Statistical Methods and Software for Substance Use and Dependence Genetic Research

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Abstract: Background: Substantial substance use disorders and related health conditions emerged during the mid-20th century and continue to represent a remarkable 21st century global burden of disease. This burden is largely driven by the substance-dependence process, which is a complex process and is influenced by both genetic and environmental factors. During the past few decades, a great deal of progress has been made in identifying genetic variants associated with Substance Use and Dependence (SUD) through linkage, candidate gene association, genome-wide association and sequencing studies. Methods: Various statistical methods and software have been employed in different types of SUD genetic studies, facilitating the identification of new SUD-related variants. Conclusion: In this article, we review statistical methods and software that are currently available for SUD genetic studies, and discuss their strengths and limitations.

Keywords: Substance dependence, linkage analysis, association analysis, interaction analysis, meta-analysis, GCTA.

1. INTRODUCTION

Substance use disorders present a significant global public health problem that places a substantial burden on individuals, health care systems, societies and economies [1, 2]. It is estimated that approximately 27 million people in the world suffered from substance use disorders in 2013, resulting in a remarkable global burden of disease that is expected to become more prevalent over time [3]. This burden is largely driven by the substance-dependence process, which is a complex disorder influenced by both genetic and environmental factors.

Substance dependence is defined as the syndrome of substance misuse that leads to adverse consequences and includes a cluster of symptoms such as tolerance, withdrawal, and inability to stop using (see DSM-IV substance dependence for the complete diagnostic criteria) [4]. The term substance often refers to products with addictive potential, such as tobacco, alcohol, cocaine and other licit and illicit drugs. The development of the substance-dependence process involves several steps: the initiation of substance use, the transition from experimental use to regular use, and the actual development of dependence [5]. Each step is influenced by both environmental and genetic factors. Twin studies have suggested a substantial genetic contribution to Substance Use and Dependence (SUD) [6]. Over the past few decades, we have successfully identified new genetic variants associated with SUD through linkage, candidate gene association, genome-wide association, animal and sequencing studies. Various statistical methods and software have played an important role in helping the identification of SUD-related variants. This article aims to give an overview of the widely used methods and software for SUD genetic data analysis, including data management, linkage analysis, association analysis and other analysis.

2. DATA MANAGEMENT

The quality-control process is an essential step in data analysis. It is used to identify and remove inaccurate information before the downstream analysis (e.g., genetic association analysis). Researchers utilize different quality control filters and criteria. While some researchers have implemented stringent criteria in order to have high-quality data, others adopt flexible standards to keep as much information as possible.

Commonly used filters are based on the call rate, Minor Allele Frequency (MAF), and Hardy-Weinberg Equilibrium (HWE) test. A fixed call rate is commonly adopted in a study to remove low-quality SNPs and samples. For instance, if a researcher sets a threshold of 1% for sample call rate in a project, then samples with more than a 1% failed genotyping call are removed; likewise, if a researcher sets a threshold of 5% for SNP call rate, individual SNPs with 5% failed genotyping call are also removed [7]. The thresholds of sample call rate and SNP call rate do not have to be the same. Besides the call rate, filters based on MAF and the HWE test are also frequently used. HWE tests are often conducted on the control samples to exclude low-quality SNPs, while an MAF threshold (e.g., MAF>1%) allows the study to focus on...
common SNPs and reduce the number of tests. Most of the above quality control procedures are implemented in the PLINK program [8]. PLINK offers several additional quality control procedures as well, such as detecting and removing duplicate samples.

Quality control procedures have also been developed for different types of studies. For population-based studies, individuals are required to be unrelated. PLINK and several other software offer a procedure to identify and remove individuals who are related. For family-based genetic data, statistical methods and software such as PREST [9] can be used to identify sample and pedigree structure errors based on the autosomal markers [10]. Misspecified familial relationships can also be detected by PREST [9], which infers the underlying relationships from the shared Identical By Descent (IBD). The reassigned family relationships can also be verified by using PREST [11, 12]. However, if the software fails to resolve the error, those families are subsequently treated as missing [13]. A Mendelian inconsistencies check is another commonly used quality control procedure for family data. Mendelian inconsistencies can be detected and set as missing by using the PedCheck [14] and Merlin [15] programs [11]. Markers with a high frequency of Mendelian segregation errors are often excluded from the analysis [13].

