

REVIEW

Challenges in reproducibility of genetic association studies: lessons learned from the obesity field

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A robust replication of initial genetic association findings has proved to be difficult in human complex diseases and more specifically in the obesity field. An obvious cause of non-replication in genetic association studies is the initial report of a false positive result, which can be explained by a non-heritable phenotype, insufficient sample size, improper correction for multiple testing, population stratification, technical biases, insufficient quality control or inappropriate statistical analyses. Replication may, however, be challenging even when the original study describes a true positive association. The reasons include underpowered replication samples, gene \times gene, gene \times environment interactions, genetic and phenotypic heterogeneity and subjective interpretation of data. In this review, we address classic pitfalls in genetic association studies and provide guidelines for proper discovery and replication genetic association studies with a specific focus on obesity.

International Journal of Obesity (2013) 37, 559–567; doi:10.1038/ijo.2012.82; published online 15 May 2012

Keywords: genetic epidemiology; false-positive association; replication; guidelines

INTRODUCTION

The prevalence of obesity has reached epidemic proportion across the world.¹ Genetic influence has a substantial role in this disease,² and has led scientists to search for the specific genetic determinants of human obesity. Candidate gene and linkage approaches have given rise to the discovery of several genes associated with monogenic or syndromic obesity, but provided limited success to identifying genetic variants that are associated with polygenic obesity.³ This situation was not exclusive to obesity, as most reported associations were not conclusive in human complex diseases.⁴ These observations raised concerns about the reliability of such genetic association studies, and high-impact publications such as the *Journal of Clinical Investigation* even stated in their editorial policy that genetic linkage and association studies related to complex disorders were unlikely to be accepted (<http://www.jci.org/kiosk/publish/policies>). This situation has changed recently with new high-throughput single-nucleotide polymorphism (SNP) genotyping array technologies enabling genome-wide association studies (GWAS),⁵ and with the establishment of international consortiums providing sample size large enough to identify gene variants with modest effect size (odds ratio (OR) < 2).⁶ More than 1000 loci have now been convincingly associated with human complex traits using GWAS and close to 70 gene variants have been associated with obesity-related traits at a stringent level of significance ($P < 5 \times 10^{-8}$).⁷ However, despite the remarkable progress made during the last decade in the genetic epidemiology field, several questions remain.⁸ How can we explain that despite promising original association reports, some genetic associations are widely replicated whereas others are not? When can we be confident that a positive finding is truly positive? How can we differentiate good genetic association studies from inadequate ones? We searched literature using the keywords 'genetic AND obesity' (11 152) or 'genetic association studies' (47 368) in the PubMed NCBI database from January 1987 to

February 2012. We selectively cited the more illustrative papers to address the classic pitfalls in genetic association studies and to provide guidelines for proper genetic association discovery and replication studies, specifically focused on obesity.

LACK OF REPLICATION DUE TO FALSE POSITIVE RESULT REPORTED IN ORIGINAL STUDY

Genetically complex diseases involve a large number of genetic, environmental factors and their interactions. In general, common risk alleles are less deterministic and more probabilistic than rare monogenic mutations, and only slightly increase the chance of disease. They are found in affected but also in unaffected people in populations.⁴ Finding an association between such a variant and disease is statistic by essence and the risk of reporting an initial positive association by chance can never be totally excluded. However, the risk of reporting a false positive association is more likely to occur in certain circumstances as listed below.

The phenotype is not heritable

Heritability reflects the proportion of total phenotypic variability caused by genetic variance in a population. When genetic variation refers to additive genetic effect only, heritability is named narrow-sense heritability or just heritability (h^2); when genetic variation includes all additive, dominant and epistatic genetic effects, heritability is defined as broad-sense heritability (H^2).⁹ Adoption and twin studies are the best study designs for heritability estimation because of their natural separation of genetic and environmental components, whereas family parent-offspring and sibling studies estimate less accurately the heritability owing to their partial similarity in genes and the same degree of environmental similarity.⁹

A phenotype with a substantial heritability is an important prerequisite to enable the identification of genetic determinants.¹⁰

Heritability studies have shown that obesity, body mass index (BMI) and BMI change are highly heritable traits (h^2 is 50–80%).^{2,11,12} Intermediary endophenotypes related to obesity, such as percentage of body fat,¹³ eating behavior¹⁴ and energy expenditure,¹⁵ have also a significant heritable component.

