	9.9 (6.2 to 14.9)
Genetic risk function score	2.3 (1.5 to 3.8)	0.55 (0.51 to 0.59)	6.4 (3.5 to 10.8)	11.4 (7.4 to 16.6)
Cambridge risk score and gene count score	9.7 (5.4 to 17.4)	0.73 (0.69 to 0.76)	20.3 (15.0 to 26.5)	33.7 (27.2 to 40.6)
Cambridge risk score and genetic risk function score	7.9 (4.5 to 13.7)	0.73 (0.69 to 0.76)	21.8 (16.3 to 28.1)	32.2 (25.8 to 39.1)
Framingham offspring risk score and gene count score	11.2 (6.1 to 20.5)	0.78 (0.75 to 0.81)	29.7 (23.5 to 36.5)	42.6 (35.7 to 49.7)
Framingham offspring risk score and genetic risk function score	11.0 (6.0 to 20.1)	0.78 (0.74 to 0.81)	31.2 (24.9 to 38.1)	40.1 (33.3 to 47.2)

*Odds ratio for top versus bottom fifth.

count or the genetic risk function) did not improve discrimination (table 1). Because part of the information included in the family history component of a risk score could reflect carriage of common genotypes, this may have undermined the incremental value of genetic information for predicting risk. However, the variants we studied explain only a small proportion of the familial aggregation of diabetes, and a formal analysis of the effect of inclusion or exclusion of the family history variable had almost no effect on the area under the receiver operating characteristics curve (data available on request).³⁶

The addition of genetic information did little to alter the accuracy of the phenotype based models when assessed with the Hosmer-Lemeshow test of calibration (table 2). Although the improvement in fit of the model was statistically significant when the Cambridge risk score was supplemented by the genetic risk function, the magnitude of the change was very small and unlikely to have any worthwhile clinical impact.

We generated four 10 year risk categories for type 2 diabetes for each of the two phenotype based risk models ($\leq 5\%$, 5-9.9%, 10-14.9%, and $\geq 15\%$) and then compared the observed and predicted event rates in each category of risk to assess if the addition of genetic information improved calibration (that is, the closeness of the observed event rate to the predicted risk) (table 3). The phenotype based risk models accurately estimated the rates of diabetes in each of the four categories of predicted risk.

Table 2 Effect of adding genetic information to global model fit and calibration of risk models to examine indices of model fit

Risk model	Hosmer-Lemeshow P value	Akaike's information criterion	Likelihood ratio test P value
Cambridge risk score	0.77	1416.1	NA
Framingham offspring risk score	0.42	1319.7	NA
Cambridge risk score plus gene count score	0.85	1414.9	0.07
Framingham offspring risk score plus gene count score	0.52	1319.7	0.17
Cambridge risk score plus genetic risk function score	0.50	1413.4	0.03
Framingham risk score plus genetic risk function score	0.55	1318.9	0.10

NA=not applicable.

Reclassification

We used the net reclassification index to assess the extent to which adding genotype data to the phenotype based risk models resulted in the movement of participants into risk categories that more accurately reflected their eventual outcome. In these analyses, we used the same four risk categories ($\leq 5\%$, 5-9.9%, 10-14.9%, and $\geq 15\%$) and did the analyses separately for people diagnosed as having type 2 diabetes on follow-up and those who remained healthy. Among the eventual cases, a shift to a higher risk category on the addition of genetic information is classed as a correct reclassification and a downward shift as an incorrect reclassification. The opposite is the case for people who remained healthy. The addition of the simple gene count to the Cambridge risk score resulted in a net reclassification improvement of -1.1% (95% confidence interval -6.0% to 3.8%) (table 4), and the addition of information from the genetic risk function produced a net reclassification improvement of 4.6% (-2.0% to 11.3%) (P=0.17) (web table L). Neither the simple gene count score (net reclassification improvement 0.2%, -5.1 to 4.7) (table 5) nor the genetic risk function (net reclassification improvement -3.2%, -9.9 to 3.5) (web table M) resulted in a net reclassification improvement when added to the Framingham offspring risk score.

