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Bayesian adjustment for exposure misclassification in
case-control studies

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Abstract

Summary: Poor measurement of explanatory variables occurs frequently in observational studies. Error-prone observations may lead to biased estimation and loss of power in detecting the impact of explanatory variables on the response. We consider misclassified binary exposure in the context of case-control studies, assuming the availability of validation data to inform the magnitude of the misclassification. A Bayesian adjustment to correct for the misclassification is investigated. Simulation studies show that the Bayesian method can have advantages over non-Bayesian counterparts, particularly in the face of a rare exposure, small validation sample-sizes, and uncertainty about whether exposure misclassification is differential or non-differential. The method

is illustrated via application to several real studies.

Keywords: Bayesian methods; case-control study; exposure misclassification; simulation-extrapolation.

1 Introduction

In biomedical studies, the misclassification problem arises when a categorical exposure variable T is not precisely recorded. Instead of T , an approximate measurement or a *surrogate*, X is obtained. Replacing T with X in data analysis without accounting for the misclassification does not generally lead to valid inference about the association between T and a health-related response Y . Hence, the goal of adjustment for mismeasurement is to achieve valid inference about the (T, Y) relationship from (X, Y) data. In this paper, we restrict ourselves to misclassification problems on a binary exposure variable ($T = 0, 1$) in case-control studies ($Y = 0, 1$ for controls, cases) and no other covariates at play. We consider the setting whereby a “validation subsample” is available, i.e, for the majority of subjects only (X, Y) data are obtained, but for a (randomly-selected) minority (T, X, Y) are obtained. Such a design can arise when X is inexpensive and/or quick to measure whereas T is expensive and/or time-consuming to measure. Table 1 described the data structure. While each cell a_{ij} in the validation data is fully specified ($i = 0, 1, j = 1, 2, 3, 4$), only margins $a_{05}, a_{06}, a_{15}, a_{16}$ in the main data are recorded.

It is sometimes sensible to assume the conditional distribution of X given T and Y does not depend on Y , which is known as *nondifferential* misclassification. In other circumstances, the sampling scheme of case-control studies (explanatory variables are retrieved after the diagnosis) may well lead to the so called *differential* measurement error, i.e. the conditional distribution of the surrogate X given the unobservable exposure T also depends on the response Y . When information about covariates is collected through some “self-report” mechanism, subjects with target clinical outcomes may tend to erroneously “blame” a set

of risk factors for their conditions, or “ignore” previous exposure to avoid any connection between behaviour and disease.

There is a large literature on correcting for exposure mismeasurement, for example Barron [1], Marshall [2], Lyles [3], Carroll et al. [4]. Most work approaches the problem from a frequentist perspective, assuming complete knowledge of whether the misclassification is nondifferential or differential. A quality indices method without explicit assumption of non-differential misclassification was proposed to estimate bias to the observed odds ratio on whole sample, when validation subsample is available [5, 6]. The simulation extrapolation method and latent class logistic regression model were also developed to tackle the problem [7, 8]. On the other hand, the dramatic improvement of computational capability and the development of indirect simulation techniques such as Markov chain Monte Carlo (MCMC) make it possible to explore misclassification problems from a Bayesian perspective [9, 10, 11, 12]. In fact, partial prior knowledge of misclassification probabilities is often accessible to medical researchers, which makes Bayesian analysis an appealing approach.

Therefore in this paper, we primarily introduce a series of Bayesian methods suitable for different misclassification assumptions. Their performance will be closely compared to those of the maximum likelihood estimates (MLEs), quality indices (QI) method and simulation extrapolation (SIMEX) method, using simulation studies and real datasets. Section 2 presents detailed methodology for the proposed Bayesian methods. Section 3 discusses the comparative behaviours of the four methods based on simulation studies. Sections 4 and 5 present the performances of Bayesian and other methods via case-control studies with misclassified exposure variables and validation sub-samples. Section 6 provides some concluding remarks.

