A Signal Detection Analysis of World Health Organization’s Pharmacovigilance Database

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Abstract
Pharmacovigilance is the primary method used to identify hazards associated with medicinal products. Using generalized likelihood ratio tests, we examined the adverse reaction signals of the drug class Monoamine oxidase inhibitors (MAOIs) from World Health Organization’s Pharmacovigilance Database. The proposed test procedure has the ability of detecting adverse reactions of multiple drugs simultaneously. Our findings suggest there are 23 common Adverse reaction signals detected within this drug class. And postural hypotension, high blood pressure, fainting, abnormal heart rhythm, dizziness, headache, drowsiness are the most strong signals for the MAOIs class. An extensive simulation study performed to evaluate the proposed test procedure also suggests the proposed test procedure works well in practice.

Keywords
Signal detection, Adverse reaction, Monte Carlo simulation, Pharmacovigilance

Introduction
Reporting of drug or medical device related adverse reactions (ARs) is usually voluntary. One of the major postmarket safety surveillance databases is the World Health Organization’s (the “WHO”) global pharmacovigilance database, which contains reports of suspected ADRs, so called Individual Case Safety Reports (ICSRs), collected by national drug authorities in over 110 countries and span over more than 100,000 different medicinal products. Clinical reviewers evaluate adverse reactions reports to look for new safety concerns that might be related to a marketed product, or for a manufacturer’s compliance to reporting regulations.

There are several statistical methods available for adverse reactions detection in and other large postmarket databases. These methods include the reporting odds ratio (ROR, [1]), proportional reporting ratio (PRR, [2]), multi-gamma Poisson shrinker (MGPS, [3,4]), Bayesian confidence propagation neural network (BCPNN, [5]), Bayesian method based on a new Information Component (IC, [6]), simplified Bayes (sB, [7]), among others. A likelihood ratio test method ([8]), that assumes the number of adverse reactions follows a Poisson distribution, was developed to identify adverse reactions for a specific drug or to identify drugs for a particular adverse event. In post-marketing surveillance, the signals of adverse reactions within a drug class, or drug signals for a group of adverse reactions may be of interest to medical reviewers. In this article, we develop a generalized likelihood ratio test to identify adverse reactions that have high reporting rates compared to other adverse reactions associated with all the drugs of the same class or with similar treatment indications. The drug class refers to a set of drugs which have similar chemical structure such as the antibiotics drug class containing Penicillins, Tetracyclines, Cephalosporins, Quinolones, Lincomycins, Macrolides, Sulfonamides, Glycopeptides, Aminoglycosides and Carbapenems, etc. A group of adverse reactions refers to a set of preferred terms such as hepatic failure, alanine aminotransferase abnormal, ascites, blood bilirubin abnormal, cholestatic liver injury, hepatic atrophy, hepatomegaly, Reye’s syndrome, and so on that are all related to hepatocellular injury.

This article is organized as followed. In section 2, a
brief overview of World Health Organization’s global pharmacovigilance database is provided. In Section 3, we give a brief review of the likelihood ratio test procedure for adverse reactions detection for a single drug, then we propose a generalized likelihood ratio test procedure, namely GLRT, to detect multiple ARs in a drug class. The performance of GLRT is evaluated using simulated datasets in Section 4. In Section 5, both the LRT and the GLRT are applied to the 2000-2005 and 2005-2010 data from WHO’s global database. Section 6 contains some discussion and concluding remarks.

World Health Organization’s Pharmacovigilance Database

The WHO’s global pharmacovigilance database consists of the individual reports with demographical information, route of administration, drug/biological information, medical history, treatment indication, therapy start dates, and end dates. For adverse reaction detection, Medical Dictionary for terminology of preferred biologic information. In studying the drug-AE association, the generic name of the drug is used, which refers to the unique chemical makeup of a drug.

The WHO’s global pharmacovigilance database includes reports since 1980, however researchers and reviewers are more interested in data from recent years. In this article we focus on cases reported to WHO between 2000 and 2010 for more than 6500 drugs and 14,000 Adverse reactions. For any particular adverse event, the investigators consider all suspect and concomitant drugs.