Insufficient sample size

A meta-analysis of 37 linkage studies for BMI did not detect significant linkage regions (LOD score > 3.6),¹⁶ suggesting that multiple susceptibility variants with small effect sizes may affect obesity predisposition. Evidence from the variants consistently associated with obesity to date confirms this view.⁷ To the rare exception of the *MC4R* I251L or *GPR120* R270H infrequent coding variants,^{17,18} polymorphisms associated with obesity display modest OR values typically lower than 1.50.⁷ This implies that large sample sizes are needed in case-control studies designed for gene discovery efforts (Table 1). Such large sample sizes have not been reported in obesity studies until recently,^{19,20} suggesting that most of the association studies reported before were underpowered to identify the effect sizes expected for obesity predisposing common variants. As small discovery studies tend to overestimate the true effect of gene variants,²¹ underpowered case-control studies are likely to prompt false positive associations and only few associations first reported from smaller studies prove to be reliable in larger replication data sets.²¹ The *GAD2* gene story is illustrative of this. In 2003, an initial study reported an association between the *GAD2* -243A > G promoter variant and obesity in a case-control design of 575 morbidly obese and 646 control French subjects.²² This sample was considered substantial at that time, but was far less than the sample requirements predicted by power calculation studies for an OR of 1.3 and a risk allele frequency of 0.17 in controls (Table 1). Logically, a replication effort in 680 class III obesity cases and 1186 lean controls of European ancestry did not replicate the original finding²³ and an adequately powered meta-analysis recapitulating the data of the two studies ($n = 3052$) did not find any association ($P = 0.28$).²³

Lack of proper correction for multiple testing

To account for multiple testing, researchers typically apply Bonferroni correction in which a threshold P of $0.05/n$ ($n =$ number of statistical tests performed in the study) is set. For instance, in a GWAS using 1×10^6 genetic markers, a threshold P of 5×10^{-8} is needed to declare an association at the corrected threshold of type I error of 5%. Testing multiple hypotheses in one study (multiple genetic markers, phenotypes and subgroup analyses) without proper adjustment for multiple comparisons increases the risk of a false positive report. In the original

Table 1. Sample sizes needed in a case-control design to detect significant association with a power of 90% and a two-sided P of 0.001 by odds ratio and allele frequency for risk allele

MAF in control	0.01	0.05	0.1	0.2	0.3	0.4
Allelic OR						
1.1	443 854	92 868	49 252	27 974	21 518	19 010
1.2	116 354	24 434	13 018	7460	5792	5162
1.3	54 110	11 404	6102	3526	2760	2480
1.5	21 208	4498	2426	1424	1132	1032
2.0	6386	1374	754	458	376	354

Abbreviations: MAF, minor allele frequency; OR, odds ratio. Calculations assume multiplicative effect on disease risk. Sample sizes presented are total number of cases and controls needed, assuming an equal number of cases and controls. The estimated sample sizes are derived assuming power = 90% and two-sided $P = 0.001$.

report showing an association between rs7566605 in *INSIG2* gene and BMI in 694 participants from the Framingham Heart Study ($P = 0.0026$), the lack of correction for multiple testing (a Bonferroni adjusted P -value threshold of 5.7×10^{-7} was needed after performing 86604 statistical tests)²⁴ in combination with the insufficient sample size of the discovery cohort may explain the lack of replication of this association in three well-powered follow-up studies,^{25–27} in a meta-analysis of 74 345 subjects²⁸ and more recently in the large-scale GIANT GWAS initiative,²⁹ even if others have pointed out a context dependent association of the rs7566605 with obesity as an alternative explanation.^{28,30}

Population stratification and confounding

In a case-control study, controls should ideally be selected from the same population as cases. Allele frequency of genetic variants can vary depending on the ethnic background or even the geographical location. When cases and controls are drawn from multiple ethnic or geographic groups whose phenotype prevalence and allele frequencies differ for the gene variants tested, confounding unintentionally occurs, leading to false positive associations.^{4,8,31} This is likely the case for obesity predisposing variants, as several of them show highly varying levels of inter-ethnic or inter-geographic allele frequency variation, possibly due to positive diversifying selection (for example, *ENPP1* rs1044498, *FTO* rs9939609 or *LCT* rs4988235).^{32–34} In addition, prevalence of obesity strongly varies according to the ethnic background in a specific country.³⁵ Even if such confounding seems to have modest consequences in case-control studies, it may generate spurious association signal if not properly taken into account, especially in large-scale association studies.³⁶ GWAS for obesity-related traits have reported modest but real evidence of population stratification.^{5,19}

Genotype misclassification and reliability in quality control

Genotyping errors are frequent in genetic association studies,³⁷ leading to non-differential genotype misclassifications (same probability of being misclassified for all study subjects) or differential genotype misclassifications (varying probability of being misclassified according to the study groups). A 1% increase in genotyping errors will add up sample size by 2–8% to keep the same type I and II errors.³⁸ Genotyping errors from batch to batch, laboratory to laboratory or preferential rejection of particular genotypes (usually heterozygotes) can result in differential genotype misclassifications and significant differences between case and control groups, leading to false positive association. A recent large-scale family-based study using TaqMan technology excluded a role of VNTR *INS* polymorphism in childhood obesity³⁹ despite previous positive association using PCR-based restriction fragment length polymorphism (RFLP).⁴⁰ As the reproducibility of RFLP genotyping data has been questioned, this method being highly subjective,⁴¹ the authors suggested that the lack of replication may be a result of previous genotype misclassification from the RFLP method.³⁹