3. LINKAGE ANALYSIS

Linkage analysis uses family data to locate genetic regions that are in linkage with a given trait. It was one of the most commonly used tools to infer the location of trait-related genes on chromosomes before the age of Genome-Wide Association Studies (GWAS) [16]. Linkage analysis can be classified into model-based linkage analysis and model-free linkage analysis. While model-based linkage analysis assumes a particular genetic model (e.g., the causal allele frequency, penetration functions) [17], the model-free linkage analysis does not require such assumptions. These two analyses are sometimes referred to as parametric and non-parametric, although both of them require the estimation of parameters [18].

Model-based linkage analysis is powerful for scenarios with prior knowledge of the underlying genetic models, and has been implemented in several software packages (e.g., Merlin, FASTLINK). However, for complex traits like SUD, we have limited knowledge about the underlying genetic model. Misspecification of model parameters in SUD model-based linkage analysis can lead to biased estimators and incorrect inferences.

Model-free linkage analysis is based on allele-sharing, which usually compares genetic similarity among affected relatives inferred from data to that from expectations. Because the model-free linkage methods do not require the specification of the inheritance parameters of a genetic model, they have been adopted in many genetic studies of SUD [17]. One of the most popularly used model-free linkage methods is the Haseman-Elston algorithm [19], which was later extended by Sham et al. [20]. The extended algorithm can perform genome-wide linkage analyses and has been implemented in the program Merlin [21]. GENEHUNTER [22], another linkage method for affected sib-pair analysis, has also been used by many SUD researchers [23].

The variance component method has also been widely used. It partitions the phenotype variations among family members into different variance components explained by the effects of the genes in the region of interest, additive genetic effects of other genes, and non-shared environmental determinants [24]. Software such as Merlin can be used to perform a variance components linkage analysis. All of the above programs can be used to conduct both two-point or multi-point linkage analyses.

Model-free linkage analysis is a robust but less powerful approach. Although including unaffected individuals generally provide more power for linkage analysis, there is a potential loss of power in the event of misclassification of affection status [23]. On the other hand, model-based linkage analysis requires knowledge of the parameters of the genetic model, and has advantages in terms of efficiency and a better location solution. Researchers, therefore, need to determine which methods to use based on the study purpose.

Although linkage analysis can be used to map genes, the SUD-related chromosome regions identified by linkage analysis can be up to several million bases and contain thousands of genes. Moreover, family data especially those with large pedigrees are difficult to collect. Compared to linkage analysis, association analysis is generally more powerful and can be used for fine mapping. Thus, linkage analysis is used less often than association analysis in the recent SUD genetic research. To borrow the strengths from both analysis, some researchers [11] have adopted a multistage design that combines linkage analysis with association analysis to identify genetic variants associated with SUD.

4. ASSOCIATION ANALYSIS

Linkage analysis has been successfully implemented in many family studies, identifying genetic regions linked to SUD. However, linkage analysis is subject to a few limitations, including low power for detecting variants with small effects and being less useful for the fine mapping of causal variants. As stated above, some of these limitations can be overcome by association studies [25]. Association analysis is based on the concept of linkage disequilibrium. It can be classified into population-based association analysis and family-based association analysis. In other words, SUD-based association studies aimed at identifying genes related with SUD can be conducted by means of a population-based design (e.g., case-control design) with unrelated individuals or a family-based design with related individuals [25]. This section provides an overview of the current statistical approaches and software for population-based and family-based SUD association studies.