Inference Procedure

Test procedure for adverse reactions detection of a single drug

After summarizing the data files, the WHO pharmacovigilance data can be presented in a tabular form with, say, adverse reactions (ARs) as the row variable and drugs as the column variable (as in Table 1), with \( n_{ij} \) as the cell count for \( j^{\text{th}} \) AR and \( j^{\text{th}} \) drug, \( n_i \) as the sum of counts for \( j^{\text{th}} \) AR \((j^{\text{th}} \text{row total})\) and \( n_j \) as the sum of counts for \( j^{\text{th}} \) drug \((j^{\text{th}} \text{column total})\).

We collapse the data structure table into multiple 3 × 3 tables. For a fixed \( j^{\text{th}} \) drug, we have \( l \) such tables (Table 2), each associated with an AR \((i = 1, ..., l)\). We assume that \( n_{ij} \sim \text{Poisson}(n_i \times p_{ij})\), where \( p_{ij} \) is the reporting rate of \( j^{\text{th}} \) drug for \( i^{\text{th}} \) AR; and \( n_j - n_{ij} \sim \text{Poisson}(n_j - n_{ij})q_j\) where \( q_j \) is the reporting rate of \( j^{\text{th}} \) drug for other ARs combined excluding \( j^{\text{th}} \) AR. We also assume \( n_i \) and \( n_j - n_{ij} \) are independent. Since drug \( j \) is fixed, unless stated otherwise, we suppressed the notational dependence of \( p_{ij} \) and \( q_j \) and on \( j^{\text{th}} \) drug. We define the null hypothesis,

\[ H_{0j}: p_{ij} = q_j = p_0 \text{ for all ARs in drug } j \text{ (ie. } H_{0j}: \bigcap_{i} p_{ij} = q_j = p_0); \]

and the alternative hypothesis (one sided) as

\[ H_{ij}: p_{ij} > q_j \text{ for at least one AR in drug } j \text{ (ie. } H_{ij}: \bigcup_{i} p_{ij} > q_j). \]

Under the null hypothesis, the maximum likelihood estimates (MLEs) for \( p_{ij} \) is \( \hat{p}_{ij} = \frac{n_{ij}}{n_i} \), and the expected number of cases for \( i^{\text{th}} \) AR and \( j^{\text{th}} \) drug is \( \hat{E}_{ij} = n_i \times \frac{n_{ij}}{n_i} \).

Under the two-sided alternative hypothesis \((p_{ij} = q_j)\), the maximum likelihood estimates (MLEs) for \( p_{ij} \) and \( q_j \) are

\[ \hat{p}_{ij} = \frac{n_{ij}}{n_i} \text{ and } \hat{q}_j = \frac{n_{ij} - n_{ij}}{n_i - n_i}. \]

The maximum likelihoods under both the null and the two-sided alternative hypotheses are obtained by replacing the parameters with their MLEs in the likelihood functions, leading to the likelihood ratio, for \( j^{\text{th}} \) AR and \( j^{\text{th}} \) drug as,

\[ LR_j = \frac{L_n(\hat{p}_{ij}, \hat{q}_j)}{L_n(\hat{p}_0)} = \frac{(\frac{n_{ij}}{n_i})^{n_{ij}}(\frac{n_i - n_{ij}}{n_i})^{n_i - n_{ij}}}{(\frac{n_i}{n_i})^{n_i}} = \frac{(\frac{n_{ij}}{n_i})^{n_{ij}}(\frac{n_i - n_{ij}}{n_i})^{n_i - n_{ij}}}{(\frac{n_i}{n_i})^{n_i}} \]

Where \( \hat{E}_{ij} = n_i \times \frac{n_{ij}}{n_i} \) is the expected number of counts for \( j^{\text{th}} \) AR and \( j^{\text{th}} \) drug. The likelihood ratio test statistic for testing \( H_{ij}: p_{ij} = q_j \) for all ARs in drug \( j \) vs. \( H_{ij}: p_{ij} > q_j \) for at least one AR, is the maximum likelihood ratio \( MLR_j = \max |LR_j| \), where the maximum is over ARs \( i = 1, ..., l \). We calculate \( LR_j = (\frac{n_{ij}}{n_i})^{n_{ij}}(\frac{n_i - n_{ij}}{n_i})^{n_i - n_{ij}} I(\hat{p}_{ij} > \hat{q}_j) \)

