Review on Databases and Bioinformatic Approaches on Pharmacogenomics of Adverse Drug Reactions

Hang Tong1,2
Nga VT Phan1,2
Thanh T Nguyen3
Dinh V Nguyen4,5
Nam S Vo3
Ly Le1,3

1School of Biotechnology, International University, Ho Chi Minh City, Vietnam; 2Vietnam National University, Ho Chi Minh City, Vietnam; 3Department of Translational Biomedical Informatics, Vingroup Big Data Institute, Hanoi, Vietnam; 4Department of Respiratory, Allergy and Clinical Immunology, Vinmec International Hospital, Hanoi, Vietnam; 5College of Health Sciences, VinUniversity, Hanoi, Vietnam

Abstract: Pharmacogenomics has been used effectively in studying adverse drug reactions by determining the person-specific genetic factors associated with individual response to a drug. Current approaches have revealed the significant importance of sequencing technologies and sequence analysis strategies for interpreting the contribution of genetic variation in developing adverse reactions. Advance in next generation sequencing and platform brings new opportunities in validating the genetic candidates in certain reactions, and could be used to develop the preemptive tests to predict the outcome of the variation in a personal response to a drug. With the highly accumulated available data recently, the in silico approach with data analysis and modeling plays as other important alternatives which significantly support the final decisions in the transformation from research to clinical applications such as diagnosis and treatments for various types of adverse responses.

Keywords: pharmacogenomics, adverse drug reactions, next generation sequencing, genome-wide association study, candidate gene approach

Adverse Drug Reactions

Adverse drug reactions (ADRs) are defined as adverse events that happen to patients after taking certain drugs in clinical treatment.1,2 ADRs can cause failure to almost every organ, but more frequent targets are skin, blood, and liver.3–16 These reactions have been reported to affect 10–20% inpatients and about 25% outpatients17–19 and becoming the major burden of healthcare globally. Edward and Aronson have characterized ADRs into six types from A-F (Table 1), in which two major causes are determined as pharmacologic (type A) and immunologic effects (type B). Some rare ADRs resulted from complementary drug metabolism and immunogenic responses.20 Type A is the most common ADRs which driven by the pharmacodynamics reactions including drug metabolism and transport. This type of ADRs is dose-dependent and predictable, therefore they can be managed by adjusting drug intake. The incidences of this ADR type depend largely on the manifestations of Phase I and Phase II liver enzymes such as cytochrome P450s and glutathione transferases. The majority of ADR type B is drug hypersensitivity reactions (DHRs) which count for about 20% of total ADR cases and are mostly driven by immune system factors such as human leukocyte antigens. The reactions may happen at a very low amount of drug compared with a normal dose, and are classified based on different sorts of...
mechanisms. DHRs are more frequently categorized as immediate and delayed reactions regarding the time course of development.21 Even though the other four types of ADRs were also classified based on the involvement of the associated factors and systematic response to the monitoring methods, the most common ADRs observed from literature are type A (on-target or intrinsic) and type B (off-target or idiosyncratic). The ADRs may appear in patients with mild symptoms from dizziness to very severe syndromes or death, causing much uncomfortability during treatment. In some specific conditions, drugs must be withdrawn and the treatments have to be switched to new therapies. Accurate diagnosis therefore becomes vital to save patients and reduce financial tension. The pharmacogenomic studies of gene–drug relationships bring relevant knowledge to add genetic factors as one to be diagnosed prior to using a drug.

Table 1 ADR Classification

| Type | Clinical Characteristics | Examples | Drugs |
|------|--------------------------|----------|-------|
| A | Pharmacological effect; Predictable; Dose dependent | Thrombolysis, Serotonin syndrome | Antiplatelets, Digoxin |
| B | Caused by Immune – mediation and non-immune –medication; Nonpredictable; Dose independent | MEPE, SCARs | Betalactams, Anticonvulsants |
| C | Mix pharmaco-immuno effect; Chronic; Cumulative dose-related; Manageable by withdrawal | Hypothalamic-pituitary-adrenal axis suppression | Corticosteroids |
| D | Dose – related; Nonmanagable by withdrawal | Teratogenesis | Diethylstilbestrol |
| E | Withdrawal effect; Manageable by slow withdrawal or reintroducing | Opiate withdrawal syndrome | Opiate |
| F | Failure; Dose – related; Often caused by drug interaction; Manageable by changing dose | Inadequate dosage of an oral contraceptive | Contraceptive drugs |

Genetic Predisposition of ADRs

Advances in pharmacogenomics and immunogenomics have revealed the involvement of multiple molecular factors in response to a drug, raising the concept of drug–gene relationship, and moreover, the mechanisms of response. Pharmacogenomics approaches developed to perceive the presentation of genes in a particular group of samples as well as the gene products under certain conditions. Upon exposure to a drug, a particular gene set will express and bring about the products that could be assessed. The completion of the human genome sequencing has brought several advantages to elucidate the relationship between a person’s genome and drug response. Drug metabolizing enzymes cytochrome P450 (CYPs) especially members 1, 2 and 3 families play an important role in drug metabolism and toxicity. CYP2D6, CYP2C19 and CYP2C9 involve in the metabolism of about 80% of therapeutic drugs today. The variation in these genes therefore accounts for most of the ADRs type A in literature. In clinical practice, CYP2D6 biomarkers were observed to link with about 18% of ADR cases reported and suggested by the FDA.22 This gene variation is ethnic-specific and required individual validation in a certain population. In drug hypersensitivity, observational studies in patients treated with some drug groups have shown the high association of genes of HLA classes I and II with the incidence of disease states. For example, HLA-B*15:01 is well-known linked with carbamazepine – inducing SCARs in Japanese and Korean people,23,24 while in South East Asian countries, SCARs to the same drug have prevalently appeared in carriers of HLA-B*15:02 alleles.25–28 Antiretroviral abacavir, on the other hand, is more consistently linked with hypersensitivity in patients carrying HLA-B*57:01 across multiple populations of different origins.29–38 The increasing size and abundance of data accumulated from studies make research in data mining now available for applications.39,40 This review presents the features of current approaches and archive, and state the availability of the data obtained for public share.

Pharmacogenomics Approaches to Study ADRs

Studies on Target Genes Candidates: Replication Approach

More than 95 percent of ADR investigations in Asia are replication approaches.41 A similar situation is also seen in
other regions. The study is based on case-control design where people using the same drug(s) are chosen for investigation. Those subjects that develop adverse reactions are carefully characterized and set as cases, whereas controls are the ones who can metabolize the same drug normally. Key variants that show strong association parameters in genome-wide association study (GWAS) would be selected for replications. The designated genes or variants might also be chosen from previous investigations in closely related populations.