4.1. Population-based Association Study

Population-based association studies are conducted on unrelated samples. The most commonly adopted population-based association studies are case-control studies, which compare the distribution of genetic variants in affected subjects with those in unaffected subjects. We summarize below methods for single-locus association analysis, multi-locus association analysis, and briefly discuss the issues of population stratification and multiple testing.
4.1.1. Methods for Single-locus Association Analysis

Assuming that a genetic variant is a Single Nucleotide Polymorphism (SNP) with three genotypes, AA, Aa and aa, where A is the minor frequent allele. One way to code the SNP is assuming an additive allelic effect (i.e., AA=2, Aa=1, aa=0). The additive model, indicating that each additional copy of the A allele increases the disease risk [26], is commonly used in SUD association analysis [27]. In certain circumstances, where researchers have prior knowledge of the underlying mode of inheritance, other models can also be used. For instance, we could assume the SNP follows a dominant mode of inheritance (i.e., AA=AAa=1, aa=0) or a recessive mode of inheritance (i.e., AA=1, Aa=aa=0) [28]. Based on the coded genetic variable, classic statistical methods can be employed to link the SNP to different types of phenotypes.

Standard contingency table methods can be used to test the null hypothesis of no association of the SNP with a categorical phenotype. While the Z test and Chi-squared test are the typical methods, the Fisher’s Exact Test can be adopted for studies with a small sample size. A contingency table method, such as Cochran-Armitage trend statistics [29] can be used to assess the association between a binary phenotype and an SNP with additive coding, which is equivalent to the score test in a logistic regression [30]. A logistic regression model is another popular method for analyzing data with a binary phenotype. It is more flexible than the Cochran-Armitage trend test and can accommodate covariates and model interaction effects [31]. A Manhattan plot is often used to visualize the results from a single-locus association analysis, and a Quantile-Quantile (Q-Q) plot is drawn to check systematic biases (e.g., bias due to population stratification) [32].

Similar to logistic regression, linear regression can be used for testing an association between a genetic marker and a quantitative phenotype, allowing for covariates adjustment [33]. Linear regression assumes that the quantitative phenotype follows a normal distribution. For non-normally distributed data to normal before model-fitting or use a non-parametric method (e.g., U statistics). Popular transformations include log transformation, Box-Cox transformation, and normal quantile transformation. Both linear regression and logistic regression are special cases of Generalized Linear Models (GLM), which have been implemented in most of existing statistical software packages.

4.1.2. Methods for Multi-locus Association Analysis

Complex diseases are often caused by multiple genetic variants and other factors (e.g., environmental determinants) [34]. Since each genetic variant plays a small role in complex diseases, a joint association analysis of multiple SNPs (e.g., all SNPs in a gene) is able to accumulate the effects from multiple SNPs and reduce the multiple testing issue [18, 31].

Methods used for single-locus analysis can also be extended for analyzing multiple SNPs. For instance, multiple logistic regression can be adopted for multi-locus association analysis. The corresponding score test is a generalization of the Armitage test and is related to the Hotelling-\(T^2\) statistic [35].

Most of the methods introduced in the multiple-SNP analysis section do not consider Linkage Disequilibrium (LD) among SNPs. To take LD into account, haplotype-based analysis can be used to infer an association of a specific haplotype or haplotypes with a disease phenotype of interest. Before we perform a haplotype test, we need to first infer the haplotype phase based on genotypes. The likelihood-based approach and the Expectation-Maximization (EM) algorithm can be used to infer possible haplotype phases. A nice feature of the likelihood-based approach is that a likelihood-ratio test can be easily formed to test the association of the inferred haplotypes with the disease phenotype. Alternatively, we can also adopt a score test, which is computationally efficient and is widely used in SUD genetic research. The score test has been implemented in the program HAPLO.STATS [36, 37]. Other programs for haplotype phase estimation include Bayesian approaches, such as those implemented in PHASE and HAPLOTYPE [18]. The Bayesian approach and the partition-ligation (PL) algorithm implemented in the program PHASE [38] are reported to be more accurate in reconstructing haplotypes than the EM algorithm [39, 40].

Similar to multiple-SNP association analysis, a haplotype analysis can potentially reduce the multiple testing burden and be more powerful than a single-locus association analysis, especially when there are strong haplotype-specific effects. Nevertheless, a haplotype-based association analysis may also be subject to issues of uncertain phasing and low power due to a large number of inferred haplotypes [31]. There are several applications of haplotype-based analysis. For instance, Wang et al. found that both CHRNA2 and CHRNA46 were significantly associated with Nicotine Dependence (ND) [41] based on the haplotype-based analysis.