Fig 2 Percentage of participants in each gene count score category among those who developed type 2 diabetes and those who remained free from diabetes. Risk of developing diabetes according to gene count shown as fitted line from regression analysis

Table 3 | Prevalence and percentage of incident cases of type 2 diabetes in categories of predicted risk calculated by using phenotype based risk models alone, and same models with addition of genetic data

	Category of predicted risk				Proportion reclassified (%)
	<5%	5-9.9%	10-14.9%	>15%	
Cambridge risk score					
Without genotype scores	64/2257 (2.8)	54/753 (7.2)	48/328 (14.6)	36/188 (19.2)	NA
With simple gene count score	63/2226 (2.8)	59/787 (7.5)	41/317 (12.9)	39/196 (19.9)	322/3526 (9.9)
With genetic risk function	57/2191 (2.6)	61/824 (7.4)	42/309 (13.6)	42/202 (20.8)	522/3526 (16.0)
Framingham offspring risk score					
Without genotype scores	60/2403 (2.5)	42/603 (7.0)	31/261 (11.9)	69/259 (26.6)	NA
With simple gene count score	54/2385 (2.3)	49/598 (8.2)	32/265 (12.1)	67/278 (24.1)	210/3526 (6.0)
With genetic risk function	55/2381 (2.3)	54/614 (8.8)	26/255 (10.2)	67/276 (24.3)	432/3526 (12.3)

NA=not applicable.

DISCUSSION

In a British cohort (the Whitehall II study), a panel of 20 genotypes associated with type 2 diabetes performed less well than the Cambridge and Framingham offspring type 2 diabetes risk scores in discriminating incident cases of type 2 diabetes. Adding the genetic panel to the phenotype based risk models did not improve discrimination and produced only minimal improvement in accuracy of risk estimation assessed by recalibration and, at best, a minor net reclassification improvement.

Over the past five years, the pace of identification of genetic loci underlying susceptibility to common diseases has increased rapidly, leading to interest in how this information might best be used to improve personal and public health. One potential application is the use of genetic information to help predict susceptibility to disease in initially healthy people, so as to focus preventive interventions on those at the highest risk of future disease. This targeted approach to prevention is exemplified by the established use of risk equations based on non-genetic variables to estimate risk of coronary heart disease and guide blood pressure lowering and cholesterol lowering treatment.³⁴ This approach to the prevention of vascular disease, for which diabetes is a major risk factor, will become more systematic in the next two years, through the Department of Health's vascular health check scheme (www.dh.gov.uk/en/

Publicationsandstatistics/Publications/PublicationsPolicyAndGuidance/DH_083822).

Preventive interventions also exist for type 2 diabetes, which motivated the recent evaluation of risk scores (including those studied here) for the prediction of type 2 diabetes. The Cambridge risk score and the Framingham offspring risk score are based on a combination of demographic, family history, anthropometric, and biochemical data, but neither includes genetic information.⁷⁸ Although these phenotype based risk models seem to perform well, an important question is whether typing a panel of validated genetic risk factors might improve their ability to predict type 2 diabetes. Some studies in this area have used case-control datasets.³⁷⁻³⁹ Although efficient for gene discovery, these are a suboptimal design for evaluating the predictive performance of a marker, as risk information is available only in relative terms and the range of metrics that can be derived to assess predictive performance is more limited than for prospective studies with incident cases of disease. Those prospective studies that have previously evaluated the performance of genetic markers have been set outside the UK, typed fewer type 2 diabetes risk alleles, or reported only some of a range of metrics available to evaluate the performance of a predictive test (table 6).³⁵⁻⁴⁰⁻⁴² In a prospective study set in the UK, we therefore tested the performance of a panel of 20 common genes associated with type 2 diabetes,

Table 4 | Net reclassification improvement based on addition of gene count score to Cambridge risk score

Predicted diabetes risk (Cambridge risk score)	No of people				Reclassified		Net correctly reclassified (%)
	<5%	5-10%	10-15%	>15%	Increased risk	Decreased risk	
Plus gene count score—people without diabetes during follow-up (n=3324)							
<5%	2088	105	0	0	167	131	-1.1
5-9.9%	75	581	43	0			
10-14.9%	0	42	219	19			
≥15%	0	0	14	138			
Plus gene count score—people with diabetes during follow-up (n=202)							
<5%	60	4	0	0	12	12	0
5-9.9%	3	49	2	0			
10-14.9%	0	6	36	6			
≥15%	0	0	3	33			

Net reclassification improvement -1.1% (95% CI -6.0% to 3.8%); P=0.66.