Replication approach can be considered to assess the prevalence of genes in certain groups of patients in the association study. It can also be used to validate a marker in multiple samples. The replication scheme is therefore designed in a variable approach such as combining different polymerase chain reaction (PCR) – based techniques such as PCR followed with restriction fragment length polymorphism (RFLP); conventional PCR with sequencing; or RT-PCR and in silico techniques.\(^{24,42-45}\) Compared with next generation sequencing, this method is much cheaper and totally affordable for research, and it can be scaled and transformed into an application with little optimization.\(^{46,47}\)

Several genetic risk factors responding to drugs were discovered in the gene candidate approach. Abacavir becomes one of the first drugs studied to date to be aware by FDA in patients carrying HLA-B*57:01 alleles (released in 2017). The allele was found more prevalently in abacavir – inducing hypersensitivity patients across continents.\(^{30,34-36,48-50}\) PCR method was repeated simply in several samples using the sequence-specific oligonucleotides followed with sequencing. The consistency of the association between abacavir hypersensitivity and HLA-B*57:01 leads to the general instruction and guideline by FDA to use HLA-B*57:01 test prior to taking this drug. The other example is antiepileptic drug carbamazepine, which has caused highly variable reactions to people in different ethnic groups. Using PCR – based genotyping, HLA-B*15:02 is shown to associate with SJS/TEN in South Asian populations including Han Chinese, Thai, Vietnamese, and Indian,\(^{25-28,35,51,52}\) but in East Asians like Korean and Japanese, the disease is tightly linked with HLA-B*15:01.\(^{11,23,24}\) In addition to the main allele mentioned above for carbamazepine-induced SCAR, HLA-A*31:01 is also associated with the severe skin disease caused by this drug. A detection test for these alleles has been developed to promisingly use as a diagnostic method in screening the carbamazepine sensitive patients before prescription.\(^{53,54}\)

### Table 2 Selected Genes in Replication Approaches

| Drug         | Gene       | Disease         | Population          | Methods                                | Ref.               |
|--------------|------------|-----------------|---------------------|----------------------------------------|-------------------|
| Abacavir     | HLA – B*57:01 | Hypersensitivity | Costa Rica Central, Australia, Italia, Argentina | PCR, sequencing | 30,34-36,48-50 |
| Nevirapine   | CYP2B6 TRAF3IP2 | Hypersensitivity | African             | PCR                                   | 55-57             |
| Carbamazepine| HLA – B*15:02 | SCAR            | Han Chinese, Thai, Vietnamese | Sequence specific oligonucleotide reverse line blots | 25,26,52,54,58 |
|              | HLA-B*15:11 | SJS/TEN         | Japanese            |                                        | 23                |
|              | HLA-A*31:01 | MPE DRESS       | Han Chinese, Vietnamese | PCR                                   | 26,54             |
|              | HLA-B*51:01 | MPE DRESS       | Han Chinese          |                                        | 26                |
| Phenytoin    | HLA - B*15:02 | SCAR            | Malaysia             | PCR                                   | 59                |
|              | HLA – B*15:13 | SCAR            | Malaysia             | PCR                                   | 59                |
|              | HLA-B*13:01, HLA- B*56:02/04, CYP2C19*3 | SCAR | Thailand            | PCR                                   | 60                |
| NSAID        | ALOX15     | Respiratory disease | Spanish            | PCR                                   | 42                |
| Co-trimoxazole | CYP2C9 2*/3* and CYP2C19 3* and NAT2 | Hypersensitivity | UK                    | PCR                                   | 61                |
Table 2 Approaches of GWAS in Drug Hypersensitivity and Outcomes

| Drugs          | Population   | Case | Control | Associated Gene – Disease                                                                 | Ref. |
|----------------|--------------|------|---------|------------------------------------------------------------------------------------------|------|
| **Antiretrovirals** |              |      |         |                                                                                           |      |
| Nevirapine     | Saharan      | 151  | 182     | HLA-C*04:01 - SJS/TEN (rs1265112 and rs746647) within CCHCR1 – skin rash                  | 73   |
|                | African Thai | 72   | 77      |                                                                                           | 74   |
|                |              |      |         |                                                                                           |      |
| **Antibiotics** |              |      |         |                                                                                           |      |
| Sulfonamide    | US           | 91   | 184     | None – hypersensitivity                                                                     | 75   |
| Dapsone        | Chinese      | 39   | 833     | HLA-B*13:01 – dapsone hypersensitivity                                                      | 76   |
| Beta lactam -  | Spain        | 387  | 1124    | Rs4958427 of ZNF300 – penicillin allergy                                                    | 77   |
| Penicillin     | Italy        | 299  | 362     | rs17612 of C5 rs7754768 and rs926832 of the HLA-DRA | HLA-DRB5 interregion rs7192 of HLA-DRA |      |
| **NSAID**      |              |      |         |                                                                                           |      |
| NSAID          | Spanish      | 112  | 124     | None but suggestive regions of RIMS1, BICCI and RAD51L 1 – urticarial/angiokeratoma        | 78   |
|                | Han Chinese  | 120  | 101     | None but suggestive regions of ABIBP - urticarial/angiokeratoma                            | 78   |
| Aspirin        | Korean       | 117  | 685     | HLA-DRB1 rs1042151 - Respiratory disease                                                    | 79   |
|                | Korean       | 117  | 691     | SBF1 – asthma rs4917014 of IKZF1 SCAR                                                      | 80   |
| Cold medicine  | Japanese     | 117  | 691     |                                                                                           | 81   |
| **Anticonvulsant** |            |      |         |                                                                                           |      |
| Lamotrigine    | UK           | 46   | 1296    | None – hypersensitivity                                                                     | 82   |
| Phenytoin      |              | 44   |         |                                                                                           | 83   |
| Phenytoin      | Taiwan       | 34   | 1214    | CYP2C9*3 – SCAR rs12668095 near CRAMP1/1M3M204/1FT140/HNL                                  | 84   |
| Lamotrigine    | Korean       | 34   | 1214    | rs79007183 near TNS3 skin rash                                                             | 85   |
| Carbamazepine  | Japanese     | 53   | 882     | HLA-A*31:01 – SCAR                                                                         | 86   |
| Carbamazepine  | European     | 65   | 3987    | HLA-A*31:01 – immediate and delayed hypersensitivity (including SCAR)                     | 87   |
| Various drugs  | Caucasian    | 96   | 198     | None – SJS/TEN                                                                              | 88   |
| Others         |              |      |         |                                                                                           |      |
| Allopurinol    | Japanese     | 589  | 3308    | SJS/TEN HLA-B*58:01 rs6021191 variant in NFATC2 rs17885382 in HLA-DRB1                    | 89   |
| Asparaginase   | US           |      |         |                                                                                           |      |

Using PCR in case–control design, several gene-drug associations have been revealed, providing the basis for syndrome – drug linkage using amplification methods (Table 2, Table 3).

Accumulated evidences from individual researches across populations reveal the geographic difference in genetic variability. The association of a gene to a drug response in different populations are varied along with variable allele frequency, haplotype frequency and linkage disequilibrium. Several diseases are multifactorial that require the combinational interpretation of many genes. The meta-analysis of whole exome data from six ethnic populations from different parts of the world shows the consistent findings in drug-related genes. About half of the functional variants of these genes are unique to one of 6 populations studied. In addition, amongst the drug-related genes analyzed, CYPs and phase II enzyme have highest difference in cumulative allele probability (CAP), indicating the variation in possibility that a functional variant affects the drug response phenotypes. Although the analysis was based on
only the exome data, the major outcome obtained from this result is crucial for each country to build the own pharmacogenomic database. The availability of this data would get us closer to personalized medicine when doctors can more precisely predict the effect of a drug to a patient and choose the best medication for a person.