and define \( MLR = \max (LR_j) \) as the test statistic. For computational convenience, we may sometimes work with the log-likelihood ratio \( \log(MLR) \) which is the

| Drug | Drug | ... | ... | Drug | n_i |
|------|------|-----|-----|------|-----|
| AR1  | n_{11} | n_{12} | ... | ... | n_{1j} | n_i |
| AR2  | n_{21} | n_{22} | ... | ... | n_{2j} | n_i |
| ...  | ...  | ...  | ... | ... | ...  | ... |
| AR1  | n_{11} | n_{12} | ... | ... | n_{1j} | n_i |
| n_j  | n_{11} | n_{21} | ... | ... | n_{nj} | n_j |

| j^{th} Drug | Other Drugs | Marginal Total |
|-------------|-------------|---------------|
| P^{th} AR   | n_j        | n_j - n_{ij}  |
| Other AR    | (n_j - n_{ij}) | n_j - n_{ij}  |
| Marginal Total | n_j - n_{ij} | n_{j} |

Table 2: 3 × 3 tables for \( j^{\text{th}} \) drug.

Table 1: Data structure.
monotone function of $LR_{ij}$.

The distribution of MLR under $H_0$ is not analytically tractable and is obtained using Monte Carlo simulation as defined below. First, the number of cases for each AR, for a given drug $j$, are simulated under $H_0$. Under $H_0$, since $n_{ij},...,n_{iL}$ given the margin totals $n_i,...,n_l$ are independent $\text{Poisson}(n_j,...,n_{l})$, $i=1,...,l$, the joint distribution of $(n_{ij},...,n_{il})$ conditioning on $n_j$ and $(n_{ij},...,n_{il})$ is $(n_{ij},...,n_{il})|n_j,n_i,...,n_l \sim \text{multinomial}(n_j,(\frac{n_{ij}}{n_j,...,n_{il}})).$

A total of 499 datasets under $H_0$ are simulated from the multinomial distribution, and 500 MLRs are calculated. The null hypothesis is rejected at the $\alpha = 0.05$ level if the value of MLR from the observed dataset is greater than the 95th percentile of the 500 MLR values (threshold, $T_\alpha$). The corresponding $p$-value is then $1-R/500$, where $R$ is the rank of the observed MLR among all the 500 MLR values. If the $p$-value of the observed MLR is less than $\alpha$ (say, 0.05), then the AR associated with this MLR is the strongest signal among all ARs for the $j^{th}$ drug under consideration. Having found the strongest signal, we can then move to the second largest $LR_{ij}$, and so on, and declare them as signals if their $LR_{ij}$ are greater than $T_\alpha$ or the corresponding $p$-values are less than $\alpha$.

The likelihood ratio test is shown, analytically and through extensive simulation study, to control type-I error and false discovery rate (FDR) while retaining good power and sensitivity ([7,8]). In the next section, we generalize the likelihood ratio test procedure to detect all AR signals in a drug class. The methods to detect drug signals for a set of prespecified ARs can be performed in a similar fashion.

**Test procedure for adverse reaction detection of multiple drugs**

In order to develop a test statistic that can identify adverse reactions of multiple drugs in a class, we assume that a drug class has $K$ different drugs (usually $K$ is a small number), and we assume that for $k^{th}$ drug the number of reports for $i^{th}$ AR and all other ARs (excluding $i^{th}$ AR) still remains a Poisson distribution:

$$n_k \sim \text{Poisson}(n,p_k)$$

$$n_k-n_{k} \sim \text{Poisson}((n_{k}-n_{k})q_{k})$$

Where $p_k$ is the reporting rate of $k^{th}$ drug for $i^{th}$ AR, and $q_{k}$ is the reporting rate of $k^{th}$ drug for the other ARs. The null and alternative hypotheses for detecting AR signal in drug $k$ are,

$$H_{0k}: p_k = q_k = p_{\bar{k}}$$

for all ARs in drug $k$ versus

$$H_{a}: p_k > q_k \quad \text{for at least one AR.}$$

The null and alternative hypotheses for detecting AR signals among this drug class with $k$ drugs are,

$$H_0: \bigcup_i p_k = q_k = p_{\bar{k}}$$

$$H_a: p_k > q_k \quad \text{for} \quad i=1,...,l \quad \text{and} \quad k=1,...,K.$$  