Inappropriate statistical analyses

Numerous studies use the χ^2 or Fisher exact test to estimate the difference of genotype distribution in cases and controls. Linear/logistic regression is therefore a more relevant approach to make comparisons between groups as it allows adjustment for confounding or mediating factors, and introduction of gene \times gene or gene \times environment interaction terms.⁴² The *FTO* identification study by Frayling *et al.*⁴³ is illustrative of the importance of properly adjusted association studies for confounding or mediating factors. The rs9939609 in *FTO* gene was found initially by GWAS for type 2 diabetes. This strong

association disappeared after further adjustment in the linear regression model for BMI, suggesting that the *FTO* intron 1 variant predisposition for diabetes was mediated through an effect on BMI.⁴³

In some association studies, when no expected main effect is observed, researchers tend to perform post-hoc subgroup analyses to dredge some apparent positive association.³¹ Positive association drawn from such analyses should be viewed with caution.

REPLICATION MAY BE CHALLENGING EVEN WHEN THE ORIGINAL RESULT IS A TRUE POSITIVE ASSOCIATION

There is a consensus that a single study, even when well-designed and powered, is not sufficient to establish a valid genotype-phenotype association and further replication is needed. The main interest of replication is not only to increase the statistical significance of the result but also to prove that the initial association is true and not the result of methodological inadequacies.⁴⁴ Given that many published associations could not be replicated, several high-impact journals such as *Nature Genetics* stipulate in their publication guidelines that replication of an original association in a second independent cohort is requested prior to consideration for publication.⁴⁵ The *FTO* gene identification story is illustrative of the interest of independent replications to be confident about a genetic discovery. In 2007, four different teams independently discovered that variation in the intron 1 of fat mass- and obesity-associated (*FTO*) gene is the major contributor to polygenic obesity in populations of European ancestry.^{43,46–48} However, replication does not always follow this ideal route, and the lack of association in follow-up studies does not automatically imply that the initial study did not pick-up a true association signal. Below we describe the main reasons why replication of a true association may be challenging.

Sample size

A well-powered replication design is crucial to avoid the risk of both false negative and false positive association reports.⁴⁹ Power estimation of the replication samples has become an issue with the emergence of large-scale discovery studies through international consortiums. The current largest GIANT meta-analysis of 249 796 individuals in obesity association study has confirmed 10 previously identified loci associated with BMI and 4 loci associated with waist circumference, and revealed 18 new loci associated with BMI.⁵⁰ Among these 32 loci, rs1558902 (*FTO*) explains 0.34% of variance in BMI, and the remaining loci harbor smaller effect size. A recent study by Zhao *et al.*⁵¹ examined all 32 loci identified from GIANT study in 1097 childhood obesity cases and 2760 controls, but the associations of only 9 SNPs out of 32 were confirmed. Phenotypic heterogeneity (adult BMI versus childhood obesity) may explain why some genetic variants are not replicated, but beyond this limitation the Zhao *et al.*⁵¹ study illustrates the difficulty to replicate association signals with small effect size issued from large GWAS meta-analyses. With the sample size used in this study, the statistical power was 99% to detect an OR of 1.5, 73% to detect an OR of 1.15, but only 19% to detect an OR of 1.05 (minor allele frequency = 0.2, one-sided $\alpha = 0.05$), whereas the range of the effect size of obesity predisposing variants is typically between 1.05 and 1.5.

Gene \times gene, gene \times environment interaction

Gene \times gene or gene \times environment interactions specific to the discovery sample may challenge the ability to replicate the initial discovery in independent study designs.⁵² If the other genetic and relevant environmental factors are not balanced in populations between initial and subsequent studies, the variance in effect size from gene \times gene and gene \times environment interactions may in

part explain the inability in replicating the association. Statistical power issues are critical in gene \times gene or gene \times environment interaction studies and meta-analytic approaches are recommended to reach sufficient power.^{53,54} Different study designs enable the study of gene \times gene or gene \times environment interactions, but power calculations have shown that population-based or nested case-control studies have a higher capacity in detecting interactions than retrospective case-control studies.^{55–57} Case-control studies may be relevant in detecting gene \times gene, gene \times environment interactions in specific situations, as several parameters (for example, allele frequency, magnitude of interactions, genetic model and linkage disequilibrium with the causal variant) impact the statistical power.^{55,58} Convincing evidence of gene \times gene or gene \times environment starts to emerge in the field of obesity. Interaction between the *FTO* intron 1 variant, the level of physical activity and BMI or obesity has been described in 14 independent studies and confirmed in a meta-analysis of 218 166 adults where physical activity attenuated the odds of obesity conferred by the variant by 27%.⁵⁹ A recent study conducted in 62 245 east Asian subjects identified a significant gene-gene interaction ($P = 2.0 \times 10^{-8}$) between two SNPs in the *KLF9* and *MSTN* loci.⁶⁰ These data support the existence of gene \times gene and gene \times environment interactions that may interfere with the replication of an initial association.