2 Bayesian adjustment for misclassification

Let us denote the true exposure prevalences amongst controls and cases by $r_i = P(T = 1|Y = i), i = 0, 1$. The retrospective odds ratio describing the correlation between the response and

explanatory variable is defined as

$$OR_T = \frac{r_1/(1-r_1)}{r_0/(1-r_0)}.$$

Sensitivity (SN) and specificity (SP) jointly measure the magnitude of exposure misclassification. In the scenarios subject to *differential* misclassification, the conditional distribution of the surrogate X given T can change with Y . The sensitivities and specificities among cases and controls can be formulated as, $SN_i = P(X = 1|T = 1, Y = i)$, $SP_i = P(X = 0|T = 0, Y = i)$, $i = 0, 1$. Prevalences of the *apparent* exposure for diseased and non-diseased individuals are denoted by $r_i^* = P(X = 1|Y = i) = r_i SN_i + (1 - r_i)(1 - SP_i)$, $i = 0, 1$. The degree of misclassification can also be expressed by the positive predictive value (PPV) and negative predictive value (NPV), where

$$PPV_i = P(T = 1|X = 1, Y = i) = \frac{SN_i r_i}{SN_i r_i + (1 - SP_i)(1 - r_i)} \quad (1)$$

$$NPV_i = P(T = 0|X = 0, Y = i) = \frac{SP_i(1 - r_i)}{SP_i(1 - r_i) + (1 - SN_i)r_i} \quad (2)$$

It is easy to justify that, in the main study the actual number of subjects of positive exposure status (b_{i1}) amongst those who are apparently exposed in either case or control group (a_{i5}) follows a Binomial distribution, i.e. $b_{i1} \sim Binomial(a_{i5}, PPV_i)$. Similarly, conditioning on the number of cases or controls with negative apparent exposure status (a_{i6}), the number of truly unexposed subjects (b_{i4}) follows $Binomial(a_{i6}, NPV_i)$, for $i = 0, 1$.

When the nondifferential misclassification condition is fulfilled, meaning the conditional distribution of $X|T, Y$ does not depend on Y , it follows immediately that $SN_0 = SN_1 = SN$, $SP_0 = SP_1 = SP$. However it is worth pointing out that, nondifferential misclassification does not imply equality of cases and controls regarding the predictive values (PPV_i , NPV_i).

2.1 Prior distributions

The exposure prevalances r_0 , r_1 , sensitivities SN_0 , SN_1 , and specificities SP_0 , SP_1 are the parameters of interest. By converting into a logit scale, $logit(x) = \log\{x/(1-x)\}$, the prior

information concerning these parameters can be modeled using bivariate normal distributions [13]. The actual exposure prevalences (r_i), sensitivities (SN_i) and specificities (SP_i) of X as a surrogate for T are assumed to be uncorrelated of one another, with,

$$\begin{pmatrix} \text{logit}(r_0) \\ \text{logit}(r_1) \end{pmatrix} \sim N \left(\begin{pmatrix} \mu_1 \\ \mu_2 \end{pmatrix}, \begin{pmatrix} \sigma_1^2 & \rho_1 \sigma_1 \sigma_2 \\ \rho_1 \sigma_1 \sigma_2 & \sigma_2^2 \end{pmatrix} \right),$$

$$\begin{pmatrix} \text{logit}(SN_0) \\ \text{logit}(SN_1) \end{pmatrix} \sim N \left(\begin{pmatrix} \nu_1 \\ \nu_2 \end{pmatrix}, \begin{pmatrix} \tau_1^2 & \rho_2 \tau_1 \tau_2 \\ \rho_2 \tau_1 \tau_2 & \tau_2^2 \end{pmatrix} \right),$$

$$\begin{pmatrix} \text{logit}(SP_0) \\ \text{logit}(SP_1) \end{pmatrix} \sim N \left(\begin{pmatrix} \gamma_1 \\ \gamma_2 \end{pmatrix}, \begin{pmatrix} \delta_1^2 & \rho_3 \delta_1 \delta_2 \\ \rho_3 \delta_1 \delta_2 & \delta_2^2 \end{pmatrix} \right).$$

It follows immediately that,

$$\text{logit}(SN_0) - \text{logit}(SN_1) \sim N(\nu_1 - \nu_2, \tau_1^2 + \tau_2^2 - 2\rho_2 \tau_1 \tau_2) \quad (3)$$

$$\text{logit}(SP_0) - \text{logit}(SP_1) \sim N(\gamma_1 - \gamma_2, \delta_1^2 + \delta_2^2 - 2\rho_3 \delta_1 \delta_2) \quad (4)$$