4.1.3. Issues in Population-based Association Analysis

Multiple testing is an important issue in association analysis, especially for a genome-wide association analysis involving millions of SNPs. Performing an association analysis on such a large number of SNPs without considering the issue of multiple testing could result in substantial false positive findings. The Bonferroni correction is an easy way to address this issue [42]. For genome-wide association data with 1 million SNPs, the Bonferroni correction at a family-wise error rate (FWER) of 5% results in a genome-wide significance level of \(5 \times 10^{-8}\) [42-44]. While the Bonferroni correction is easy to implement, it tends to be conservative and does not consider the LD among SNPs [45]. Nevertheless, the Bonferroni correction is still the most popular approach in SUD genetic research. Another popular approach for multiple testing correction is the False Discovery Rate (FDR) approach, which controls the expected proportion of false positives among all rejected hypotheses [46]. FDR tends to be less conservative than the Bonferroni correction.

In population-based association studies, spurious associations could be caused by a variety of confounding factors. Confounding bias has been of concern in SUD genetic studies using samples from multiple ethnic groups (i.e., population stratification), especially in large-scale national/international studies. For genome-wide association studies, Q-Q plot is a useful tool for visualizing a systematical bias due to factors such as population stratification. Deviation
| Software Packages | URL | Platform         | Programming Language | Limitations                                                                 | Computational Cost | Public/ Commercial |
|-------------------|-----|------------------|----------------------|-----------------------------------------------------------------------------|--------------------|--------------------|
| PLINK             | https://www.cog-genomics.org/plink2 | Windows/ Linux/ Mac  | C, C++               | --                                                                          | --                 | Public             |
| PREST             | http://www.atstat.toronto.edu/sun/software/prest | Linux              | C++                  | --                                                                          | --                 | Public             |
| PedCheck          | https://watson.hgen.pitt.edu/register/docs/pedcheck.html | Windows/ Linux/ Mac | C, C++               | PedCheck performs single-locus analysis only.                               | --                 | Public             |
| Merlin            | http://csg.sph.umich.edu/abecasis/merlin/download/ | Windows/ Linux/ Mac | C++                  | --                                                                          | --                 | Public             |
| FASTLINK          | https://cran.r-project.org/web/packages/fastLink | Windows/ Linux/ Mac | R                    | FASTLINK is likely to have difficulty of producing high-quality matches when the overlap between two data sets is small. | --                 | Public             |
| GENE-HUNTER       | http://www-genome.wi.mit.edu/ftp/distribution/software/genehunter | Windows/ Linux/     | C                    | It is not suitable for analyzing large multi-generational pedigrees.         | --                 | Public             |
| HAPLOSTATS        | https://cran.r-project.org/web/packages/haplo.stats | Windows/ Linux/ Mac | R                    | --                                                                          |                    | Public             |
| PHASE             | http://www.stat.washington.edu/stephens/software.html | Windows/ Linux/ Mac | C++                  | When haplotype analysis is conducted on a large number of loci, many of the resulting haplotypes will have small frequencies. The power of haplotype analysis is hence significantly reduced due to the large number of degrees of freedom [18]. |                    | Public             |