**Genome-Wide Association Studies**

Genome-wide association study (GWAS) has shown to be one of the very effective approaches to screen for the risk factors associated with a certain disease.63 To date, there are 4054 publications and almost 140 thousand associations have been published in the GWAS catalog.64 The approaches are designed with a case–control model followed with next generation sequencing and SNP calling against the reference genome. At this present time, the enormous number of SNPs would be obtained and sent for further evaluation of association analysis. Since GWAS evaluated a large number of SNPs, it requires a much larger number of samples to achieve the statistical reliability.65–69 The correlation between the number of SNP and sample size is calculated. It is estimated that testing a single SNP marker requires 248 cases, while testing 500,000 SNPs and 1 million markers requires 1206 cases and 1255 cases, respectively, under the assumption of an odds ratio of 2, 5% disease prevalence, 5% minor allele frequency, complete linkage disequilibrium (LD), 1:1 case/control ratio, and a 5% error rate in an allelic test.70

Recruitment of the large sample in research requires big financial support as well as expertise in data analysis. For this reason, GWAS is preferably used for emerging diseases or complex traits that cannot be simply understood by a single gene phenotype. Specific variants were found in different cohorts providing that risk factors could be thoroughly scanned and validated. GWAS can be designed to integrate with pharmacodynamics to study the drug response in either prospective or retrospective approaches. The suspected drug is administered at common or tested doses and given to individuals. People are recruited based on their response and their genotypes are scanned for all SNPs, and phenotypes are observed through the response of each individual under drug conditions.71 In common design for ADR study, patients are selected with a similar phenotype that demonstrating the characteristics of the same disease in response to a certain drug, compared with the control subjects who take the same prescription and well improved without adverse reactions (drug tolerant). The statistical analysis is then performed to find the association based on genome wide significance, predictive values, et etc.72 The anticonvulsants, antimicrobial and NSAID are among the most causative drugs of ADRs, and were among the first drugs studied and analyzed in GWAS (Table 2).

Several factors have been discussed for the better improvement of the GWAS significance. The sample size is one of the important factors. In the standard approach mentioned, the number of subjects recruited in the study should reach the required quantity in which it could cover the number of possible variants.90 However, in many rare syndromes, recruiting adequate samples seems impossible in a given time. It is therefore requiring consecutive phases of research in which sample collection needs to be completed prior to implementing all other research steps. Multiple attempts have been made to improve the reliability and accuracy of the genome-wide analysis even with a small reference population.91 The findings discussed the adjustment of the model, in which after testing in reference data set of population, both theory and empirical observation from simulation agreed well in the population samples with a high degree of relatedness. These results suggested that higher significance would be obtained in the subjects with the same ethnic origin than in the mixed groups such as meta-analysis. For that, the accuracy and significance at genome-wide scale are still obtained even in the cohort with a small number of subjects (as 50).

Design is another important feature besides sample size added to the success of GWAS. Several samples such as some examples explained above cannot get the genome wide significance assigned by the International Hapmap consortium92 assuming that an OR of 2, MAF 5%, disease prevalence 5% and complete linkage disequilibrium. The outcome might be due to the low allele frequency. The associated p-value suggested 5 × 10⁻⁸ is valid for common variants with MAF ≥ 5%. When analyzing variants with lower MAF values such as 1%, 0.5% or 0.1%, the model showed the genome wide significance with p values are 3 × 10⁻⁸, 2 × 10⁻⁸ and 1 × 10⁻⁸, respectively. The inclusion of LD was indicated not necessary as the model can analyze all variants even they have complete linkage disequilibrium (r² =1).93

**Data Mining and in silico Approach**

Although replication approaches have contributed to the mass data accumulation through scanning several cohorts, adding to better understanding the genetics of population,
the research design should be utilized with thorough consideration in different aspects.

First, the selected candidate gene should be chosen from the combinational results of genome scanning and/or tested gene of the whole population. The high association of a single gene with the disease might not completely interpret the cause–effect relationship. Pan reported an HLA-B*15:02 negative - case in Taiwan population who developed hypersensitivity in response to carbamazepine, suggesting that, the insight about the genetics of a common disease contributes as part of disease development. In the Thai cohort, screening of the HLA genes could help to protect only 22% of SCAR Thai patients (mainly allopurinol - and carbamazepine – inducing) whereas using drug-induced IFN-γ-specific cells scan, approximately 46% of patients with SCAR were detected positive. Second, with the complication of the metabolism, approaching a real system to study mechanism sometimes is not possible. The virtual platform may be one of the very good options to try.

The in silico study of carbamazepine and SCAR using pooled data from Asian populations have proved that the polymorphic alleles themselves may not be sufficient to explain the clinical outcomes, instead, the proteins or complex combination of multiple factors could efficiently help to increase the predictive value. For example, when aligning available HLA – B75 protein members on the crystal structure of HLA – B*15:01, all except HLA – B*15:21 imposed and fit on each other. The molecular docking of the chemical structure of carbamazepine to the antigen-presenting sites of tested HLAs could determine the drug binding amino acids on HLA protein. This finding explains for the case reported, in which the HLA – B*15:02 negative but HLA – B*15:21 positive patients still developed SCAR when taking carbamazepine. When both alleles belong to the HLA-B75 family, the presence of one and/or another causes hypersensitivity. More investigation combining gene detection and in silico studies have been implemented for dapsone or NSAID.

Analyzing ADRs Using Next Generation Sequencing Data

Data Generation and Collection

In recent years, high-throughput technologies have accelerated genomics/pharmacogenomics studies and resulted in large-scale data. Next-generation sequencing technologies (NGS) such as those provided by Ion Torrent or Illumina platforms are becoming the most common way to get genomics data. Such technologies enabled genomic sequencing on a massive scale at a low-cost and high-quality, which enabled genome-wide studies in large-scale cohorts. Recently, third-generation sequencing technologies (TGS) such as those provided by Pacific Biosciences or Oxford Nanopore Technologies are also exploited in some recent genomics/pharmacogenomics studies. Such technologies could give read lengths of around tens to even hundreds of thousands of bases although with high-cost and somewhat low-quality compared to second-generation technologies. By generating long reads, TGS has clear advantages in resolving highly repetitive or polymorphic regions which are quite common in pharmacogenomics and immunogenomics studies.

Available Databases for ADRs Studies

Large-scale data from pharmacogenomics studies have been collected and managed under large consortia and networks such as Clinical Pharmacogenetic Consortium (CPIC), Pharmacogenomics Research Network (PGRN), or The South East Asian Pharmacogenomic Network (SEAPHARM). More details of these consortia and networks are shown in Table 3. Some recent work such as The Observational Medical Outcomes Partnership–Common Data Model (OMOP-CDM) provides clinical data sources such as electronic health records (EHR) from which ADR-related information can be extracted. Specific databases such as FDA Adverse Event Reporting System (FAERS), Side Effect Resource (SIDER), or Healthcare Cost and Utilization Project (HCUP) provide public datasets that can be used to analyze ADRs and support controlling ADEs. Currently, many databases have been integrated into larger ones such as PharmGKB. PharmGKB also provides a number of datasets and annotations for drugs that have shown adverse reactions, many of them have been pharmacogenetically tested. Some others such as CTD, KEGG, or SuperTarget provide non-clinical data for metabolic pathways, or drug metabolism, interactions in the molecular structures of proteins, or associations of drugs-drugs, drugs-genomes. In addition, immunogenomics databases such as HLA-ADR provide allele frequency and haplotype of HLA genes that have been associated with ADRs. These databases can be used to help minimize ADRs in drug design or patient treatment. Some general databases such as dbSNP or the Database of Genomic Variants can be also used as
**Table 3 PGx Consortia and Networks**