Table 5 | Net reclassification improvement based on addition of gene count score to Framingham offspring risk score

Predicted diabetes risk (Framingham offspring risk score)	No of people				Reclassified		Net correctly reclassified (%)
	<5%	5-10%	10-15%	>15%	Increased risk	Decreased risk	
Plus gene count score—people without diabetes during follow-up (n=3324)							
<5%	2295	48	0	0	121	64	-1.7
5-9.9%	36	482	43	0			
10-14.9%	0	19	181	30			
≥15%	0	0	9	181			
Plus gene count score—people with diabetes during follow-up (n= 202)							
<5%	52	8	0	0	14	11	1.5
5-9.9%	2	37	3	0			
10-14.9%	0	4	24	3			
≥15%	0	0	5	64			
Net reclassification improvement 0.2% (95% CI -5.1 to 4.7); P=0.94.							

each of which confers a small to moderate increase in the risk of type 2 diabetes, and compared prediction based on genetic information alone, phenotypic information alone, and both, by using a range of metrics to assess predictive performance.

We found that risk functions based on routinely measured clinical variables better discriminated incident type 2 diabetes cases than did a panel of 20 diabetes associated single nucleotide polymorphisms. The inclusion of genetic information in the risk models did not improve the discrimination of cases of type 2

diabetes, and nor did it provide clinically important improvement in the accuracy of these models when assessed by calibration. The addition of genetic data to phenotype based risk models also provided minimal net reclassification improvement. The addition of genetic information resulted in the reassortment of people into different risk categories, but not all the shifts were helpful. Although some eventual cases were upgraded to higher risk categories, almost as many had their risk downgraded, and the opposite was true for many of those who remained healthy.

Table 6 | Comparison of published studies that have used genetic information with or without non-genetic risk factors to discriminate between people with and without type 2 diabetes, in case-control, cross sectional, or prospective settings

Study	Study design	No of cases	Non-genetic risk factors	No of risk alleles studied	Other metrics of prediction reported
Including Diabetes UK Warren 2 repository and people with early onset type 2 diabetes and 1958 birth cohort as controls ⁴⁴	Case-control	2406 type 2 diabetes, 3668 controls	None	3	None
DESIR ³⁷	Prospective	187 type 2 diabetes, 336 impaired glucose tolerance, 2732 non-type 2 diabetes	Age, sex, body mass index	3	None
DESIR plus DIAB2.NEPHRO-GENE and people with normal glucose tolerance from SU.VI.MAX ³⁸	Case-control	4232 type 2 diabetes, 4595 glucose tolerant	Age, sex, body mass index	15	None
Rotterdam Study ³⁹	Prospective	686 type 2 diabetes, 5221 non-type 2 diabetes	Age, sex, body mass index	18	None
Framingham Offspring Study ³⁵	Prospective	255 type 2 diabetes, 2122 non-type 2 diabetes	Age, sex, family history of type 2 diabetes, body mass index, fasting glucose, systolic blood pressure, triglycerides, high density lipoprotein cholesterol	18	Net reclassification index
Malmö Preventive Project and Botnia Study ⁴⁰	Prospective	2063 type 2 diabetes, 12 210 non-type 2 diabetes; 138 type 2 diabetes, 2632 non-type 2 diabetes	Age, sex, family history of type 2 diabetes, body mass index, blood pressure, triglycerides, fasting glucose	16	Integrated discrimination improvement
GoDARTS41	Case-control	2309 cases, 2598 controls	Age, sex, body mass index	18	None
Co-Laue ⁴²	Cross sectional	356 type 2 diabetes, 5004 non-type 2 diabetes	Age, body mass index, family history of type 2 diabetes, waist: hip ratio, triglyceride:high density lipoprotein cholesterol ratio, physical activity	15	Integrated discrimination improvement
Inter99 study, ADDITION study, plus type 2 diabetes cases and people with normal glucose tolerance ⁴³	Cross sectional	4093 type 2 diabetes, 5302 glucose tolerant	Age, sex, body mass index	19	None