(Table 1) contd.…. 
| Software Packages | URL | Platform | Programming Language | Limitations | Computational Cost | Public/Commercial |
|-------------------|-----|----------|----------------------|-------------|-------------------|------------------|
| HAPLO-TYPER       | http://www.people.fas.harvard.edu/~junliu/Haplo/click.html | Windows/Linux/Mac | C++ | Efficiency may be lost because of incorrect prior information. When the sample size is small, the power of haplotype analysis is reduced [110]. | It is computationally faster than the PHASE program. | Public |
| EIGENSTRAT        | http://genepath.med.harvard.edu/~reich/EIGENSTRAT.htm | Linux | C++ | Researchers should pay attention to the experimental design, matching the ancestry and laboratory treatment of cases and controls to the fullest extent possible [111]. | -- | Public |
| STRUCTURE         | http://web.stanford.edu/group/pritchardlab/structure_software/release_versions/v2.3.4/html/structure.html | Windows/Linux/Mac | C, Java | -- | -- | Public |
| SKAT              | http://cran.r-project.org/web/packages/SKAT | Windows/Linux/Mac | R | -- | -- | Public |
| FBAT, PBAT        | http://www.biostat.harvard.edu/~clange/default.htm | Windows/Linux/Mac | C, C++ | FBAT are designed for analyzing nuclear families, but in practice general pedigrees are commonly collected [112]. | It is computationally fast. | Public |
| METAL             | http://www.sph.umich.edu/csg/abecasis/metal/ | Windows/Linux/Mac/Unix | C++ | -- | The analysis of 15 studies and 36 million association tests requires <6 min computing time. | Public |

(Table 1) contd....
from the diagonal line indicates possible population stratification and inflation of spurious associations. One of the most common methods for controlling population stratification is the principal component analysis. In a principal component analysis, each individual receives scores on each principal component. These scores, representing continuous variations in race and ethnicity, can be used to control for the effects of population stratification [47]. The EIGENSTRAT can estimate Principal Components (PCs) on a sufficient number of SNPs, which can then be used as covariates in a regression model to control for population stratification [35]. Several other methods have also been used, including the program STRUCTURE. STRUCTURE first classifies the samples into different clusters and then tests the association within each cluster [48]. The genomic control approach, which estimates genomic inflation factor λ, is another fast and easy way to detect and adjust for population stratification [37]. Unfortunately, most of the above methods are designed to adjust for global ancestry, so they could be inadequate when local population structure is an important confounding factor.

4.2. Rare Variants Association Test

While many common variants have been identified to be associated with SUD, much of the genetic contribution to SUD remains unexplained. There is an increasing interest in studying SUD-related rare variants [49-51]. For such a purpose, methods for rare-variants analysis have been developed.

Methods developed for single-locus analysis can also be used to detect rare variants. However, due to high dimensional and low frequent features of rare variants, single-locus analysis is subject to low power [52]. Single-locus analysis is useful only if the sample size is large enough and/or the effect sizes are large [52].

Compared to single-locus tests, multiple-locus test is generally more powerful for rare variants analysis. Numerous multiple-locus methods and software have been developed for analyzing rare variants. Shrinkage methods, such as LASSO [53] and Bayesian Lasso [54], can be incorporated into the regression framework to select disease-related SNPs and to deal with the high-dimensional genetic data. In addition to these methods, collapsing methods, weighted sum methods and linear mixed methods (e.g., SKAT) have also been developed to detect rare variants. While several SUD-related rare variants have been identified, overall they have a limited contribution to SUD [55].

4.3. Family-based Association Analysis

Compared to population-based association studies, family-based association studies provide robust protection against population stratification at the design stage. In a family-based association study, a typical Transmission Disequilibrium Test (TDT) compares the alleles that are transmitted to an affected child from parents to the alleles that are not transmitted. Therefore, it matches the ancestry background of samples within families, and provides robustness against population stratification at a locus-specific level. Many methods have been proposed for family-based association analysis. Some of these methods are extensions of population-based association methods while accounting for familiar correlations. We only review a few of methods commonly used for SUD genetic research. There are several popularly adopted family-based study designs, such as case-parent trio design and case-sibling design. We briefly introduce these designs and then summarize methods associated with these designs (Table 1).

A case-parent trio design is one of the most popularly used family-based study designs, in which cases and their parents are recruited. The well-known method for case-parent trio design is the Transmission Disequilibrium Test (TDT) [56]. The classic TDT is essentially a McNemar test that compares the distribution of alleles transmitted from parents to affected offspring with that of the non-transmitted
alleles [57]. The original form of TDT can only be applied to one affected child with known parental genotypes. It has been further extended to continuous phenotype, multiple offspring, and missing parental genotypes.