Our prior beliefs can be reflected through the *hyperparameters*, μ_i , σ_i , ν_i , τ_i , γ_i , δ_i and ρ_j . For instance, we proceed to set the prior distributions on the misclassification parameters as follows. We set $\nu_1 = \nu_2$, $\gamma_1 = \gamma_2$, $\tau_1^2 = \tau_2^2$, $\delta_1^2 = \delta_2^2$ to reflect an absence of knowledge about the “direction” of possible differentiability in the exposure assessment, with the assigned values to these quantities then reflecting prior belief about the extent of exposure misclassification. Put another way, we are expressing *exchangeable* prior beliefs about the misclassification of controls versus the misclassification of cases.

As a result, setting $\rho_2 = \rho_3 = 1$ implies that $SN_0 = SN_1$ and $SP_0 = SP_1$, which corresponds to nondifferential misclassification. Conversely, setting $\rho_2 = \rho_3 = 0$ implies independence of SN_0 and SN_1 , and independence of SP_0 and SP_1 . This intuitively reflects the notion that sensitivities or specificities are free to vary by themselves, and can be interpreted as “fully differential” misclassification. We will describe situations in between ($0 < \rho_j < 1$, $j = 2, 3$) as corresponding to “nearly nondifferential” misclassification, particularly when

each ρ_j is close to one. This setting is useful when investigators postulate that the nondifferential assumption might hold, and that should it be violated, the extent of violation is not likely to be severe.

Similarly, we set $\mu_1 = \mu_2$ and $\sigma_1 = \sigma_2$ to be “unbiased”, a *priori* concerning the direction of any exposure-disease association. The particular choice of values is dictated by belief about plausible values for exposure prevalence. We can then choose ρ_1 to obtain plausible prior for the effect size.

2.2 Posterior simulation

As is common in problems with “latent structure”, we can implement Bayesian inference via simulation from the distribution of parameters and unobservables given observables. In the fully-differential and nearly-nondifferential cases, this amounts to sampling from the distribution of parameter $\theta = (r_0, r_1, SN_0, SN_1, SP_0, SP_1)$ and latent variables b_{ij} given observed data a_{ij} . It is easy to verify that in the related problem where the prior on θ is comprised of independent uniform distributions (or more generally independent beta distributions) for each parameter, that Gibbs sampling is possible. That is, in the related problem each component of θ has a standard “full conditional” distribution. Gibbs sampling has the nice features that (i) no tuning constants are involved, and (ii) proposed moves are always accepted. Therefore we adapt this approach to the actual problem at hand by implementing a Metropolis-Hastings algorithm, *using the full-conditionals for the related problem to generate proposals*. Thus tuning is still not needed. Moreover, the acceptance probability for each proposal will depend only on the ratio of prior densities, i.e., the specified prior based on bivariate normal distributions versus the uniform prior in the related problem. Thus we find high acceptance rates, and in general this algorithm performs well. Note also that the same computational strategy can be adopted in the nondifferential case, via the smaller parameter vector $\theta = (r_0, r_1, SN, SP)$. The Bayesian method is implemented in R and downloadable from <http://www.stat.ubc.ca/People/Home/index.php?person=gustaf>.

3 Simulation Studies

3.1 Data Simulation

In order to demonstrate the comparative performance of Bayesian adjustment against other statistical approaches, we conduct a simulation study for three choices of odds ratio (1.25, 1.8 and 2.5) and two choices of exposure prevalence in the control group ($r_0 = 0.25$ and 0.04). At each combination, four misclassification scenarios concerning different levels of differentiability are built. Data in scenario 1 are simulated under nondifferential misclassification, with increasing degree of differentiability in scenarios 2, 3, and 4. To mimic the occurrence of erroneously “blaming” or “ignoring” a risk factor, we let the misclassification arise across scenarios, as follows.

