NSIDES: DRUG EFFECT DISCOVERY USING THE FDA ADVERSE REPORTING SYSTEM

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Adverse drug events are a leading cause of morbidity and mortality around the world. Regulatory agencies, such as the Food and Drug Administration (FDA), maintain large collections of adverse event reports. These spontaneous reporting systems provide an opportunity to retrospectively study drug effects from patient population data. Using an updated version of our previously developed algorithm to conduct high dimensional propensity score matching, we use the FDA Adverse Event Reporting System (FAERS) to develop a a database drug effects, known as nSides. FAERS contains millions of reports, thousands of drugs and thousands of effects. In order to calculate side effect significances over the complete set of drugs reported in FAERS, we present a computational architecture which is distributed. In addition, we present an on-demand interface for the members of the public to request side effect significances for drug interactions.

1. Introduction

Spontaneous reporting systems such as the FDA Adverse Event Reporting System (FAERS) are important resources for detecting drug adverse events after a drug is approved (pharmacovigilance). However, pharmacovigilance algorithms often lead to many false positive and false negative findings due to issues of confounding, and detection of drug-drug interactions is an even greater challenge. We previously developed databases for off-label drug effects (OFFSIDES) and drug interactions (TWOSIDES) that account for these limitations using a novel Statistical Correction for Uncharacterized Bias (SCRUB).[1] These databases are available for download to the public. In addition to updates to the algorithm, we have developed a gateway to access the databases, known as nSides. nSides aims to make these databases accessible to researchers, clinicians, and patients alike and contains additional features related to drug safety. Since it is not feasible to generate models for every possible drug combination, we develop a novel model request system which submits jobs to the Open Science Grid and appends the results to the databases for future access.

2. Data Sources

There are several data sources which are involved in nSides. We use a curated version of the FDA Adverse Event Reporting System (FAERS) known as Adverse Event Open Learning through Universal Standardization (AEOLUS).[2] AEOLUS aims to clean and normalize the data by removing duplicate cases. This is done by applying standardized vocabularies in the form of RxNorm to map drug names and SNOMED-CT to map outcomes. The AEOLUS dataset is publicly available.

   The algorithm used to develop the databases used for nSides are similar to the previously developed OFFSIDES and TWOSIDES databases using raw FAERS data.[1] A standard signal detection algorithm involves conducting a disproportionality analysis by comparing the observed reporting frequency of a drug and outcome to the expected reporting frequency of all other drugs and the outcome. The metric is known as a Proportional Reporting Ratio (PRR). If the outcome occurred by chance, the frequencies will be equal and the PRR will be one. If the PRR is much larger than one, the null hypothesis is rejected. To reduce sampling variance and selection bias, propensity score matching is implemented on the FAERS data to form the groups used in the disproportionality analysis. This procedure, known as SCRUB, matches cases and controls between patients exposed and not exposed to a particular drug (OFFSIDES) or two drugs (TWOSIDES) to mitigate confounding biases.

3. Methods

3.1. *Computational Challenge*

The AEOLUS dataset contains ≈4,500 drugs, ≈4,500,000 reports and ≈7,500 effects. Since SCRUB is performed to find appropriate controls for a set of cases on a particular drug, a model needs to be developed for every unique drug. It is easily seen that modeling drug interactions becomes intractable quickly, with 20 million possible interactions of 2 drugs. However, many of the 2 drug combinations do not have enough reports in the AEOLUS dataset to perform

meaningful case-control matching.

References

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2. J. M. Banda, L. Evans, R. S. Vanguri, N. P. Tatonetti, P. B. Ryan and N. H. Shah, *Scientific data* **3** (2016).