Case-sibling design, in which each proband is matched with one or more unaffected siblings, is an alternative design when the parental genotypes are missing. Case-sibling design requires at least one affected offspring and one unaffected offspring, and their genotypes can’t be identical. Although case-sibling designs are generally robust to population stratification, they are not as powerful as unrelated case-control design because of the similarity of siblings [58].

A Family-Based Association Test (FBAT) is the most commonly adopted method for SUD family-based association analysis. FBAT can be considered an extension of TDT, which allows for any type of genetic model, affected and unaffected siblings, qualitative and quantitative phenotypes and multiple markers [59-62]. FBAT has also been extended to incorporate haplotypes and gene-environment interaction analysis [63]. A software package for FBAT is freely available online. An association test implemented in the FBAT program has the option of computing the p-value from the Z statistic using Monte Carlo sampling under the null hypothesis of no linkage and no association [63]. P-value from FBAT can be computed either by asymptotic theory or by permutation test. FBAT has been successfully used in many studies. For instance, Shirley et al. identified 6 SNPs associated with alcohol dependence by using FBAT [64]. The TDT and FBAT approaches are usually applied to nuclear family data, but in practice, general pedigrees are commonly collected. A Pedigree Disequilibrium Test (PDT) is a method specifically designed for general pedigrees [65], and has been widely adopted in family-based association analysis of SUD. It is also worthwhile mentioning that the integrated software package PBAT (v. 3.5), which contains tools for the design of family-based association studies as well as tools for data analysis [66].

5. G-G/G-E INTERACTION ANALYSIS

Significant progress has been made toward the identification of genetic variants contributing to SUD. Despite these achievements, a large proportion of SUD phenotype variations remain unexplained. Gene-gene and gene-environment (G-G/G-E) interactions could play an important role in the SUD development process. The identification of these G-G/G-E interactions not only explain additional SUD variations, but also elucidate how genetic variants interplay with environmental determinants to cause SUD. A G-G/G-E interaction study, either using a population-based design or a family-based design, takes into account the complex relationship between genetic variants and environmental determinants, and could lead to novel G-G/G-E interactions contributing to SUD.

Most of G-G/G-E interaction studies incorporate a population-based design. While conventional methods, such as contingency-table-based methods and regression-based methods [67, 68], are very useful for G-G/G-E interaction analysis on a handful of variables, they are subject to low performance when there are a large number of genetic variants. Efforts have been made to extend the classic methods to high-dimensional genetic data, such as the development of stepwise logistic regression [69-71] and LASSO [72, 73], but the implementation of these methods on genome-wide data remains a great challenge [74]. One of the popular methods for SUD G-G/G-E interaction analysis is the Multifactor Dimensionality Reduction (MDR) [75]. The fundamental principle of MDR is pooling multi-locus genotypes into high-risk and low-risk groups based on case-control ratio, which effectively reduces the data dimensions. The Generalized Multifactor Dimensionality Reduction (GMDR) method further extends the original MDR method for handling various phenotypes, taking covariates into consideration [76]. MDR, as well as GMDR, employ an exhaustive search algorithm, which could be computationally expensive when considering high-order interactions or a larger number of genetic variants. GMDR-GPU, developed by Zhu et al., not only allows for GWAS data but also runs much faster than the earlier version of the GMDR program by utilizing a more computationally efficient implementation [77, 78]. GMDR has been successfully used in detecting an ND-association interaction between CHRNA4 and CHRNA2 [79]. Other advanced methods, such as a neural network and random forest, have been used in the G-G/G-E interaction analysis, though none of them have been widely used in SUD G-G/G-E studies.

Methods have also been developed for family-based G-G/G-E interaction analysis. FBAT, which allows various pedigree structures, various phenotypes and multiple markers, has been extended to G-G/G-E interaction analysis [57]. Similarly, the extensions of MDR and GMDR, MDR-PDT [80] and PGMDR [81], have also been developed for family-based G-G/G-E interaction analysis. PGMDR allows for covariate adjustment and could attain more power than MDR-PDT.