| PGx Consortia and Networks | Description | URL |
|-----------------------------|-------------|-----|
| PharmGKB                    | PharmGKB is a database that collects and curates knowledge of human genetic variation on drug responses. The database provides information about 706 annotated drugs and 149 curated pathways. There are currently 155 annotations for clinical guidelines and 753 annotations for drug labeling. The database also provides 4,570 clinical annotations and 23,938 variant annotations (accessed on July 15th). | https://www.pharmgkb.org/ |
| Clinical Pharmacogenetics Implementation Consortium (CPIC) | CPIC is an assessment organization with updated information on clinical findings and laboratories in the field of pharmacogenomics. CPIC has provided 24 guidelines of 20 genes and 62 drugs to address and breakdown barriers in clinical implementation, reducing “one size fit all” status, and optimizing drugs in precision medicine (accessed on July 15th). | https://cpicpgx.org/ |
| Dutch Pharmacogenetics Working Group (DPWG) | The aim of DPWG consortium is to provide well-known PGx clinical testing to translate genotype to phenotype. With more than 90 clinical guidelines, annotation validated by DPWG will be formulating hypotheses to support clinical implementation or in silico related pharmacogenomics. | https://www.pharmgkb.org/page/dpwg |
| Ubiquitous Pharmacogenomics (U-PGx) | U-PGx was established by European experts to implement a pre-emptive pharmacogenomics approach. A panel of 13 PGx genes with 50 variants helps to study the genetic factors that influence the patient’s response to medication, with the aim of improving the quality of life, reducing costs, and giving better results for patients. | http://upgx.eu |
| ClinGen PGx Working Group | As a data center to support clinical practitioners, researchers with genomic and phenotypic information to help interpret gene factors. There are currently more than 1750 curated genes, 50 expert groups and 11,413 experts for the development of bioinformatics tools and increasing accuracy in the fairly healing process. | https://www.clinicalgenome.org/working-groups/ |
| PGRN-RIKEN | PGRN-RIKEN is a collaboration between Pharmacogenomics Research Network (PGRN) and RIKEN in the use of patient samples and drug response for collaborative research, involving adverse drug response. | https://www.pgrn.org/pgrn-riken.html |
| Canadian Pharmacogenomics Networks for Drug Safety (CPNDS) | CPNDS was founded in 2004 with the goal of building guidelines (8 guidelines – updated 07/03/2020) related to PGx response and ADR. Learn and assess risks to genetic factors, develop PGx clinical implementation tools to support and optimize drug use. | http://cpnds.ubc.ca/ |
| PharmVar | PharmVar is a repository of pharmacogenomics variation that supporting the defined haplotype and alleles, focusing on human cytochrome P450 families and NUDT15. A comprehensive database providing information for pharmacogenomics Knowledge (PharmGKB) and the Clinical Pharmacogenetics Implementation Consortium (CPIC). | https://www.pharmvar.org/ |
| European Pharmacogenetic Implementation Consortium (EU – PIC) | A group for clinical implementation of many European countries to improve the treatment from pharmacogenomics guideline into clinical care. | http://www.eu-pic.net |
| Southeast Asian Pharmacogenomic Research Network (SEAPHARM) | SEAPharm was established to enable PGx research among the various communities within but not limited to countries in South East Asia, with the ultimate goal to support PGx implementation in the region. | http://www.pharmagtc.org/seapharm/ |
| Database genomic variant | DGV provides the archiving, accessioning and distribution of public available genomic structural variant in all species. | https://www.ebi.ac.uk/dgva/ |

(Continued)
reference for pharmacogenomics studies, eg, to check allele frequency, genotype, and annotation. Figure 1 shows an overview of data, methods and resources for ADR studies.

**Data Analysis**
With the advances in high-performance computing and big data analytics, we can take advantage of large-scale chemical, biological, and biomedical information to discover complex genetic mechanisms of ADRs. Nevertheless, while sequencing data are now easier and cheaper to produce, analyzing such data is still the bottleneck. In the following paragraphs, we will focus on two types of analyses: detecting genomics/pharmacogenomics variants/haplotypes and annotating them for ADR studies. State-of-the-art bioinformatics tools/pipelines for such analyses and emerging machine learning techniques to classify genomic variants into ADRs were summarized in Table 4.

**Genetic Variant Calling**
The GATK best practices are often recommended to use for variant calling on NGS data, together with BWA-MEM for reading alignment. However, some others such as DeepVariant or Novoalign (http://novocraft.com) can be used as complementary for BWA-MEM or even replacement in some cases to improve performance. This strategy can be widely applied for WGS, WES or sequencing data generated from gene panels. The called variants often serve as a starting point for downstream analyses to study ADRs.

**PGx Haplotyping**
Due to the complexity of PGx regions, general haplotype callers often have limitations in determining PGx star alleles. As a result, specific tools such as Astrolabe, Cyrius, Aldy, or Stargazer are often used. However, most of the current PGx haplotyping tools are still limited to detecting star alleles in a subset of PGx genes, eg, Cyrius works on only gene Cytochrome P450 2D6 (CYP2D6), while Stargazer covers only 51 PGx genes. Developing more comprehensive and accurate tools is still an urgent need.

**HLA Typing**
The complexity and high polymorphisms of HLA regions make HLA typing always challenging, especially for NGS data. HLA typing tools for NGS data such as PHLAT, Polysolver, OptiType, xHLA, or Kourami can get HLA alleles with 4-digits, 6-digits, up to 8-digits using WGS/WES data. These tools are often limited to detecting known HLA alleles. For RNA-seq data, seq2HLA can be used to get 4-digits resolution. Despite a lot of efforts, current tools for HLA typing still suffer from detecting novel or class II HLA alleles.

**PGx Annotation**
Variant annotation is the process of determining the effects of genetic variants on disease and genes. Annotation for PGx variants can be obtained using general tools for annotation of genetic variants such as VEP, SnpEff/SnpSift, Annovar, or Intervar. More informative annotation of PGx genes could be obtained using specific tools such as PharmCAT, which was built based on CPIC’s guidelines. Such guidelines link genotypes to phenotypes and prescribing recommendations based on genotype/phenotype. Recently, PGx guidelines are provided by CPIC, DPWG, CPNDS, and PharmVar are still the gold standard. Due to the limited number of guidelines, such annotation tools can cover only a small subset of PGx genes. The current version of PharmCAT can detect only 12 guidelines out of 64 very important pharmacogenomics (VIPs) of more than 100 known PGx genes.

**Classifying Genomic Variants into ADRs**
The American College of Medical Genetics and Genomics and Association of Molecular Pathology (ACMG-AMP) has developed guidelines for standardizing and improving disease classification based on genomic variants, which
can be used for studying ADRs.\textsuperscript{125,126} Some tools aim to predict phenotype from genotype using activity score assigned based on allele frequency information from CPIC.\textsuperscript{127} Some others used hierarchical or k-means clustering to detect the correlation between genotype and phenotype.\textsuperscript{128} Some recent tools such as Hubble have applied deep learning techniques to predict the functions of PGx alleles.\textsuperscript{129} Nevertheless, classifying genomic variants to explore the correlation of genes related to ADRs is still challenging and needs significant improvement.

**Put-All-Together**

A number of data analysis pipelines have been developed to detect and annotate ADR-associated variants and haplotypes. A pipeline for analyzing three genes CYP2C9, CYP2C12, and HLA using WGS/WES/genotyping data have been carried out in.\textsuperscript{130} They have extracted 39 variants from whole exome sequencing data of 1585 individuals, then haplotype was assigned based on U-PGx translation. Currently the pipeline developed by PGRN seems to be a gold standard for PGx data analysis.\textsuperscript{131} Meanwhile, some other groups such as RIKEN-Pharmacogenomics Laboratory, SEAPHARM also built their own pipelines for pharmacogenomics data analysis (personal discussion).