- *Scenario 1:* $(SN_0, SN_1)=(0.80, 0.80)$, $(SP_0, SP_1)=(0.90, 0.90)$
- *Scenario 2:* $(SN_0, SN_1)=(0.80, 0.75)$, $(SP_0, SP_1)=(0.90, 0.85)$
- *Scenario 3:* $(SN_0, SN_1)=(0.80, 0.70)$, $(SP_0, SP_1)=(0.90, 0.80)$
- *Scenario 4:* $(SN_0, SN_1)=(0.80, 0.65)$, $(SP_0, SP_1)=(0.90, 0.75)$

For each scenario, 2000 datasets are generated to assure that simulation standard error for a true 95% interval is 0.005. Three sample sizes (500, 960, and 1760) are considered in data generation: (a) $\sum_{j=1}^4 a_{ij} = 50$, $a_{i5} + a_{i6} = 200$; (b) $\sum_{j=1}^4 a_{ij} = 80$, $a_{i5} + a_{i6} = 400$; and (c) $\sum_{j=1}^4 a_{ij} = 80$, $a_{i5} + a_{i6} = 800$.

Three Bayesian methods, adopting nondifferential, nearly nondifferential and differential prior distributions respectively, are applied to each dataset, to adjust for possible misclassifications and assess the association between the true exposure and outcome. In this study, data generation and analysis of simulated and real examples are all implemented in **R**.

3.2 Choice of Hyperparameters

According to Section 2, under the assumptions that $\mu_1 = \mu_2 = \mu$, $\sigma_1 = \sigma_2 = \sigma$, $\nu_1 = \nu_2$, $\gamma_1 = \gamma_2 = \gamma$, $\tau_1 = \tau_2 = \tau$ and $\delta_1 = \delta_2 = \delta$, we assign $\mu = -1.946$, $\sigma = 0.993$ to model the prior information that the logit true exposures are normally distributed with central 95%

probability between $\text{logit}(0.02)$ and $\text{logit}(0.5)$. Mild correlation between r_0 and r_1 ($\rho_1 = 0.3$) is selected to allow a relatively large prior standard deviation of 1.175 for $\log OR$ around mean 0. Similarly, we set $\nu = \gamma = 1.675$, $\tau = \delta = 0.648$ to represent the prior knowledge that the logit sensitivity and logit specificity are normally distributed within $\text{logit}(0.6)$ and $\text{logit}(0.95)$ with 95% probability. As discussed in Section 2, we set $\rho_2 = \rho_3 = 1$ to reflect nondifferential misclassification; $\rho_2 = \rho_3 = 0$ to express prior belief in differential misclassification.

The choice of ρ_2, ρ_3 for nearly nondifferential misclassification requires extra work. We note that by setting $\rho_2 = \rho_3 = 0.95$ we attain:

$$P\{|\text{logit}(SN_1) - \text{logit}(SN_0)| < 0.1\} = P\{|\text{logit}(SP_1) - \text{logit}(SP_0)| < 0.1\} = 0.3746,$$

and (by simulation)

$$P\{|SN_1 - SN_0| < 0.01\} = P\{|SP_1 - SP_0| < 0.01\} = 0.32252.$$

This seems reasonable as an encapsulation of the notion that deviations from nondifferentiability are not likely severe.

3.3 Model comparison

Bayesian statistical inferences are conducted based on samples drawn from 10000 MCMC iterations, after we discard the first 1000 simulations to diminish the effect of initial distributions.

The performance of Bayesian methods is contrasted with maximum likelihood (ML), quality indices and SIMEX methods. MLEs for model parameters under differential misclassification are calculated using closed-form expressions given by Lyles Lyles [3]. A numerical optimizer (function “`optim()`” in **R**) is adopted to maximize the log likelihood under nondifferential misclassification. The asymptotic variance of the log odds-ratio estimator is attainable by the multivariate Delta method in these cases.

The quality indices (QI) method was developed to assess misclassification in case-control

studies [5]. It provides a simple formula for calculating the actual odds ratio OR_T , taking advantage of the relationship between the true exposure T and surrogate X observed in the validation subsample: $OR_T = OR_X \times \frac{OR'_T}{OR'_X}$, where OR'_T and OR'_X are odds ratios for T and X in the validation data. The asymptotic variance of $\log OR_T$ is calculated via Delta method.