6. META-ANALYSIS

GWAS allows us to identify SUD-associated variants with large or modest effect sizes. In order to discover SUD-associated variants with smaller effect sizes or low frequency, large-scale genetic studies with a much larger sample size are needed [32]. Meta-analysis has become popular because it does not require individual-level data and is capable of integrating information (e.g., p-values and odd ratios) from multiple independent studies with a substantially increased sample size [82]. Based on a much larger sample, it increases the power of association studies, leading to the discovery of novel small-effect variants and low-frequency variants [28, 83]. Classic meta-analysis methods typically based on combining effect sizes (i.e., fixed effects or random effects), p-values, or Z scores [28]. In this section, we summarize the classic methods available for SUD meta-analysis.

Prior to conducting a meta-analysis, we need to test heterogeneity to make sure that individual studies are similar to each other. I² is a statistic metric that has been widely used for evaluating heterogeneity. It measures the proportion of total variation between studies attributed to heterogeneity [84]. Many programs, such as METAL [85] and GWAMA [86], can estimate I² for the evaluation of heterogeneity.

P-value-based meta-analysis methods have been used for decades but have become less popular due to their limi-
tions (e.g., including weights is not a straightforward process) [87]. Among these methods, the Fisher’s method [82] and the Z-score-based method are often adopted and are closely related to each other [34]. In a typical meta-analysis study, each site generates $p$-values or $Z$ test statistics, and then uploads the results to a server. A meta-analysis method can be implemented to combine the information from all sites for the discovery of low-frequent variants or small-effect variants [82].

Methods for combining effect sizes can be based on two types of models: fixed effect models and random effect models. A mixed effect model, such as the DerSimonian and Laird model [88], is preferred when heterogeneity exists among studies. Otherwise, a fixed effect model should be used [32]. The major advantage of fixed effect models over random effect models is that they maximize discovery power [89]. Inverse variance weighting [90] and Cochran-Mantel-Haenszel [91] are commonly used approaches that provide similar results for fixed-effect-based meta-analyses [34]. Popularly packages for fixed-effect-based meta-analyses include METAL, GWAMA, R and PLINK, among which GWAMA and R can also be used for random-effect meta-analysis.

7. GENOME-WIDE COMPLEX TRAIT ANALYSIS

The genetic variants that have been discovered so far only account for a small fraction of SUD heritability [92]. We should point out that the heritability discussed in this section is the narrow-sense heritability, i.e., the proportion of phenotypic variance due to additive genetic variance. Genome-wide Complex Trait Analysis (GCTA) can help us to study “missing heritability” and to identify which parts of a genome or categories of variants contribute to SUD heritability [93]. Yang et al. [94] developed a versatile tool, GCTA, to study heritability. GCTA is a user-friendly software with four main functions: data management, genetic relationships estimation from SNPs, mixed linear model analysis of variance and linkage disequilibrium structure estimation [93].

GCTA was developed based on a linear mixed method to address the “missing heritability” problem [94]. Yang et al. believe that “missing heritability” can be partially explained by small or medium-effect-size SNPs that don’t reach the stringent genome-wide significance level. GCTA first calculates a Genetic Relationship Matrix (GRM), which estimates the genetic relationships between all pairs of individuals based on the genetic data. Given the calculated GRM, it further estimates the variance components by using a Restricted Maximum Likelihood (REML) algorithm from the Linear Mixed Model (LMM) [93], and then infers the heritability. In LMM, covariates (e.g., sex, age, principle components) can also be easily adjusted. Using the actual genetic data, the estimated heritability from GCTA is expected to be close to that from twins or segregation studies. For example, the estimation of heritability of nicotine dependence is close to 50% [95, 96] provided by twin and family studies. GCTA can not only provide a better heritability estimate but also allows researchers to investigate which parts of the genome or categories of variants contribute to SUD [93].

To better model known disease-related variants, Yang et al. developed a GCTA-COJO module by treating the effects of known variants as fixed effects and excluding these variants from GRM [97]. This module has been applied to SUD data. Clarke et al. used the GCTA-COJO module for a stepwise conditional analysis, in which disease-associated SNPs are gradually added to the model until there is no SNP significantly associated with SUD [98]. Other functions, such as the one computing LD scores and the one calculating the eigenvectors of GRM, are also included in GCTA [93, 99, 100].