Heterogeneity

Genetic/ allelic heterogeneity exists where multiple alleles at the same locus are associated with the same disease, so that different studies may find different alleles associated with the same disease. Linkage disequilibrium varies with the ethnic background and most of the disease-associated SNPs issued from GWAS are proxy (a genetic marker of the disease in a specific population) but not causal, which results in misclassification of the functional risk alleles. Some SNPs are causal but may be associated in an ethnic-specific manner.⁶¹ If a negative replication is obtained in a different population, this might lead to reject a true association in that specific population as a consequence of ignoring allelic heterogeneity. Another consequence of genetic heterogeneity is that different polymorphisms can be associated with obesity-related traits by independent teams, and it can puzzle researchers when they select genetic markers for replication studies. In 2007, four SNPs in high linkage disequilibrium (rs9939609, rs1421085, rs9930506, rs1121980) were described as lead signals for the association between *FTO* and obesity by different teams.^{43,46–48} As no consensus has been made to select a unique lead SNP for replication, follow-up studies using any of the four SNPs have been published, puzzling subsequent meta-analysis initiatives.⁶²

The lack of replication in follow-up studies may sometimes be explained by phenotypic heterogeneity (different phenotypes used in the discovery and the follow-up studies). For instance, BMI can be studied as a quantitative or a categorical trait. Several thresholds can be used to define the status (overweight, class I, II or III obesity). Although highly correlated, these different phenotypes cannot be considered as identical. Following the initial report of association between the *GAD2* -243 A>G variant and morbid adult obesity among the French,²² Groves *et al.*⁶³ assessed its role in 573 UK pedigrees ascertained for type 2 diabetes, and found a nominal but directionally inconsistent association between *GAD2* -243 A>G and BMI. The authors concluded that their data 'did not replicate the previous associations' while acknowledging the phenotypic heterogeneity in the two studies.⁶³

Depending on the study design and recruitment strategy, ascertainment heterogeneity can partly explain the lack of consistency between initial and replication association studies. Even if the same phenotype is used to classify subjects as cases

(for example, $BMI \geq 30 \text{ kg m}^{-2}$), different subpopulations of patients may be collected according to the recruitment procedure (population-based, pedigrees selected for a family history of obesity, hospital, type 2 diabetes cases, incident cases). The age window of the collected sample is also crucial. Obesity in childhood, adolescence, adulthood and agedness are likely to harbor a complex pattern of inheritance, as recently shown for *FTO* intron 1 gene variant and BMI variation.⁶⁴

Inheritance model

Information on the inheritance model is useful in replication studies or biological experiments.⁶⁵ When a formal procedure to attribute the best-fitting inheritance model for obesity was applied to the *ENPP1* K121Q gene variant, a significant departure from the additive to the recessive model was observed,⁶⁶ and later confirmed for the association between K121Q SNP and type 2 diabetes.⁶⁷ Association under a different genetic model from that reported in the original study cannot be considered as a formal replication.

Subjective interpretation of data

Subjective overstatement or understatement in the interpretation of replication data can be observed.⁶⁸ Researchers may over-interpret positive findings in the follow-up studies when attempting to replicate their own original findings, or negatively interpret follow-up studies when attempting to refute their competitors' original claim of association, especially if the original claim was published in a prestigious journal (the Proteus phenomenon).⁶⁹

GUIDELINES FOR PROPER DISCOVERY AND REPLICATION ASSOCIATION STUDY DESIGNS

Discovery association study designs

Study designs. Four designs are classically used in genetic association studies: prospective cohort, case-control, family-based and quantitative trait association study. The prospective cohort study is often considered as the gold standard in epidemiology, but the rates of collecting disease cases (other than common ones) and their follow-up are slow. Consequently, the statistical power is usually weak.⁴⁹

A nested case-control can be designed for a prospective study. Disease cases collected during the follow-up are matched to non-disease controls selected from a portion of the entire cohort subjects. This design minimizes the recall bias, selection bias and inadequate/unreliable records of the environmental exposure from a retrospective case-control study, particularly when gene \times environment hypotheses are being tested.

Family-based design is optimal in specific situations, such as the identification of disease-associated variants subjected to parental imprinting, or in haplotype studies (the reconstruction of the haplotype phase is improved by availability of parental genotypes). It is robust to population stratification bias,⁷⁰ but its main limitation is the lack of power, especially if the effect sizes are small.⁷¹ If the power is 90%, two-sided $P = 0.001$ and control allele frequency is 20%, 1765 trios will detect an OR of 1.30, and for an OR of 1.20, 3731 trios will be necessary, representing 50% more participants than for a case-control study.