Where $i=1,...,l$ and $k=1,...,K$, versus

$$H_a: \bigcup_k p_{ik} > q_{ik}$$

Another way to write the null and alternative hypotheses are

$$H_0: \bigcap_k H_{0k} \quad \text{versus} \quad H_a: \bigcup_k H_{ak}.$$  

Under the null hypothesis $H_0$, the MLE for $p_{ik}$ is $\hat{p}_{ik} = \frac{n_{ik}}{n_i}$ and the expected number of counts for $i^{th}$ AR and $k^{th}$ drug is $E_{ik} = n_i \times \frac{n_k}{n}$. Under the two-sided alternative hypothesis $(\bigcup_k p_k \neq q_k)$, the MLEs for $p_{ik}$ and $q_{ik}$ are $\hat{p}_{ik} = \frac{n_{ik}}{n_i}$ and $\hat{q}_{ik} = \frac{n_k-n_{ik}}{n-n_i}$.

The maximum likelihood under both the null and the two-sided alternative hypotheses are obtained by replacing the parameters with their MLEs in the likelihood functions, leading to the likelihood ratio for $i^{th}$ AR in $k^{th}$ drug as:

$$LR_{ik} = \frac{L(\hat{p}_{ik},\hat{q}_{ik})}{L(\hat{p}_i,\hat{q}_i)} = \frac{n_{ik}^{n_i - n_k} (n_k-n_{ik})^{n_i-n_k}}{n_i^{n_i} (n-n_i)^{n-n_i}} = \frac{n_{ik}^{n_i - n_k}}{n_i^{n_i-n_k} \langle n_{ik} \rangle^{n_i-n_k}}$$  

(2)

The likelihood ratio test statistic for testing $H_{0k}: p_k = q_k = p_{\bar{k}}$, versus, $H_{ak} p_k \neq q_k$ for at least one AR, is the maximum likelihood ratio $MLR_k = \max \{LR_{ik} \}, \ i=1,...,l$. The test statistic for testing $H_0: \bigcap_k p_k = q_k = p_{\bar{k}}$ versus $H_a: \bigcup_k p_k > q_k$ is

$$MLR = \max_k (MLR_k) = \max_k (\max_i (LR_{ik}(\hat{p}_{ik} > \hat{q}_{ik})))$$

Where $i=1,...,l$ and $k=1,...,K$.

Because the distribution of MLR under $H_0$ is not analytically tractable, we still use a Monte Carlo simulation to obtain its distribution. For each drug $k$ in the drug class under $H_0$ we generate 499 datasets using

$$(n_{i1},...,n_{il})|n_i;n_i,...,n_l \sim \text{multinomial}(n_j,(\frac{n_{i1}}{n_j,...,n_{il}}))$$  

(3)

and compute 500 values of MLR including the one from the real data, for $k = 1,...,K$. This results into 500 $\times K$ MLR values. The null hypothesis is rejected at $\alpha = 0.05$ level if the value of MLR from the observed dataset is greater than the $(1 - \alpha)^{th}$ percentile of the 500 $\times K$ MLR values $T_\alpha$. After AR associated with the largest $LR_{ik}$ is identified as signal ($LR_{ik} > T_\alpha$), we move to the AR with the second largest value of $LR_{ik}$ determine if it is a signal and so on. This way, the generalized likelihood ratio test procedure controls Type-I error. It also controls the false discovery rate (FDR) with $FDR \leq \alpha$.

**Applications**

In the following, we present the results from applying the likelihood ratio test procedure discussed
in Section 3 to the “Monoamine oxidase inhibitors” (MAOIs). The MAOIs are used to treat several conditions. They include, but are not limited to: Depression, generalized anxiety disorder, agitation, obsessive compulsive disorders (OCD), manic-depressive disorders, childhood enuresis (bedwetting), major depressive disorder, diabetic peripheral neuropathic pain, neuropathic pain, social anxiety disorder, post-traumatic stress disorder (PTSD) etc. The drug class includes Nardil (phenelzine), Parnate (tranylcypromine), Marplan (isocarboxazid), Emsam (selegiline), etc. We select four MAOIs labeled as MAOI1, MAOI2, MAOI3, MAOI4 and MAOI5 (not in any specific order) from the four MAOIs drugs; while using the generalized likelihood ratio test procedure to each of the four drugs in the drug class, there are 61, 32, 68, 39 ARs detected for the four MAOIs drugs; while using the generalized likelihood ratio test procedure to each of the four drugs in the drug class, there are 66, 37, 74, 45 ARs detected for the four MAOIs drugs. We apply the likelihood ratio test (LRT) and generalized likelihood ratio test (GLRT) for detecting Adverse Reactions.