See Forrest plots at www.ucl.ac.uk/genetic-epidemiology/WebMaterial.

Our findings are consistent with the nine previous published reports of 10 study populations (table 6),^{35 37 38 40-44} even though the number and range of genotypes and the phenotype based risk models used for prediction varied across studies (see Forrest plots at www.ucl.ac.uk/genetic-epidemiology/WebMaterial). All models included age, body mass index, and sex, and much of the predictive information in any phenotype based model is likely to be encompassed in these terms.

The relations shown in figure 2 and the web figure illustrate one reason for the poor predictive performance of a panel of single nucleotide polymorphisms associated with common diseases. Although people carrying multiple risk alleles are at more extreme risk of type 2 diabetes than those carrying fewer copies, they represent only a small proportion of the population, because the inheritance of each risk allele is an independent event—the probability of inheriting multiple risk alleles is a function of the frequency of each allele in the population. For example, the probability of inheriting 10 independent risk alleles with frequencies around 0.3 is 0.3^{10} (about 6×10^{-6}). People with an intermediate number of risk alleles would therefore be expected to account for the major portion of cases of type 2 diabetes, because of the large number of people at intermediate risk in the population. This explains the substantial overlap of the distribution of risk alleles among people who developed diabetes and those who remained disease-free, which makes it difficult to set a cut-off point of a gene count (or genetic risk function) that reliably discriminates later cases of type 2 diabetes. Although genetic tests for type 2 diabetes, based on a subset of the alleles studied here, can already be purchased in the commercial sector, our findings suggest that much more rigorous evaluation of their use as a health technology is needed before such tests should be adopted by healthcare organisations.

As a technology, however, genotype based tests have several inherent advantages over non-genetic tests. Genotype based assays are cheap, have high fidelity, and can be multiplexed, in contrast to multiple phenotypic risk factors and biomarkers, many of which require different methods for their measurement, and which are more affected by biological variability and measurement error than is genotyping. Moreover, because genotype is invariant it offers the prospect of risk assessment from much earlier in life than is possible with phenotype based tests. In the case of cardiovascular risk factors, evidence shows that greater benefits accrue from earlier intervention among people at higher risk (for example, in the form of smoking cessation or cholesterol lowering).^{45 46} The findings of our study should thus not lead to the premature dismissal of genotype based risk prediction as a health technology. Rather, increased efforts should be made to understand the strengths and limitations of such tests as well as their optimal place in health care, a conclusion highlighted in the recent House of Lords Science and Technology Committee's report on genomic medicine (www.publications.parliament.uk/pa/ld200809/ldselect/ldstech/107/107i.pdf).

Limitations of study

Some limitations of our study should be noted. Although prospective, the Whitehall II study is workplace based and therefore not necessarily representative of the general population. However, the excellent performance in Whitehall II of the non-genetic risk functions for type 2 diabetes, both of which were developed and validated in general populations, suggests that this is unlikely to bias our conclusions substantially. Moreover, our findings are consistent with those of prospective studies set in representative general populations. Our findings are also not generalisable to people of non-European ancestry, who we excluded from this analysis. Although DNA was collected some time after baseline, which could have introduced a survivor bias, we think that this is unlikely to have affected our results given the modest effect of the alleles we studied on risk of diabetes and the long natural history of the development of the life threatening complications of diabetes.