Nevertheless, accurate identification of haplotype in pharmacogenomics is still challenging, especially with highly polymorphic regions such as HLA or CYP2D6. Many haplotypes/diplotypes of PGx genes are still unknown. Annotation of PGx genes including novel alleles is still challenging, and there is a need for developing better annotation tools. The gene–drug interaction guidelines are now still very limited, eg, the CPIC database provides only 12 specific guidelines to support optimizing drug therapy. Furthermore, the genotype-phenotype association is still uncertain in many cases due to many factors.
Table 4 Common Tools Used for Data Analysis

| PGx Variants and Haplotypes Calling | Description | Ref. |
|-------------------------------------|-------------|------|
| GATK                               | A standard tool for variant calling, support Whole Genomes/Exomes, Gene Panels, RNA-seq and Targeted Sequencing. | [https://gatk.broadinstitute.org/hc/en-us](https://gatk.broadinstitute.org/hc/en-us) |
| BWA                                | A standard tool for aligning short genomic sequences to large reference sequences such as human genome. | [http://bio-bwa.sourceforge.net/bwa.shtml](http://bio-bwa.sourceforge.net/bwa.shtml) |
| DeepVariant                         | A variant calling tool which applies convolutional neural network approach for identifying genomic variants. | [https://github.com/google/deepvariant/](https://github.com/google/deepvariant/) |
| Novoalign                          | An accurate tool for aligning short reads to large reference genomes | [http://www.novocraft.com/](http://www.novocraft.com/) |
| Astrolabe                          | A tool for star allele calling which was initially developed for the CYP2D6 gene, then extended to CYP2C9 and CYP2C19 and other genes | [https://www.nature.com/articles/npgenmed201639](https://www.nature.com/articles/npgenmed201639) |
| Stargazer                          | A tool for calling star alleles (haplotypes) in PGx genes using data from NGS or SNP array. | [https://stargazer.gs.washington.edu/stargazer.web/](https://stargazer.gs.washington.edu/stargazer.web/) |
| HLA typing                         | | |
| Seq2HLA                            | RNA-seq; iterative allele inference (greedy); 4-digit resolution | [https://bio.tools/seq2hla](https://bio.tools/seq2hla) |
| Kourami                            | WGS; discovery of novel alleles; up to 6-digit resolution | [https://bio.tools/kourami](https://bio.tools/kourami) |
| Polysover                          | WES; k-mer seeding to get HLA reads; Bayesian inference for inferring best alleles: up to 6-digit resolution | [https://github.com/researchapps/polysover](https://github.com/researchapps/polysover) |
| HLA-HD                             | WGS, WES, RNA-seq; discovery of novel alleles; up to 6-digit resolution | [https://www.genome.med.kyoto-u.ac.jp/HLA-HD/](https://www.genome.med.kyoto-u.ac.jp/HLA-HD/) |
| Optitype                           | RNA-seq, WGS, WES; ILP solving to aligned reads for best alleles; 4-digit resolution | [https://github.com/FRED-2/OptiType](https://github.com/FRED-2/OptiType) |
| PGx annotation                     | | |
| VEP                                | A tool for predicting effects of variants including SNPs, Indels, CNVs or Structural Variants; work with genes, transcripts, protein sequences and regulatory regions. | [https://asia.ensembl.org/info/docs/tools/vep/index.html](https://asia.ensembl.org/info/docs/tools/vep/index.html) |
| Annovar                            | An annotation tool which can identify protein coding changes through the transformation of SNVs, CNVs | [http://wannovar.wglab.org/](http://wannovar.wglab.org/) |
| SnpEff/SnpSift                      | SnpEff is a tool for variant effect annotation and prediction, particularly on genes and proteins. SnpSift is a tool for genomic variant annotation, using annotated databases. The latter is often used after the former to find the most significant variants. | [https://pcingolga.github.io/SnpEff/](https://pcingolga.github.io/SnpEff/) |
| Intervar                           | A tool for clinical interpretation of genetic variants based on ACMG/AMP guidelines. | [http://wintervar.wglab.org/](http://wintervar.wglab.org/) |
| PharmCAT                           | A tool for translation of genotype to phenotype using genotyping and sequencing data based on CPIC guidelines. | [http://pharmcat.org/](http://pharmcat.org/) |

such as age, gender, the interaction between drug molecules. Developing tools to capture such uncertainty is pretty necessary but very challenging.

Suggestion for Clinical Applications

We have reviewed the typical works in pharmacogenomics to detect adverse drug reactions, a key task in the post-market drug safety surveillance. It is well-known that datadriven and data integration approach is essential in solving that problem. Advance in next generation sequencing technology and abundance of genomic data has led to the great outcomes from pharmacogenomic research in drug response to the transformation into clinical applications. A number of drugs have been labeled which concern patients to take the genetic test prior to prescription. Specific programs have also been implemented in particular countries where population
Pharmacogenomics have been well analyzed. The US FDA Sentinel Initiative (https://www.sentinelinitiative.org) or The UK CPRD (https://www.cprd.com) are among the programs which have been implemented. These are successful examples for translating research into clinical guidelines for patients or populations who suspect to use specific drugs in the future. For emerging countries, the introduction and improvement of technologies as well as the gradual enlarging of available data bring a number of options of choice. Our review has presented the characteristics, requirements and possible outcomes of some popular and powerful approaches for future applications. Using either one or another depends a lot on the prevalence of current data, design and expectation, however, they can be combined to have the best interpretation of the mechanism of drug response or association of genetic components with a disease, therefore suggest the most effective risk factors for diagnosis or strategic therapy.

Enriching knowledge and attitude towards implementation of pharmacogenomic services among clinicians and administrative staff are crucial for clinical application. In addition, expanding the list of drug-gene pairs covered by national health policy is feasible goal for the near future when sufficient pharmacogenomic and health economic data are generated to aid in the decision-making process. Efforts to implement pharmacogenomics into clinical practice are proceeding at different rates in different countries. It is important to realize the potential benefits of pharmacogenomic implementation on the healthcare system. Effective data and experience sharing at international levels would reduce unnecessary healthcare costs from inefficient or inappropriate drug therapies, maximize the effectiveness of drugs and minimize adverse drug events.

**Abbreviations**

ADR, adverse drug reaction; DHR, drug hypersensitivity reaction; Ig, immunoglobulin; MPE, maculopapular exanthema; FDE, fixed drug eruption; SCAR, severe cutaneous adverse reaction; DIHS, drug induced hypersensitivity syndrome; DRESS, drug reaction with eosinophilia and systemic symptoms; SJS, Steven Johnson syndrome; TEN, toxic epidermal necrolysis; AGEP, acute generalized exanthematous pustulosis; NSAID, nonsteroidal anti-inflammatory drug; HLA, human leucocyte antigen; GWAS, genome wide association study; WGS, whole genome sequencing; WES, whole exom sequencing; PGx, pharmacogenes; CYP, cytochrome; MAF, minor allele frequency; HWE, Hardy Weinberg equilibrium; OR, odd ratio; LD, linkage disequilibrium; SNP, single nucleotide polymorphism; PCR, polymerase chain reaction; FDA, Food and Drug Administration; RFLP, restriction fragment length polymorphism.

**Acknowledgment**

This work is partially funded by Vingroup Big Data Institute and Vietnam National University under grant number GEN2019-28-01/HD-KHCN.

**Disclosure**

The authors declare that they have no conflicts of interest.