Quantitative trait study (for example, BMI) in large-scale population-based samples has proved to be an efficient method to identify novel susceptibility loci.²⁹ However, it requires larger sample size than case-control studies to reach the same statistical power,⁷² and this limitation can become critical if expensive technologies are used (for example, genome-wide DNA arrays).

Thus, a case-control study represents the most powerful and cost-efficient method to perform genetic association studies,⁷³ if the two study groups are selected properly. Population-based

obesity cases can be recruited, but the power of a case-control study can be increased potentially by applying an enrichment sampling strategy.⁷⁴ Obesity cases coming from hospital clinics, having a strong family background of the disease, an early age of onset or a more severe phenotype are likely to be enriched for genetic susceptibility in comparison with obese subjects randomly selected in the general population. Though enrichment sampling strategy is useful to improve power in genetic association studies, it inflates the relative risk and population attributable risk of the associated gene variants, and population-based follow-up cohort studies will be needed to obtain a reliable estimation of these parameters.⁴⁹

To decrease potential sources of heterogeneity between the two study groups in case-control studies, controls and cases need to be selected from the same population. Potential confounders need to be ascertained in both and adjusted. Ideally, subjects in two study groups should be matched for ethnic/geographical origin, age, sex and additional variables for obesity (for example, level of physical activity) to adjust the regression model for confounding factors and to reduce residual error. 'Super control' subjects can also be selected (normal-weight subjects with no familial background of obesity⁷⁵ or extremely lean phenotypes).⁷⁶

Phenotype. An accurate and specific phenotype of interest is critical in the design of genetic association studies.⁸ A phenotype with a substantial heritability ($h^2 > 50\%$) should be favored and may allow the finding of etiological genetic variants in realistically achievable sample sizes. The ideal phenotype should be clinically and biologically relevant, not too rare, inexpensive and easy to identify in different places, thus allowing large-scale and feasible discovery and replication studies. It should be well defined so that measurement error, misclassification and heterogeneity can be minimized. BMI as a quantitative or qualitative trait is the more commonly used phenotype in genetic association studies for obesity. It is a highly heritable trait, easy and inexpensive to measure. As BMI is available in large data sets, genetic association meta-analyses have been performed in up to 250 000 subjects.²⁹ However, it also presents several limitations. Subjects in the higher BMI range tend to under-report their weight, resulting in inaccurate BMI measurement and misclassification in case-control studies. Thus, self-reported BMI data by the participants need to be considered with caution, and BMI data measured by medical staff is preferred.⁷⁷ The value of BMI to estimate the degree of adiposity may be questioned. Although it is strongly correlated with fat mass in obese subjects, in normal weight and underweight subjects the correlation is weaker.^{78,79} At a given BMI, substantial variation in fat mass was reported.⁷⁸ Because phenotypic precision is critical for genetic investigations, the use of 'deep phenotyping' information (for example, body fat content and percentage of fat mass estimated by dual-emission X-ray absorptiometry or bioimpedance, behavioral food intake measured by *ad libitum* meal test⁸⁰ and energy expenditure estimated by room respiration calorimetry⁸⁰) in large sample sizes may dissect the genetic influences more accurately on obesity.⁸¹ Consistent with this hypothesis, a recent GWAS for body fat percentage in 32 626 individuals, with a replication of the best association signals in 39 576 individuals, identified three loci associated at a genome-wide level of significance: *FTO*, *IRS1* and *SPRY2*. Two out of three loci were not identified by previous large-scale GWAS meta-analysis for BMI.⁸²

Recent GWAS for obesity-related traits have collected phenotype information in individuals living in widely heterogeneous environments. Although successful, this approach may have missed gene variants associated with BMI in specific environmental exposures. As the genetic basis of body weight regulation is unlikely to be fully discernible in individuals who are at stable body weight,⁸³ it may be of particular interest to perform genetic association studies for adiposity change in response to a major

environment modification (antipsychotic drug use, smoking cessation, rapid change in the level of physical activity or diet habits, caloric restriction, obesity surgery). Such phenotypes may provide a more comprehensive molecular basis for genetic predisposition to obesity.

Gene identification strategies. In the field of genetic epidemiology, we usually distinguish hypothesis-driven candidate gene and hypothesis-free genome-wide studies.⁸⁴ Candidate genes are selected based on the prior evidence of its role in the disease or phenotype of interest. Arguments to select a specific candidate gene may come from biology, functional genomics, pharmacology, animal models or genetics. The success rate of candidate gene approaches has been poor in the obesity field.³ We consider the candidate gene approach as a relevant strategy as far as a careful gene selection process is applied. For instance, candidate gene studies focusing on *MC4R* and *PCSK1* genes led to the identification of common variants reproducibly associated with obesity risk/protection.^{17,85,86} However, the candidate gene selection was based on three strong prior arguments: (1) an obesity phenotype in genetic mouse models; (2) the involvement in monogenic human obesity; (3) a likely functional role of the selected polymorphisms.