The two risk tools studied, based on non-genetic markers, performed better than genotype based tests despite the fact that the models, which were developed in different datasets, were not specifically recalibrated for the Whitehall II population. The common diabetes associated single nucleotide polymorphisms we studied might have greater incremental value in the prediction of type 2 diabetes when evaluated against some of the other validated risk models. However, we chose the Framingham and Cambridge risk scores because they are contemporary (which could be important, given the recent increase in the incidence and prevalence of type 2 diabetes), were developed in populations with a similar profile to the Whitehall II participants, and were based on studies set in the United States and the UK, where many of the genetic studies were done. Moreover, both include variables that are routinely measured in clinical practice. We did not evaluate QDRisk, which is based on routinely collected primary care data (including deprivation scores, ethnicity, and current drug treatment for hypertension or cardiovascular disease and corticosteroid use), which was reported during the preparation of this manuscript.⁴⁷

Because part of the information included in the family history component of a risk score will reflect common genotypes, this may have undermined the incremental value of genetic information for risk prediction. However, the variants we studied explain only a small proportion of the familial aggregation of diabetes.³⁶ Whether genotypes have greater predictive utility in particular categories of patient (such as among leaner people or those of a particular ancestry) could be assessed by pooling participant level data from a large number of prospective studies with the relevant information to ensure adequate power. Our current analysis is limited to the 20 common risk alleles for type 2 diabetes identified by large association or genome-wide studies. However, sequence variants of intermediate frequency but larger effect size are likely to be uncovered by future research, so our interpretation on the

WHAT IS ALREADY KNOWN ON THIS TOPIC

Several routinely used anthropometric and biological measures are included in the validated Cambridge and the Framingham offspring type 2 diabetes risk models

Common single nucleotide polymorphisms associated with susceptibility to type 2 diabetes have been identified from whole genome and candidate gene association studies

The extent to which a comprehensive panel of genotypes will help in predicting incident diabetes in the UK is not known

WHAT THIS STUDY ADDS

A panel of 20 type 2 diabetes associated genotypes performed less well than the Cambridge and Framingham offspring type 2 diabetes risk scores in discriminating incident cases of type 2 diabetes

Including the genetic panel did not improve discrimination, calibration, or reclassification when added to the Cambridge or Framingham risk models

The major translational application of currently known common type 2 diabetes associated genotypes is likely to arise from the insight they provide on causes of disease and therapeutic targets

predictive utility of genotype should be regarded as interim. Moreover, as the actual causal variants at each gene/region remain for the most part uncertain, the predictive utility of genetic markers may also have been underestimated.

Our conclusions about the performance of genetic testing for type 2 diabetes are confined to the use of single common alleles at each locus. Other common risk alleles are likely to exist at the same genetic loci (including the causal variants), which could provide additional information relevant to prediction. Our conclusions are also not transferrable to other common diseases. For example, genetic variants underlying the susceptibility to age related macular degeneration have been identified, at least one of which is both common and large in its effect on risk.⁴⁸ We previously examined the predictive utility of a common single nucleotide polymorphism associated with the risk of coronary heart disease at the 9p23.1 chromosomal locus (rs10757274) when added to a risk function that included variables incorporated in the Framingham coronary heart disease risk equation.⁴⁹ Although this genotype added minimally to the ability of the Framingham risk score to discriminate future events, improving the area under the receiver operating characteristics curve by only 3%, it did significantly improve reclassification of risk of coronary heart disease, albeit modestly. Moreover, for some disorders, including age related macular degeneration, few if any non-genetic biomarkers or risk factors exist that can be used to estimate risk of future disease.

Conclusions

Phenotype based risk models (the Framingham offspring and Cambridge risk scores) provided greater discrimination for type 2 diabetes than did models based on 20 common independently inherited alleles associated with risk of type 2 diabetes. The addition of 20 common genotypes associated with modest risk to phenotype based risk models produced only minimal

improvement in the accuracy of risk estimation assessed by recalibration and at best a minor net reclassification improvement. The major translational application of the currently known common, small effect genetic variants influencing susceptibility to type 2 diabetes is likely to come from the insight they provide on causes of disease and potential therapeutic targets.