**References**

1. Wheatley LM, Plaut M, Schwaninger JM, et al. Report from the National Institute of Allergy and Infectious Diseases workshop on drug allergy. *J Allergy Clin Immunol*. 2015;136(2):262–71.e2. doi:10.1016/j.jaci.2015.05.027
2. Utrecht J. Idiosynhrastic drug reactions: current understanding. *Annu Rev Pharmacol Toxicol*. 2007;47(1):513–539. doi:10.1146/annurev.pharmtox.47.120505.105150
3. Hunziker T, Künzi U-P, Braunschweig S, et al. Comprehensive hospital drug monitoring (CHDM): adverse skin reactions, a 20-year survey. *Allergy*. 1997;52(4):388–393. doi:10.1111/j.1398-9995.1997.tb01017.x
4. Valeyrue-Allanore L, Sassolas B, Roujeau JC. Drug-induced skin, nail and hair disorders. *Drug Saf*. 2007;30(11):1011–1030. doi:10.2165/00002098-200730110-00003
5. Sidenius S, PCT-P, Dam M. Lamotrigine-induced rash–worth a rechallenge. *Acta Neurol Scand*. 2005;111(3):191–194. doi:10.1111/j.1600-0404.2005.00381.x
6. Vonakis BM, Saini SS. New concepts in chronic urticaria. *Curr Opin Immunol*. 2008;20(6):709–716. doi:10.1016/j.coi.2008.09.005
7. Pereira FA, Mudgil AV, Rosmarin DM. Toxic epidermal necrolysis. *J Am Acad Dermatol*. 2007;56(2):181–200. doi:10.1016/j.jaad.2006.04.048
8. Downey A, Jackson C, Harun N, et al. Toxic epidermal necrolysis: review of pathogenesis and management. *J Am Acad Dermatol*. 2012;66(6):995–1003. doi:10.1016/j.jaad.2011.09.029
9. Tang YH, Mockenhaupt M, Henry A, et al. Poor relevance of a lymphocyte proliferation assay in lamotrigine-induced Stevens-Johnson syndrome or toxic epidermal necrolysis. *Clin Exp Allergy*. 2012;42(2):246–254. doi:10.1111/j.1365-2222.2011.03875.x
10. Wei C-Y, Chung W-H, Huang H-W, et al. Direct interaction between HLA-B and carbamazepine activates T cells in patients with Stevens-Johnson syndrome. *J Allergy Clin Immunol*. 2012;129(6):1562–1565. doi:10.1016/j.jaci.2011.12.990
11. Bjornsson E. Review article: drug-induced liver injury in clinical practice. *Aliment Pharmacol Ther*. 2010;32(1):3–13. doi:10.1111/j.1365-2036.2010.04320.x
12. Clay KD, et al. Brief communication: severe hepatotoxicity of telithromycin: three case reports and literature review. *Ann Intern Med*. 2006;144(6):415–420. doi:10.7326/0003-4819-144-6-200503210-00121
13. Orman ES, Conjeevaram HS, Vuppallanchi R, et al. Clinical and histopathologic features of fluoroquinolone-induced liver injury. *Clin Gastroenterol Hepatol*. 2011;9(6):517–523.e3. doi:10.1016/j.cgh.2011.02.019
49. Moragas M, Belfoso WH, Baquedano MS, et al. Prevalence of HLA-B*57:01 allele in Argentinian HIV-1 infected patients. Tissue Antigens. 2015;86(1):28–31. doi:10.1111/tan.12575

50. Dello Russo C, Lisi L, Lofaro A, et al. Novel sensitive, specific and rapid pharmacogenomic test for the prediction of abacavir hypersensitivity reaction: HLA-B*57:01 detection by real-time PCR. Pharmacogenomics. 2011;12(4):567–576. doi:10.2217/ pg.10.208

51. Aggarwal R, Sharma M, Modi M, et al. HLA-B*1502 is associated with carbamazepine induced Stevens-Johnson syndrome in North Indian population. Hum Immunol. 2014;75(11):1120–1122. doi:10.1016/j.humin.2014.09.022

52. Nguyen DV, Chu HC, Nguyen DV, et al. HLA-B*1502 and carbamazepine-induced severe cutaneous adverse drug reactions in Vietnamese. Asia Pac Allergy. 2015;5(2):68–77. doi:10.5415/apallergy.2015.5.2.68

53. Mushiroda T, et al. Association of HLA-A*31:01 screening with the incidence of carbamazepine-induced adverse reactions in a Japanese population. JAMA Neurol. 2018;75(7):842–849.

54. Nguyen DV, Vidal C, Chi HC, et al. A novel multiplex polymerase chain reaction assay for detection of both HLA-A*31:01/ HLA-B*15:02 alleles, which confer susceptibility to carbamazepine-induced severe cutaneous adverse reactions. HLA. 2017;90(6):335–342. doi:10.1111/tan.13143

55. Cicciacci C, et al. Association between CYP2B6 polymorphisms and Nevirapine-induced SJS/TEN: a pharmacogenetics study. Eur J Clin Pharmacol. 2013;69(11):1909–1916. doi:10.1007/s00228-013-1549-9

56. Cicciacci C, Rufini S, Mancinelli S, et al. A pharmacogenetics study in Mozambican patients treated with nevirapine: full re-sequencing of TRAF3IP2 gene shows a novel association with SJS/TEN susceptibility. Int J Mol Sci. 2015;16(3):5830–5838. doi:10.3390/ijms16035830

57. Carr DF, Chappona M, Cornejo D, et al. CYP2B6 c.983T>C polymorphism is associated with nevirapine hypersensitivity in Malawian and Ugandan HIV populations. J Antimicrob Chemother. 2014;69(12):3329–3334. doi:10.1093/jac/dku315

58. Cheung Y-K, Cheng S-H, Chan EJM, et al. HLA-B alleles associated with severe cutaneous reactions to antiepileptic drugs in Han Chinese. Epilepsia. 2013;54(7):1307–1314. doi:10.1111/epi.12217

59. Chang C-C, Ng C-C, Too C-L, et al. Association of HLA-B*15:13 and HLA-B*15:02 with phenytoin-induced severe cutaneous adverse reactions in a Malay population. Pharmacogenomics J. 2017;17(2):170–173. doi:10.1007/s13289-016-0120-4

60. Tassaneeyakul W, Prabmeechai N, Sukasem C, et al. Associations between HLA class I and cytochrome P450 2C9 genetic polymorphisms and phenytoin-related severe cutaneous adverse reactions in a Thai population. Pharmacogenet Genomics. 2016;26(5):225–234. doi:10.1097/FPC.0000000000000211

61. Pirmohamed M, Alfirevic A, Vilar J, et al. Association analysis of drug metabolizing enzyme gene polymorphisms in HIV-positive patients with co-trimoxazole hypersensitivity. Pharmacogenetics. 2000;10(8):705–713. doi:10.1097/00008571-200011000-00005

62. Scharfe CPI, Tremmel R, Schwab M, et al. Genetic variation in human drug-related genes. Genome Med. 2017;9(1):117. doi:10.1186/s13073-017-0502-5

63. Low S-K, Takahashi A, Mushiroda T, et al. Genome-wide association study: a useful tool to identify common genetic variants associated with drug toxicity and efficacy in cancer pharmacogenomics. Clin Cancer Res. 2014;20(10):2541–2552. doi:10.1158/1078-0432.CCR-13-2755

64. Buniello A, et al., The NHGRI-EBI GWAS Catalog of published genome-wide association studies, targeted arrays and summary statistics 2019. Nucleic Acids Res. 2019. 47(D1):D1005–D1012.

65. Klein RJ. Power analysis for genome-wide association studies. BMC Genet. 2007;8(1):58. doi:10.1186/1471-2156-8-58

66. Zondervan KT, Cardon LR. Designing candidate gene and genome-wide case-control association studies. Nat Protoc. 2007;2(10):2492–2501. doi:10.1038/nprot.2007.366

67. Spencer CC, et al. Designing genome-wide association studies: sample size, power, imputation, and the choice of genotyping chip. PLoS Genet. 2009;5(5):e1000477.