Hypothesis-free GWAS have been exceptionally successful and are to date the most efficient way to identify common variants (minor allele frequency > 1%) associated with obesity-related traits. The main limitation of genome-wide approaches in comparison with the candidate gene strategy is the high level of significance needed to adjust for the multiple tests performed and the situation may be even worth in the context of whole-exome or whole-genome studies. Testing the association of 30 000 genes with obesity rather than only one is certainly more exhaustive, but as a result a higher level of statistical evidence for genes with little supporting biological information is needed before significance is attributed. As pinpointed by John Ioannidis, 'the greater the number and the lesser the selection of tested relationships in a scientific field, the less likely the research findings are to be true'.⁶⁸

A hybrid approach, the hypothesis-driven genome-wide association analysis, may lead to an overall decreased number of statistical tests and to more relaxed significance thresholds by the prior statement of specific hypotheses to be tested. Genome-wide analyses restricted to a specific biological pathway,⁸⁷ to a linkage region,⁸⁸ to genes specifically expressed in one important tissue,⁸⁹ to genes showing different expression patterns related to the disease,⁹⁰ to prioritized candidate genes,¹⁹ to potentially damaging SNPs, or to SNPs harboring an evolutionary signature, may increase success of picking up association signals that have been missed by conventional GWAS analyses.

Genotyping methodology and quality control procedures. Quality control to prevent, detect and minimize biases and errors during genotyping should be one of the top priorities in genetic association studies.⁹¹ From earlier genotyping method of RFLP to PCR-based high-throughput genotyping, the genotyping error rate has significantly decreased, but errors and biases still exist. Sample tracking methods are encouraged to ensure a perfect correspondence in DNAs, genotypes and phenotypes, and an accurate analysis. Differences in extraction methods and storage of DNA samples collected from cases and controls or in different study centers should be mentioned. DNA quality and concentration need to be first documented to exclude low-quality DNA samples from further genotyping. Highly reliable and objective genotyping methods are recommended and genotyping protocols need to be described. Internal controls, duplicated samples and blank controls should be distributed in plates to ensure correct orientation and absence of DNA contamination. Genotyping quality control procedures include call rate calculation

and deviations from Hardy-Weinberg equilibrium (HWE) separately in cases and controls. Call rate higher than 95% and HWE with P higher than 0.005 in the controls must be targeted, as an association may induce a modest deviation from HWE in affected subjects.⁹² The assay reproducibility should be estimated by the calculation of a concordance rate in a sufficiently representative sample size (usually > 10% of the whole genotyped sample). A concordance rate higher than 99% is needed to allow further analyses of the genetic markers. If a family-based design is used, the consistency of genotypes with mendelian expectations should be checked. Additional quality control procedures for the most critical results include the comparison of marker allele frequency in the experiment to public human genome databases and to individuals of comparable ethnicity, careful examination of the genotyping cluster plots to avoid a technical artifact, validation by an independent genotyping method or genotyping of additional genetic markers in strong linkage disequilibrium to confirm the association with the phenotype of interest.⁴⁶ A flow chart scheme summarizing the quality control procedures should be provided at least as supplementary material.

Statistical analyses. Prior statistical analyses should be stated and a table with the estimated power of the study design to detect a wide range of effect and minor allele frequency should be provided. As only few associations first reported from small studies are confirmed in larger replication data sets,²¹ the scientific community must be warned against underpowered discovery studies and promote the generalization of large-sized and well-powered discovery study. A limited number of additional analyses of subgroups such as age, sex, or categories of obesity should be specified in a prior hypothesis in addition to tests in the whole sample, and interpretation of results from subgroup analyses should be done cautiously and accompanied by interaction tests to prove significant between-group heterogeneity. We recommend using statistical tests that enable the adjustment for confounding variables. In multiple SNP haplotype analyses, the investigators should detail the methodology used. Although the importance of adjusting for multiple comparisons in genetic association studies is generally accepted, there is no universal criterion on how it can be achieved. False discovery rate, Bayesian procedures and adjusted Bonferroni correction are acceptable.^{31,93,94} Application of the Bonferroni correction tends to be over-conservative because the tests performed in a genetic association study are unlikely to be fully independent (lack of independence of the three inheritance models, correlation between different phenotypes and linkage disequilibrium between SNPs). In GWAS, we recommend a more relaxed P threshold of 5×10^{-6} to select a limited number of promising genetic markers for further large-scale replication.²⁹ In that situation, a Bonferroni corrected P of 5×10^{-8} should be applied to the discovery plus replication (stage 1 + stage 2) joint analysis to confer significance at the genome-wide level.⁹⁵ In the recent GIANT GWAS meta-analysis for BMI in 123 865 individuals, the application of a more relaxed threshold of $P < 5 \times 10^{-6}$ for SNP follow-up rather than a threshold of P of 5×10^{-8} induced a modest burden of false positive association signals (24%) but substantially increased the harvest of disease-associated validated loci (from 68%).²⁹ For most original associations of common gene variants with common diseases, the genetic model of inheritance is not properly addressed.⁶⁵ A formal procedure to attribute the best-fitting inheritance model for the phenotype studied is recommended from the discovery study and should be used for replication efforts. If several SNPs are associated at the same locus, conditional regression analyses should be applied to determine if these SNPs are independent and to select the lead SNP signal(s) to follow up in replication efforts.