The results of MAOIs drug class using both the LRT and GLRT are listed in Table 3. By using the likelihood ratio test procedure to each of the four drugs in the drug class, there are 66, 37, 74, 45 ARs detected for the four MAOIs drugs; while using the generalized likelihood ratio test, there are 61, 32, 68, 39 ARs detected, respectively. Across the four drugs, the GLRT detects less ARs than the LRT. By cross-checking the ARs in the four MAOIs drugs, there are 23 common ARs detected within this drug class. The top ARs are listed in Table 4 and Table 5, and postural hypotension, high blood pressure, fainting, abnormal heart rhythm, dizziness, headache, drowsiness are the most strong ARs for this MAOIs class.

### A Simulation Study

#### Data simulation

We then study the performance of the generalized likelihood ratio test (GLRT) using simulated datasets. We simulate datasets based on the four drugs in the monoamine oxidase inhibitors drug class in WHO’s global pharmacovigilance database.

Under the null hypothesis, the data are simulated from multinomial distribution (3). Under the alternative hypothesis, data are generated as follow,

\[
(a_i, a_2, ..., a_k) | n; n_1, ..., n_k \sim \text{multinomial}(n, (r_{i0} + \gamma_{i0} \cdot n_i, ..., (r_{k0} + \gamma_{k0} \cdot n_k)) \ (4)
\]

Where \( k = 1, ..., 5 \), and \( r_{i0}, ..., r_{k0} \) are the relative reporting rates for \( AE_{i'}, ..., AE_{k'} \) in \( K \) drugs with constraints

\[
0 \leq r_{i0} \ast r_{j0} \ast \frac{n_i}{n_j} \quad \text{and} \quad \sum_{i=1}^{k} r_{i0} \ast r_{j0} \ast \frac{n_i}{n_j} = 1.
\]

Relative reporting rates \( r_{i0} \) are specified as follows: \( r_{i0} \) are assigned a value; higher than 1 for ARs selected as signals and 1 for all other ARs not selected as signals. \( r_{i0} \) can be regarded as baseline risks for drug \( k \), and \( r_{i0} \) can be different from one drug to another.

We evaluate how the relative reporting rate \( (rr) \), the sample size \( (n_i) \) and the number of signals affect the performance of the GLRT through the following four

### Table 3: Number of signals detected by the LRT and GLRT in Monoamine oxidase inhibitors (MAOIs) drug class.

| Drug   | 2000-2005 | 2005-2010 |
|--------|-----------|-----------|
|        | LRT | GLRT | LRT | GLRT |
| MAOI1  | 66  | 61   | 95  | 88   |
| MAOI2  | 37  | 32   | 52  | 45   |
| MAOI3  | 74  | 68   | 90  | 89   |
| MAOI4  | 45  | 39   | 73  | 68   |

### Table 4: Total common reactions found for the five drugs in MAOIs class in 2000-2005 data set.

| Drug   | Common Signals                                                                 |
|--------|-------------------------------------------------------------------------------|
| MAOI1  | Postural hypotension, fainting, abnormal heart rhythm, dizziness, headache, drowsiness, insomnia, constipation, nausea, diarrhea, sexual dysfunction, weight gain or weight loss, seizures, rash, blurred vision, hepatitis |
| MAOI2  | High blood pressure, fainting, abnormal heart rhythm, dizziness, headache, drowsiness, insomnia, anxiety, nausea, diarrhea, sexual dysfunction, weight gain or weight loss, blurred vision, hepatitis |
| MAOI3  | Postural hypotension, dizziness, headache, drowsiness, insomnia, anxiety, constipation, nausea, diarrhea, sexual dysfunction, weight gain or weight loss, seizures, rash, blurred vision, hepatitis |
| MAOI4  | Fainting, headache, drowsiness, postural hypotension, high blood pressure, insomnia, anxiety, constipation, nausea, diarrhea, sexual dysfunction, weight gain o weight loss, hepatitis |