68. Wu Z, Zhao H, Walsh B. Statistical power of model selection strategies for genome-wide association studies. PLoS Genet. 2009;5(7):e1000582. doi:10.1371/journal.pgen.1000582

69. Park J-H, Wacholder S, Gail MH, et al. Estimation of effect size distribution from genome-wide association studies and implications for future discoveries. Nat Genet. 2010;42(7):570–575. doi:10.1038/ng.610

70. Hong EP, Park JW. Sample size and statistical power calculation in genetic association studies. Genomics Inform. 2012;10(2):117–122. doi:10.5808/GI.2012.10.2.117

71. Wu R, Tong C, Wang Z, et al. A conceptual framework for pharmacodynamic genome-wide association studies in pharmacogenomics. Drug Discov Today. 2011;16(19–20):884–890. doi:10.1016/j.drudis.2011.09.001

72. Hayes B. Overview of statistical methods for Genome-Wide Association Studies (GWAS). Methods Mol Biol. 2013;1019:149–169.

73. Carr DF, Bourgeois S, Chaponda M, et al. Genome-wide association study of nevirapine hypersensitivity in a sub-Saharan African HIV-infected population. J Antimicrob Chemother. 2017;72(4):1152–1162. doi:10.1093/jac/dku545

74. Chantarangsu S, Mushiroda T, Mahasirimongkol S, et al. Genome-wide association study identifies variations in 6p21.3 associated with nevirapine-induced rash. Clin Infect Dis. 2011;53(4):341–348. doi:10.1093/cid/cir403

75. Reinhart JM, Motisinger-Reif A, Dickey A, et al. Genome-wide association study in immunocompetent patients with delayed hypersensitivity to sulfonamide antimicrobials. PLoS One. 2016;11(6):e0156000. doi:10.1371/journal.pone.0156000

76. Zhang F-R, Liu H, Irwanto A, et al. HLA-B*13:01 and the Dapsone Hypersensitivity Syndrome. N Engl J Med. 2013;369(17):1620–1628. doi:10.1056/NEJMoa1213096

77. Gueant JL, Romano A, Cornejo-Garcia J-A, et al. HLA-DRA variants predict penicillin allergy in genome-wide fine-mapping genotyping. J Allergy Clin Immunol. 2015;135(1):253–259. doi:10.1016/j.jaci.2014.07.047

78. Cornejo-Garcia JA, Liu L-B, Blanca-López N, et al. Genome-wide association study in NSAID-induced acute urticaria/angioedema in Spanish and Han Chinese populations. Pharmacogenomics. 2013;14(15):1857–1869. doi:10.2217/pg.13.166

79. Park BL, Kim T-H, Kim J-H, et al. Genome-wide association study of aspirin-exacerbated respiratory disease in a Korean population. Hum Genet. 2013;132(3):313–321. doi:10.1007/ s00439-012-1247-2

80. Kim J-H, Park B-L, Cheong HS, et al. Genome-wide and follow-up studies identify CEP68 gene variants associated with risk of aspirin-intolerant asthma. PLoS One. 2010;5(11):e13818. doi:10.1371/journal.pone.0013818

81. Ueta M, Sotozono C, Nakano M, et al. Association between prostaglandin E receptor 3 polymorphisms and Stevens-Johnson syndrome identified by means of a genome-wide association study. J Allergy Clin Immunol. 2010;126(6):1218–25e10. doi:10.1016/j.jaci.2010.08.007

82. McCormack M, Urban TJ, Shiami KV, et al. Genome-wide mapping for clinically relevant predictors of lamotrigine- and phenytoin-induced hypersensitivity reactions. Pharmacogenomics. 2012;13(4):399–405. doi:10.2217/pg.11.165
Su SC, Chen C-B, Chang W-C, et al. HLA Alleles and CYP2C9*3 as predictors of phenytoin hypersensitivity in East Asians. Clin Pharmacol Ther. 2019;105(2):476–485. doi:10.1002/cpt.1190

Jang HW, Kim SW, Cho YJ, et al. GWAS identifies two susceptibility loci for lamotrigine-induced skin rash in patients with epilepsy. Epilepsy Res. 2015;115:88–94. doi:10.1016/j.eplepsyres.2015.05.014

Ozeki T, Mushiroda T, Yowang A, et al. Genome-wide association study identifies HLA-A*3101 allele as a genetic risk factor for carbamazepine-induced cutaneous adverse drug reactions in Japanese population. Hum Mol Genet. 2011;20(5):1034–1041. doi:10.1093/hmg/ddr537

McCormack M, Alfrevic A, Bourgeois S, et al. HLA-A*3101 and carbamazepine-induced hypersensitivity reactions in Europeans. N Engl J Med. 2011;364(12):1134–1143. doi:10.1056/NEJMoa1103297

Shen Y, Nicolleti P, Floratos A, et al. Genome-wide association study of severe blistersing skin rash caused by drugs. Pharmgenco J. 2012;12(2):96–104. doi:10.1038/tpj.2010.84

Tohkin M, et al. A whole-genome association study of major determinants for allopurinol-related Stevens-Johnson syndrome and toxic epidermal necrolysis in Japanese patients. Pharmgenco J. 2013;13(1):60–69. doi:10.1038/tpj.2011.41

Fernandez CA, Smith C, Yang W, et al. Genome-wide analysis links NFATC2 with asparaginase hypersensitivity. Blood. 2015;126(1):69–75. doi:10.1182/blood-2015-02-628800

Nishino J, Ochi H, Kochi Y, et al. Sample size for successful genome-wide association study of major depressive disorder. Front Genet. 2018;9:227. doi:10.3389/fgene.2018.00227

Lee SH, Clark S, van der Werf JH. Estimation of genomic prediction accuracy from reference populations with varying degrees of relationship. PLoS One. 2017;12(12):e0189775. doi:10.1371/journal.pone.0189775

International HapMap C. A haplotype map of the human genome. Nature. 2005;437(7063):1299–1320.

Fadista J, Manning AK, Florez JC, et al. The (in)famous GWAS P-value threshold revisited and updated for low-frequency variants. Eur J Hum Genet. 2016;24(8):1202–1205. doi:10.1038/ejhg.2015.269

Klaewsongkram J, Sukasem C, Thanitworasit P, et al. Analysis of HLA-B allelic variation and IFN-gamma ELISPOT responses in patients with severe cutaneous adverse reactions associated with drugs. J Allergy Clin Immunol Pract. 2019;7(1):219–227.e4. doi:10.1016/j.jaip.2018.05.004

Watanabe H, Watanabe Y, Tashiyo Y, et al. A docking model of dapsone bound to HLA-B*13:01 explains the risk of dapsone hypersensitivity syndrome. J Dermatol Sci. 2017;88(3):320–329. doi:10.1016/j.jdermsci.2017.08.007

Schäfer D, Brumen M, Dobovítek A, et al. Dynamic Systems of eicosanoid production with special reference to non-steroidal anti-inflammatory drug-triggered hypersensitivity. IET Systems Biology. 2015;9(5):204–215. doi:10.1049/iet-syb.2014.0037

Ammar R, Paton TA, Torti D, et al. Long read nanopore sequencing for detection of HLA and CYP2D6 variants and haplotypes [version 2; peer review: 2 approved]. F1000Research. 2015;4(17):17. doi:10.12688/f1000research.6037.2

Laver TW, et al. Pitfalls of haplotype phasing from ampiclon-based long-read sequencing. Sci Rep. 2016;6:21746. doi:10.1038/srep21746

Ambardar S, Gowda M. High-resolution full-length HLA typing method using third generation (Pac-Bio SMRT) sequencing technology. Methods Mol Biol. 2018;1802:135–153.