8. POST-GENOMICS AND INTEGRATIVE ANALYSIS

The advent of high-throughput technologies makes it feasible to study multi-level omic data, which includes not only genotype data but also other types of data, such as gene expression and DNA methylation data. An expression Quantitative Trait Loci (eQTL) is genetic loci that regulate gene expression [101]. eQTLs mapping is a useful tool to find genomic locations likely regulating transcript expression. Methods of eQTL mapping have been developed to dissect genetic variants affecting gene expression [102]. The statistical methods, such as single-marker regression analysis, interval mapping, and multiple interval mapping [103], have been widely used for eQTL mapping. The main difference between the eQTL mapping and the QTL mapping method is that a large number of gene expressions are analyzed simultaneously in eQTL mapping. Therefore, the issue of multiple testing need to be considered. Typically, $p$-values corresponding to the peaks of LOD score curves from each transcript are obtained and are adjusted for multiple testing by using FDR [104]. However, this approach only takes LOD score peaks into account [105]. In order to address this limitation, statistical methods, such as Empirical Bayes methods, are developed to control the overall FDR [106]. eQTL mapping provides a list of transcripts that can regulate transcriptional expression. The identification of the hot spots is usually the next task. The easiest way to determine eQTL hot spots is to directly count the number of localized transcripts. However, it can lead to spurious identification or “ghost” hotspots under a number of situations. Therefore, statistical tests, such as a Poisson-based test, are used to determine the true hot spots [104]. Additionally, Kendzierski et al. summarize evidence across every transcript to identify significant eQTL hot spots [106].

In addition to eQTL analysis, integrative analysis combining genetic data with other data sources (e.g. transcriptional, epigenetic, and metabolite data) can provide a comprehensive view of disease biological mechanism [107]. For example, by integrating GWAS and Epigenome-Wide Association Study (EWAS) data, we can not only identify variants at the DNA and epigenetic level but also identify the modulation caused by epigenetic change of genetic variants leading to gene expression alterations. Given the rapid development in both technology and analytical capability, we anticipate that the integrative approach will grow rapidly to provide novel means to study SUD [108, 109].

CONCLUSION

This paper provides an overview of statistical methods and software for data management, linkage analysis, association
analysis and other analysis involved in SUD genetic research. The quality control process is used to identify and remove inaccurate information before the downstream analysis. Stringent quality control ensures the results from the downstream analysis are reliable. In this paper, we summarized some filters commonly used in SUD genetic research and discussed various quality control filters and criteria used by researchers.

The initial understanding of the genetic influences of SUD is assisted by linkage mapping, resulting in the identification of several SUD-associated regions. However, regions identified through linkage analysis could harbor many genes, which makes fine-mapping of the disease-related gene difficult. This problem can be overcome by an association analysis [25]. Depending on the study design, association analysis can be classified into population-based association analysis and family-based association analysis. Population-based association studies have many advantages, such as the easy collection of study samples and high power. Nevertheless, population-based association studies are subject to population stratification issues. Although association analysis has been widely used in SUD genetic research, the identified SUD-related variants only explain a small fraction of the heritability [93]. GCTA is a versatile and user-friendly tool for estimating the heritability explained by all SNPs. The application of GCTA to the SUD genetic dataset suggests that heritability can be explained by small- to medium-effect variants and their possible interactions [94]. With the increased sample size, we are able to detect these small- to medium-effect variants. Meta-analysis is an attractive strategy to combine information from different studies to detect small- to medium-effect variants [32]. Recently, the development of sequencing and other omics technologies have promoted the development of advanced statistical methods and software. There is a pressing need for the development of new analytical tools to keep pace with fast-growing technology, and the increasing amount of SUD omic data. We anticipate that more advanced methods and software will be developed in the future, facilitating the SUD genetic discovery process.

CONSENT FOR PUBLICATION

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CONFLICT OF INTEREST

The authors declare no conflict of interest, financial or otherwise.

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Statistical Methods and Software for SUD

Current Genomics, 2019, Vol. 20, No. 3

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