### Table 5: Total common reactions found for the five drugs in MAOIs class in 2005-2010 data set.

| Drug   | Common Signals                                                                 |
|--------|-------------------------------------------------------------------------------|
| MAOI1  | Postural hypotension, dizziness, fainting, abnormal heart rhythm, headache, drowsiness, insomnia, constipation, nausea, diarrhea, sexual dysfunction, weight gain or weight loss, seizures, rash, blurred vision, hepatitis |
| MAOI2  | Abnormal heart rhythm, dizziness, high blood pressure, fainting, headache, drowsiness, insomnia, anxiety, nausea, diarrhea, sexual dysfunction, weight gain or weight loss, blurred vision, hepatitis |
| MAOI3  | Dizziness, headache, postural hypotension, drowsiness, insomnia, anxiety, diarrhea, sexual dysfunction, constipation, nausea, weight gain or weight loss, seizures, rash, blurred vision, hepatitis |
| MAOI4  | Fainting, drowsiness, postural hypotension, headache, high blood pressure, insomnia, nausea, anxiety, constipation, diarrhea, sexual dysfunction, weight gain o weight loss, hepatitis |
across the drug class, but signals are not necessarily common between drugs.

- Scenario 4: We take a similar process as Scenario 3, randomly select 30 signals for each drug independently, but we use inhomogeneous $rr$. A rate of $2 \times rr$ is assigned to those AR signals for which $n_i$ (the total num-

### Table 6: Illustration of false discovery rate (FDR)

| $H_0$ is true | $H_a$ |
|---------------|-------|
| Positive result | V | S |
| Negative result | U | T |

### Table 7: Simulation results of GLRT.

| Marginal Count | $n_i$ | $rr$ | Power | Sensitivity | FDR  |
|----------------|-------|------|-------|-------------|------|
| Scenario 1: Select one signal in one drug with marginal total as 28,216 or 4,362 |
| 28216          | 500   | 3    | 0.073 | 0.13        | 0.0565 |
| 28216          | 1500  | 3    | 0.315 | 0.278       | 0.0442 |
| 28216          | 3000  | 3    | 0.83  | 0.753       | 0.044  |
| 28216          | 5000  | 3    | 0.992 | 0.983       | 0.0368 |
| 28216          | 8000  | 3    | 1     | 1           | 0.0257 |
| 28216          | 12000 | 1    | 0.061 | NA          | 0.053  |
| 28216          | 12000 | 2    | 0.758 | 0.751       | 0.043  |
| 28216          | 12000 | 3    | 1     | 1           | 0.0351 |
| 28216          | 12000 | 5    | 1     | 1           | 0.0326 |
| 28216          | 12000 | 7    | 1     | 1           | 0.0316 |
| 4362           | 12000 | 1    | 0.077 | NA          | 0.077  |
| 4362           | 12000 | 2    | 0.084 | 0.005       | 0.090  |
| 4362           | 12000 | 3    | 0.152 | 0.079       | 0.073  |
| 4362           | 12000 | 4    | 0.50  | 0.364       | 0.0569 |
| 4362           | 12000 | 5    | 1     | 1           | 0.0393 |
| 4362           | 12000 | 7    | 1     | 1           | 0.0397 |
| Scenario 2: Randomly select 30 common signals with homogeneous $rr$ |
| NA             | AS    | 1    | 0.062 | NA          | 0.064  |
| NA             | AS    | 2    | 0.744 | 0.02107    | 0.0381 |
| NA             | AS    | 3    | 1     | 0.1347     | 0.0057 |
| NA             | AS    | 5    | 1     | 0.4794     | 0.0014 |
| NA             | AS    | 7    | 1     | 0.7315     | 0.0011 |
| NA             | AS    | 9    | 1     | 0.8520     | 0.0009 |
| Scenario 3: Randomly select 30 signals (not necessarily common) with homogeneous $rr$ |
| NA             | AS    | 1    | 0.085 | NA          | 0.081  |
| NA             | AS    | 2    | 0.80  | 0.0163     | 0.0293 |
| NA             | AS    | 3    | 1     | 0.1426     | 0.0050 |
| NA             | AS    | 5    | 1     | 0.4450     | 0.0018 |
| NA             | AS    | 7    | 1     | 0.6829     | 0.0013 |
| NA             | AS    | 9    | 1     | 0.8018     | 0.0010 |
| Scenario 4: Randomly select 30 signals with inhomogeneous $rr$ |
| NA             | AS    | 1    | 0.056 | NA          | 0      |
| NA             | AS    | 3    | 1     | 0.93912    | 0.0009 |
| NA             | AS    | 5    | 1     | 0.98467    | 0.006  |
| NA             | AS    | 7    | 1     | 0.99887    | 0.018  |