Population stratification. The first step to lower population stratification is to collect cases and controls in the same geographic area,

Table 2. Guidelines in designing and implementing discovery and replication studies

<i>Discovery association study</i>	
<i>Study designs</i>	
Prospective cohort study	Gold standard Slow process Low statistical power Expensive to follow-up Possible high withdrawal rate
Family-based design	Useful in the context of parental imprinting Useful in the context of haplotype studies Robust to population stratification bias Low power when effect size is small Expensive design
Quantitative trait study	Large sample size needed Unable to detect genetic effects at the extremes of the trait distribution
Case-control study	Most powerful and cost-efficient Cautions needed in selecting cases and controls to lower heterogeneity
<i>Phenotype selection</i>	
BMI	High heritability Clinically and biologically relevant Common Non-negligible measurement error/ risk of misclassification if self-reported Not accurate to measure adiposity body fat content and percentage of fat mass
'deep phenotyping'	Food intake/ behavior Energy expenditure Adiposity response to environmental changes
<i>Gene identification strategies</i>	
Candidate genes	Hypothesis-driven Prior evidence from other disciplines such as biology, functional genomics, pharmacology, animal models or genetics
Genome-wide studies	Hypothesis-free Efficient in identifying novel common variants without previous candidacy High level of significance needed
Hypothesis-driven genome-wide association	Lower number of statistical tests Useful to pick up the association signals missed by conventional GWAS Focus on biological pathways, linkage regions, transcriptomics data
<i>Genotyping methodology and quality control procedures</i>	
DNA sample handling	
DNA quality	
Description of genotyping protocol	
Genotyping quality control: DNA contamination, call rate, Hardy-Weinberg equilibrium, concordance rate, external control from public human genome databases, genotyping cluster plots, mendelian inconsistencies, validation by another technology and analysis of neighboring SNPs in strong linkage disequilibrium	
<i>Statistical analyses</i>	
Analytic methodology selected according to the study design	
Prior statement of hypotheses to be tested	
Multiple testing correction	
Best-fitting inheritance model	
Selection of lead SNP for replication efforts	
<i>Population stratification</i>	
Cases and controls selected from the same population and geographic area	
Adjustment for population stratification	
<i>Replication studies</i>	
<i>Systematic replication of promising original reports</i>	
Systematic follow-up of promising associations	
Objective interpretation of data	
<i>Statistical power</i>	
Sample size calculation corrected by the 'Winner's curse effect'	
<i>Homogeneity</i>	
Same ethnicity and similar age, sex, geographic origin	
Same phenotype, measurement method	
Same inheritance model	
Same study design, ascertainment strategy	
Same genetic marker, statistical analysis, adjustment for confounding variables	
<i>Meta-analysis</i>	
Powerful to confirm genetic associations at a stringent level of significance	
Estimation of publication bias and between-study heterogeneity	
<i>Additional studies</i>	
Extension of confirmed variants to other populations (different ethnic backgrounds, age, environmental risks)	
Fine-mapping	
Gene \times environment, gene \times gene interaction studies	
Functional assay to confirm the biological role of gene variants	
Additional biological evidence	

Abbreviations: BMI, body mass index; GWAS, genome-wide association studies; SNP, single-nucleotide polymorphism.

first using self-reported ethnicity to equilibrate cases and controls in multi-ethnic studies, or restricting the analysis to one ethnic background if a particular ethnic group predominates. As relying solely on self-reported ethnicity is not sufficient to exclude population stratification⁹⁶ and subtle population stratification is observed within ethnically homogeneous populations,⁹⁷ several methods (genomic control, EIGENSTRAT, PCA-L and LAPSTRUCT) have been developed to correct for population stratification⁹⁸ and are now routinely used.^{6,19} These methods are based on using a set of ancestry informative markers,⁹⁹ which are easy to derive from GWAS data sets,¹⁰⁰ but require further genotyping in candidate gene studies. These markers may also be useful to account for cryptic relatedness, another potential source of spurious association in case-control designs.¹⁰¹ Alternatively, family-based association tests can be used to confirm an association derived from a case-control study, as they are insensitive to population stratification biases.¹⁰²

Replication study designs

Systematic replication of promising original reports. Several teams working in the field of obesity should be commended for systematically replicating promising association reports in their own cohorts. However, other replication efforts may not have been as systematically conducted even if the original report is well designed and statistically sound. To move on the right direction, researchers, but also editors of influential journals, need to consider well-designed initial and follow-up (negative or positive) reports of association as equally important. Replication studies need to be conducted more systematically and less selectively, and an objective interpretation of data is crucial.