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Competing interests: ADH is on the editorial board of *Drug and Therapeutics Bulletin*, a BMJ Group publication, has provided non-remunerated advice to GlaxoSmithKline and London Genetics, and has received honorariums for speaking at educational meetings with sponsorship from the pharmaceutical industry, which have been donated in whole or large part to charity.

Ethical approval: The study was approved by the UCL/UCLH ethics committee, and all participants gave written informed consent.

Data sharing: Whitehall II data, protocols, and other metadata are available to the whole of the scientific community. Please refer to the Whitehall II data sharing policy on www.ucl.ac.uk/whitehallII/data_sharing/index.htm.

- 1 Bonora E, Kiechl S, Willeit J, Oberhollenzer F, Egger G, Meigs JB, et al. Population-based incidence rates and risk factors for type 2 diabetes in white individuals: the Bruneck study. *Diabetes* 2004;53:1782-9.
- 2 Mathers CD, Loncar D. Projections of global mortality and burden of disease from 2002 to 2030. *PLoS Med* 2006;3:e442.
- 3 Wilson PW, D'Agostino RB, Levy D, Belanger AM, Silbershatz H, Kannel WB. Prediction of coronary heart disease using risk factor categories. *Circulation* 1998;97:1837-47.
- 4 Assmann G, Cullen P, Schulte H. Simple scoring scheme for calculating the risk of acute coronary events based on the 10-year follow-up of the prospective cardiovascular Munster (PROCAM) study. *Circulation* 2002;105:310-5.
- 5 Hippisley-Cox J, Coupland C, Vinogradova Y, Robson J, Minhas R, Sheikh A, et al. Predicting cardiovascular risk in England and Wales: prospective derivation and validation of QRISK2. *BMJ* 2008;336:1475-82.
- 6 Gillies CL, Abrams KR, Lambert PC, Cooper NJ, Sutton AJ, Hsu RT, et al. Pharmacological and lifestyle interventions to prevent or delay type 2 diabetes in people with impaired glucose tolerance: systematic review and meta-analysis. *BMJ* 2007;334:299.
- 7 Rahman M, Simmons RK, Harding AH, Wareham NJ, Griffin SJ. A simple risk score identifies individuals at high risk of developing type 2 diabetes: a prospective cohort study. *Fam Pract* 2008;25:191-6.
- 8 Wilson PW, Meigs JB, Sullivan L, Fox CS, Nathan DM, D'Agostino RB Sr. Prediction of incident diabetes mellitus in middle-aged adults: the Framingham offspring study. *Arch Intern Med* 2007;167:1068-74.