Suzuki S, Ranade S, Osaki K, et al. Reference Grade Characterization of Polymorphisms in Full-Length HLA Class I and II genes with short-read sequencing on the Ion PGM system and long-reads generated by single molecule, real-time sequencing on the pacbio platform. Front Immunol. 2018;9:2294. doi:10.3389/fimmu.2018.02294

Buermans HJP, Vossen RHAM, Anvar SY, et al. Flexible and scalable full-length CYP2D6 long amplicon PacBio sequencing. Hum Mutat. 2017;38(3):310–316. doi:10.1002/humu.23166

Hripsak G, Ryan PB, Duke JD, et al. Characterizing treatment pathways at scale using the OHDSI network. Proc Natl Acad Sci. 2016;113(27):7329–7336. doi:10.1073/pnas.1510502113

Dey S, Luo H, Fokoue A, et al. Predicting adverse drug reactions through interpretable deep learning framework. BMC Bioinform. 2018;19(821):476. doi:10.1186/s12859-018-2544-0

Poudel DR, Acharya P, Ghimire S, et al. Burden of hospitalizations related to adverse drug events in the USA: a retrospective analysis from large inpatient database. Pharmacoeconom Drug Saf. 2017;26(6):635–641. doi:10.1002/pds.4184

Barbarino JM, Whirl-Carrillo M, Altman RB, et al. PharmGKB: a worldwide resource for pharmacogenomic information. WIREs Syst Biol Med. 2018;10(4):e1417. doi:10.1002/wsbm.1417

Davis AP, Grondin CJ, Johnson RJ, et al. The comparative toxigenomics database: update 2019. Nucleic Acids Res. 2019;47(D1):D948–D954. doi:10.1093/nar/gky868

Kanehisa M, Goto S. KEGG: kyoto encyclopedia of genes and genomes. Nucleic Acids Res. 2000;28(1):27–30. doi:10.1093/nar/28.1.27

Hecker N, et al. SuperTarget goes quantitative: update on drug-target interactions. Nucleic Acids Res. 2012;40(Database issue):D1113–D1117.

Ghataoraia GS, et al. A web resource for mining HLA associations with adverse drug reactions: HLA-ADR. Database. 2016:2016.

Cai M-C, Xu Q, Pan Y-J, et al. ADReCS: an ontology database for aiding standardization and hierarchical classification of adverse drug reaction terms. Nucleic Acids Res. 2015;43(Database issue):D907–D913. doi:10.1093/nar/gku1066

Zh L, Han LY, Yap CW, et al. Drug ADVERSE Reaction Target Database (DART): proteins related to adverse drug reactions. Drug Safety. 2003;26(10):685–690. doi:10.2165/0002283-20032610-00002

Van der Auwera GA, et al. From FastQ to data at high confidence variant calls: the Genome Analysis Toolkit best practices pipeline. Curr Protoc Bioinformatics. 2013;43(11.01):10.11–10.33.

Li H, Aligning sequence reads, clone sequences and assembly contigs with BWA-MEM. arXiv preprint arXiv:1303.3997. 2013.

Poplin R, Chang P-C, Alexander D, et al. A universal SNP and small-indel variant caller using deep neural networks. Nat Biotechnol. 2018;36(10):983–987. doi:10.1038/nbt.4235

Twist GP, Gaedigk A, Miller NA, et al. Erratum: constellation: a tool for rapid, automated phenotype assignment of highly polymorphic pharmacogene, CYP2D6, from whole-genome sequences. Npj Genomic Med. 2017;2(1):16039. doi:10.1038/npjgenmed.2016.39

Chen X, et al., Accurate CYP2D6 genotyping using whole genome sequencing data. bioRxiv, 2020: p. 2020.

Numanagic I, Malikic S, Ford M, et al. Allelic decomposition and exact genotyping of highly polymorphic and structurally variant genes. Nat Commun. 2018;9(1):828. doi:10.1038/s41467-018-03273-1

Lee S-B, Wheeler MM, Thummel KE, et al. Calling star alleles with stargazer in 28 pharmacogenes with whole genome sequences. Clin Pharmacol Ther. 2019;106(6):1328–1337. doi:10.1002/cpt.1552

Boegel S, Löwer M, Schäfer M, et al. HLA typing from RNA-Seq sequence reads. Genome Med. 2012;4(12):102. doi:10.1186/gm403

Szołek A, Schubert B, Mohr C, et al. OptiType: precise HLA typing from next-generation sequencing data. Bioinformatics (Oxford, England). 2014;30(23):3310–3316. doi:10.1093/bioinformatics/btu548
121. Kawaguchi S, Higasa K, Shimizu M, et al. HLA-HD: an accurate HLA typing algorithm for next-generation sequencing data. Hum Mutat. 2017;38(7):788–797. doi:10.1002/humu.23230

122. Lee H, Kingsford C. Kourami: graph-guided assembly for novel human leukocyte antigen allele discovery. Genome Biol. 2018;19(1):16. doi:10.1186/s13059-018-1388-2

123. Boegel S, et al. In silico HLA typing using standard RNA-Seq sequence reads. In: Bugert P, editor. Molecular Typing of Blood Cell Antigens. New York: Springer New York; 2015:247–258.

124. McCarthy DJ, Humburg P, Kanapin A, et al. Choice of transcripts and software has a large effect on variant annotation. Genome Med. 2014;6(3):26. doi:10.1186/gm543

125. Jarvik GP, Browning BL. Consideration of cosegregation in the pathogenicity classification of genomic variants. Am J Human Genetics. 2016;98(6):1077–1081. doi:10.1016/j.ajhg.2016.04.003

126. Richards S, Aziz N, Bale S, et al. Standards and guidelines for the interpretation of sequence variants: a joint consensus recommendation of the American College of Medical Genetics and Genomics and the Association for Molecular Pathology. Genetics Med. 2015;17(5):405–424. doi:10.1038/gim.2015.30

127. Gaedigk A, Sangkuhl K, Whirl-Carrillo M, et al. Prediction of CYP2D6 phenotype from genotype across world populations. Genetics Med. 2017;19(1):69–76. doi:10.1038/gim.2016.80

128. Kim E-Y, Shin S-G, Shin J-G. Prediction and visualization of CYP2D6 genotype-based phenotype using clustering algorithms. Transl Clin Pharmacol. 2017;25(3):147–152. doi:10.12793/ tcep.2017.25.3.147

129. McInnes G, et al. Transfer Learning Enables Prediction of CYP2D6 Haplotype Function. bioRxiv; 2020:684357.

130. van der Lee M, Allard WG, Boilén S, et al. Repurposing of diagnostic whole exome sequencing data of 1583 individuals for clinical pharmacogenetics. Clin Pharmacol Ther. 2020;107(3):617–627. doi:10.1002/cpt.1665

131. Srinivasan BS, Chen J, Cheng C, et al. Methods for analysis in pharmacogenomics: lessons from the Pharmacogenetics Research Network Analysis Group. Pharmacogenomics. 2009;10(2):243–251. doi:10.2217/14622416.10.2.243

### Pharmacogenomics and Personalized Medicine

**Publish your work in this journal**

Pharmacogenomics and Personalized Medicine is an international, peer-reviewed, open access journal characterizing the influence of genotype on pharmacology leading to the development of personalized treatment programs and individualized drug selection for improved safety, efficacy and sustainability. This journal is indexed on the American Chemical Society’s Chemical Abstracts Service (CAS). The manuscript management system is completely online and includes a very quick and fair peer-review system, which is all easy to use. Visit http://www.dovepress.com/testimonials.php to read real quotes from published authors.

Submit your manuscript here: https://www.dovepress.com/pharmacogenomics-and-personalized-medicine-journal