Statistical power. When the original association report describes a true association signal, the effect size is often overestimated due to the Winner's curse effect.¹⁰³ Subsequent follow-up studies often fail to replicate the association owing to underestimation of the required sample size. Hence, when calculating the statistical power of the replication design, the investigators may correct the OR from initial study by taking into account the Winner's curse effect.¹⁰³

Heterogeneity. If both original and replication studies are well designed and well executed, the inability to replicate an original finding indicates a genuine difference that may be due to heterogeneity from many sources.¹⁰⁴ Heterogeneity among studies represents a major problem in replicating a true original signal. Moonesinghe *et al.*⁵² have shown that in the presence of large between-study heterogeneity, true associations may not be possible to replicate with consistency, no matter how large the study is. When designing a study in the initial rounds of replication, investigators need to avoid all the potential sources of heterogeneity and focus on one main question to increase the chance to reproduce the original association finding. To reduce the genetic and environmental heterogeneity, populations in the initial and in the follow-up studies should share the same ethnicity and ideally come from the same country. Correction for population stratification should be applied as well. Same age window and sex ratios should be targeted. The study design, genetic marker, statistical analysis and genetic model should be the same as those in the initial study. To lower phenotypic heterogeneity, the same phenotype, measurement method, recruitment strategy and adjustment for confounding variables should be applied.

Meta-analysis. There are always limitations in individual association study. Common genetic variants harboring modest OR or rare variants with higher OR rarely reach a conclusive threshold of association in a single study. Meta-analysis of data sets from comparable studies improves the power to confirm genetic

associations at a stringent level of significance. Unfortunately, the heterogeneity in the design of follow-up studies (genetic marker, phenotype and genetic model) and insufficient data description may reduce the number of eligible studies and the statistical power in subsequent meta-analyses. Hence, Gallo *et al.*¹⁰⁵ recently provided detailed guidelines to accurately report the findings of epidemiological studies involving biomarkers through the STrengthening the Reporting of OBservational studies in Epidemiology - Molecular Epidemiology (STROBE-ME) statement. Estimation of the degree of between-study heterogeneity and of potential publication bias are important for the interpretation of meta-analyses.¹⁰⁶

Additional studies. Once the risk of false positive association has been ruled out by initial replication studies, the focus of the association can be extended to different age windows, different study designs and ascertainment criteria for cases and controls, additional phenotypes related to obesity, or to the study of gene \times gene/ gene \times environment interactions. The association can be studied in different ethnic backgrounds to estimate the worldwide contribution of a given locus, and to fine-map the association signal. Using a dense SNP map in the associated interval is strongly recommended, as the initial SNP associated with obesity is probably in linkage disequilibrium with the causal locus, while linkage disequilibrium patterns vary according to ethnicity. Once the putative causal susceptibility variant(s) has been identified, functional assays should be used to examine whether a SNP variant alters protein expression or function. As an association does not always imply causality, biological insights are essential in increasing the credibility of the observed genetic association,¹⁰⁷ as recently illustrated by the *FTO* gene. In 2007, a link between *FTO* and obesity was established in Europeans through association studies when *FTO* was 'a gene of unknown function in an unknown pathway'.^{43,46} Subsequently, more than 400 articles have been published on *FTO*, providing strong biological evidence that *FTO* is a causal gene underlying obesity.¹⁰⁸

CONCLUSION

We have reviewed the reasons for the lack of reproducibility in obesity genetic association studies and offered guidelines for proper discovery and replication association designs (see Table 2). 'Learn from yesterday, live for today, hope for tomorrow': a quote attributed to the physicist Albert Einstein perfectly depicts our position in the young field of genetic epidemiology of obesity. Genetic association studies may first appear simple but actually face numerous challenges. After some initial disappointments, methodological, technological and scientific developments have recently led to more robust associations and renewed attention to genetic epidemiology. We expect a prolific period of discovery in human obesity genetics ahead and hope that this review will contribute to the advancement of these noble goals.

CONFLICT OF INTEREST

The authors declare no conflict of interest.

ACKNOWLEDGEMENTS

We thank Dipika Desai and Emeritus Professor Patricia Chang for reviewing and editing of the manuscript, and the reviewers for their helpful comments.

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