- 9 Sladek R, Rocheleau G, Rung J, Dina C, Shen L, Serre D, et al. A genome-wide association study identifies novel risk loci for type 2 diabetes. *Nature* 2007;445:881-5.
- 10 Saxena R, Voight BF, Lyssenko V, Burt NP, de Bakker PI, Chen H, et al. Genome-wide association analysis identifies loci for type 2 diabetes and triglyceride levels. *Science* 2007;316:1331-6.
- 11 The Wellcome Trust Case Control Consortium. Genome-wide association study of 14,000 cases of seven common diseases and 3,000 shared controls. *Nature* 2007;447:661-78.
- 12 Scott LJ, Mohlke KL, Bonnycastle LL, Willer CJ, Li Y, Duren WL, et al. A genome-wide association study of type 2 diabetes in Finns detects multiple susceptibility variants. *Science* 2007;316:1341-5.
- 13 Horikawa Y, Oda N, Cox NJ, Li X, Orho-Melander M, Hara M, et al. Genetic variation in the gene encoding calpain-10 is associated with type 2 diabetes mellitus. *Nat Genet* 2000;26:163-75.
- 14 Scott LJ, Bonnycastle LL, Willer CJ, Sprau AG, Jackson AU, Narisu N, et al. Association of transcription factor 7-like 2 (TCF7L2) variants with type 2 diabetes in a Finnish sample. *Diabetes* 2006;55:2649-53.
- 15 Song Y, Niu T, Manson JE, Kwiatkowski DJ, Liu S. Are variants in the CAPN10 gene related to risk of type 2 diabetes? A quantitative assessment of population and family-based association studies. *Am J Hum Genet* 2004;74:208-22.
- 16 Weedon MN, Owen KR, Shields B, Hitman G, Walker M, McCarthy MI, et al. A large-scale association analysis of common variation of the HNF1alpha gene with type 2 diabetes in the U.K. Caucasian population. *Diabetes* 2005;54:2487-91.
- 17 Zeggini E, Weedon MN, Lindgren CM, Frayling TM, Elliott KS, Lango H, et al. Replication of genome-wide association signals in U.K. samples reveals risk loci for type 2 diabetes. *Science* 2007;316:1336-41.
- 18 Zeggini E, Scott LJ, Saxena R, Voight BF, Marchini JL, Hu T, et al. Meta-analysis of genome-wide association data and large-scale replication identifies additional susceptibility loci for type 2 diabetes. *Nat Genet* 2008;40:638-45.
- 19 Cook NR. Comments on "Evaluating the added predictive ability of a new marker: From area under the ROC curve to reclassification and beyond" by M. J. Pencina et al., *Statistics in Medicine* (DOI: 10.1002/sim.2929). *Stat Med* 2008;27:191-5.
- 20 Marmot MG, Smith GD, Stansfeld S, Patel C, North F, Head J, et al. Health inequalities among British civil servants: the Whitehall II study. *Lancet* 1991;337:1387-93.
- 21 Marmot M, Brunner E. Cohort profile: the Whitehall II study. *Int J Epidemiol* 2005;34:251-6.
- 22 Cooper GR. Methods for determining the amount of glucose in blood. *CRC Crit Rev Clin Lab Sci* 1973;4:101-45.
- 23 Alpert L. Instrument series: Model 23A Glucose Analyzer. *Lab World* 1976;27:8-13.
- 24 Hanning I, Home PD, Alberti KG. Measurement of free insulin concentrations: the influence of the timing of extraction of insulin antibodies. *Diabetologia* 1985;28:831-5.
- 25 Tunstall-Pedoe H, Kuulasmaa K, Amouyel P, Arveiler D, Rajakangas AM, Pajak A. Myocardial infarction and coronary deaths in the World Health Organization MONICA project: registration procedures, event rates, and case-fatality rates in 38 populations from 21 countries in four continents. *Circulation* 1994;90:583-612.
- 26 Rose GA, Blackburn H, Gillum RF, Prineas RJ. Cardiovascular survey methods. 2nd ed. WHO, 1982.
- 27 World Health Organization. Definition, diagnosis and classification of diabetes mellitus and its complications. WHO, 1999.
- 28 Hoshino T, Takahashi Y, Suzuki M. Application of high-performance liquid chromatography in establishing an accurate index of blood glucose control. *J Chromatogr* 1990;515:531-6.
- 29 Matthews DR, Hosker JP, Rudenski AS, Naylor BA, Treacher DF, Turner RC. Homeostasis model assessment: insulin resistance and beta-cell function from fasting plasma glucose and insulin concentrations in man. *Diabetologia* 1985;28:412-9.