AS: Actual sample size; NA: Not applicable.
FDR is defined as $\frac{V}{V+S}$, the expected proportion of rejected null hypotheses which are erroneously rejected. It’s estimated by,

$$FDR = \frac{1}{L} \sum_{i=1}^{L} \frac{\text{Number of reactions falsely detected in } i^{th} \text{ simulated data (V)}}{\text{Total number of reactions detected in the } i^{th} \text{ simulated data (V+S)}}$$

All power, ST and FDR have values between 0 and 1. As we shall see in next section, GLRT have high sensitivity, low FDR, and to control Type-I error $\alpha$ which indicates its superiority over the conventional likelihood ratio test.

### Simulation results

The simulation results shown in Table 7 include power, sensitivity, and false discovery rate for the different scenarios described in Section 5.1.

In Scenario 1, one signal was assigned to the ARs with the relative large or moderate marginal counts (28,216 and 4,362). With fixed $r = 3$, $n_i = 28216$ and sample size $n_j = 500$, the power is 0.073, ST is 0.13 and FDR is 0.0565. As the sample size $n_j$ increases, the power and ST increase to 1, and FDR decreases from 0.06 to 0.03. When the sample size of AR is fixed at $n_i = 28216$, with the increase of $r$ from 1 to 7, the power increases from 0.06 to 0.75, and then to 1. The same increasing trend is also observed for ST. FDR decreases from 0.05 to 0.03, a value much lower than

### Table 8: The effect of the number of true signals on the performance of the GLRT.

| Marginal Count | $n_j$ | $r$ | Power | Sensitivity | FDR |
|----------------|------|----|-------|-------------|-----|
| **Scenario 2.1: Randomly select 10 common signals with homogeneous $r$** | | | | | |
| NA | AS | 1 | 0.071 | NA | 0.068 |
| NA | AS | 2 | 0.246 | 0.076 | 0.059 |
| NA | AS | 3 | 0.996 | 0.14081 | 0.021 |
| NA | AS | 5 | 1 | 0.45008 | 0.006 |
| NA | AS | 7 | 1 | 0.69561 | 0.0030 |
| NA | AS | 9 | 1 | 0.80329 | 0.00265 |
| **Scenario 2.2: Randomly select 20 common signals with homogeneous $r$** | | | | | |
| NA | AS | 1 | 0.059 | NA | 0.058 |
| NA | AS | 2 | 0.535 | 0.01639 | 0.005 |
| NA | AS | 3 | 1 | 0.1601 | 0.0082 |
| NA | AS | 5 | 1 | 0.5631 | 0.0029 |
| NA | AS | 7 | 1 | 0.72354 | 0.00145 |
| NA | AS | 9 | 1 | 0.85018 | 0.00137 |
| **Scenario 2.3: Randomly select 30 common signals with homogeneous $r$** | | | | | |
| NA | AS | 1 | 0.063 | NA | 0.063 |
| NA | AS | 2 | 0.837 | 0.01416 | 0.0449 |
| NA | AS | 3 | 1 | 0.1629 | 0.0052 |
| NA | AS | 5 | 1 | 0.5048 | 0.0012 |
| NA | AS | 7 | 1 | 0.7393 | 0.0009 |
| NA | AS | 9 | 1 | 0.9461 | 0.0008 |

AS: Actual sample size; NA: Not applicable.
vides a useful tool to identify potential adverse reactions in pharmacovigilance database. However, the final discovery of the true adverse reactions should also be based on a thorough review of all available medical records.

**Competing Interests**

The author declares that he has no competing interests.

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