- 30 Frayling TM, Timpson NJ, Weedon MN, Zeggini E, Freathy RM, Lindgren CM, et al. A common variant in the FTO gene is associated with body mass index and predisposes to childhood and adult obesity. *Science* 2007;316:889-94.
- 31 Keating BJ, Tischfield S, Murray SS, Bhangale T, Price TS, Glessner JT, et al. Concept, design and implementation of a cardiovascular gene-centric 50 k SNP array for large-scale genomic association studies. *PLoS ONE* 2008;3:e3583.
- 32 Perry JR, Frayling TM. New gene variants alter type 2 diabetes risk predominantly through reduced beta-cell function. *Curr Opin Clin Nutr Metab Care* 2008;11:371-7.
- 33 Hosmer DW, Lemeshow S. Confidence interval estimation of interaction. *Epidemiology* 1992;3:452-6.
- 34 Pencina MJ, D'Agostino RB Sr, D'Agostino RB Jr, Vasan RS. Evaluating the added predictive ability of a new marker: from area under the ROC curve to reclassification and beyond. *Stat Med* 2008;27:157-72.
- 35 Meigs JB, Shrader P, Sullivan LM, McAteer JB, Fox CS, Dupuis J, et al. Genotype score in addition to common risk factors for prediction of type 2 diabetes. *N Engl J Med* 2008;359:2208-19.
- 36 Lyssenko V, Groop L. Genome-wide association study for type 2 diabetes: clinical applications. *Curr Opin Lipidol* 2009;20:87-91.
- 37 Cauchi S, Meyre D, Durand E, Proenca C, Marre M, Hadjadj S, et al. Post genome-wide association studies of novel genes associated with type 2 diabetes show gene-gene interaction and high predictive value. *PLoS ONE* 2008;3:e2031.
- 38 Lango H, Palmer CN, Morris AD, Zeggini E, Hattersley AT, McCarthy MI, et al. Assessing the combined impact of 18 common genetic variants of modest effect sizes on type 2 diabetes risk. *Diabetes* 2008;57:3129-35.
- 39 Weedon MN, McCarthy MI, Hitman G, Walker M, Groves CJ, Zeggini E, et al. Combining information from common type 2 diabetes risk polymorphisms improves disease prediction. *PLoS Med* 2006;3:e374.
- 40 Vaxillaire M, Veslot J, Dina C, Proenca C, Cauchi S, Charpentier G, et al. Impact of common type 2 diabetes risk polymorphisms in the DESIR prospective study. *Diabetes* 2008;57:244-54.
- 41 Van Hoek M, Dehghan A, Witteman JC, van Duijn CM, Uitterlinden AG, Oostra BA, et al. Predicting type 2 diabetes based on polymorphisms from genome-wide association studies: a population-based study. *Diabetes* 2008;57:3122-8.
- 42 Lyssenko V, Jonsson A, Almgren P, Pulizzi N, Isomaa B, Tuomi T, et al. Clinical risk factors, DNA variants, and the development of type 2 diabetes. *N Engl J Med* 2008;359:2220-32.
- 43 Lin X, Song K, Lim N, Yuan X, Johnson T, Abderrahmani A, et al. Risk prediction of prevalent diabetes in a Swiss population using a weighted genetic score—the CoLaus study. *Diabetologia* 2009;52:600-8.
- 44 Sparso T, Garup N, Andreassen C, Albrechtsen A, Holmkvist J, Andersen G, et al. Combined analysis of 19 common validated type 2 diabetes susceptibility gene variants shows moderate discriminative value and no evidence of gene-gene interaction. *Diabetologia* 2009;52:1308-14.
- 45 Doll R, Peto R, Boreham J, Sutherland I. Mortality in relation to smoking: 50 years' observations on male British doctors. *BMJ* 2004;328:1519.
- 46 Baigent C, Keech A, Kearney PM, Blackwell L, Buck G, Pollicino C, et al. Efficacy and safety of cholesterol-lowering treatment: prospective meta-analysis of data from 90,056 participants in 14 randomised trials of statins. *Lancet* 2005;366:1267-78.
- 47 Hippisley-Cox J, Coupland C, Robson J, Sheikh A, Brindle P. Predicting risk of type 2 diabetes in England and Wales: prospective derivation and validation of QDScore. *BMJ* 2009;338:b880.
- 48 Thakkinian A, Han P, McEvoy M, Smith W, Hoh J, Magnusson K, et al. Systematic review and meta-analysis of the association between complement factor H Y402H polymorphisms and age-related macular degeneration. *Hum Mol Genet* 2006;15:2784-90.
- 49 Talmud PJ, Cooper JA, Palmen J, Lovering R, Drenos F, Hingorani AD, et al. Chromosome 9p21.3 coronary heart disease locus genotype and prospective risk of CHD in healthy middle-aged men. *Clin Chem* 2008;54:467-74.

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