The simulation extrapolation (SIMEX) method originates in continuous measurement error settings [14]. The method introduces artificial extra measurement error to the data in question, in order to infer a relationship between the magnitude of measurement error and the estimate of the exposure-disease relationship. This relationship is then extrapolated back to the point of zero measurement error, to give an estimate which is adjusted for this error. Recently, Küchenhoff et al. extended the SIMEX procedure to the case of misclassified categorical data [7]. In brief, extra misclassification is introduced by raising the misclassification matrix to a power $\lambda > 1$, for multiple values of λ . The relationship between the point estimate of interest and λ is then extrapolated back to the $\lambda = 0$ setting of no misclassification (i.e., misclassification matrix equal to the identity matrix). The corresponding software package [15] allows different choices of extrapolation function and different methods for the calculation of standard errors. We therefore reports multiple sets of results for SIMEX. Note also that the SIMEX procedure is operationalized by “plugging in” estimates of sensitivity and specificity obtained from the validation data. Thus by pooling or not-pooling the validation data across controls and cases, one can implement SIMEX under the nondifferential or differential misclassification assumptions respectively.

Results for all inferential schemes (3 sample sizes by 3 true OR s by 2 levels of exposure prevalence) are reported in terms of mean-squared error, bias and sample variance of point estimators, and coverage and average width of nominal 95% interval estimators. Results for two particular schemes are provided in Table 2 and 3.

Results under the higher setting of exposure prevalence are explained through an example in Table 2, where $r_0 = 0.25$, $OR=1.25$, $\sum_{j=1}^4 a_{ij} = 80$ and $a_{i5} + a_{i6} = 800$. Within MLE,

Bayes and SIMEX approaches, the nondifferential estimates of $\log OR$ have smaller MSE and sample variance, when the data truly are nondifferentially misclassified (scenario 1). Differential methods are not as efficient under truly nondifferential misclassification, because sensitivity and specificity are estimated separately, and therefore via less data, for controls and cases. On the other hand, coverage of nondifferential methods deteriorates rapidly as the true misclassification mechanism becomes more differential. The performance of Bayes and ML methods is somehow comparable, with the former having slightly smaller overall error rate and sample variance, yet often bigger bias when differential misclassification is correctly assumed.

The performance of QI method is adequate when level of misclassification between case and control group is substantial, for QI is advantageous in controlling bias to a minimal level. Nevertheless, simulation results suggest QI point estimator tends to have larger sample variance and MSE, followed by the ML-DF estimator then Bayes-DF estimator, when differential misclassification is correctly (or incorrectly) specified. When misclassification is nondifferential and the MLE-NDF or Bayes-NDF procedure is applied, the QI estimate has a 50 to 100 per cent larger sample variance compared to the alternatives, although the property of small bias remains in this case. In general, the QI method produces accurate but less precise estimator regardless of misclassification assumption between groups.

Note also that in comparing SIMEX to other methods in Table 2, even the empirically better choice of extrapolation function and variance estimation (quadratic form with asymptotic variance) in SIMEX gives much larger bias and sample variance, shorter confidence interval (CI) and lower coverage proportion than the corresponding Bayes or ML procedure, for the differential misclassification scenarios. For nondifferential misclassification, SIMEX has point estimates similar to Bayes and ML methods, yet shorter CI and smaller coverage. This is not necessarily surprising. While the SIMEX approach is intuitively appealing, it does not carry the large-sample efficiency guarantees that come with likelihood-based procedures.

Results for ML and Bayes procedures in the lower exposure prevalence setting are illus-

trated via Table 3 where $r_0 = 0.04$, $OR=1.25$, $\sum_{j=1}^4 a_{ij} = 80$ and $a_{i5} + a_{i6} = 800$. The combination of rare exposure, relatively high sensitivity, and relatively small validation sample size implies that for some generated datasets no subject is truly exposed to the risk factor in the case or control group, i.e. $a_{i1} = 0$ and $a_{i2} = 0$. For differential misclassification, this leads to $\widehat{PPV}_i = 0$, $\widehat{NPV}_i = 1$, hence $\hat{r}_i = 0$ and undefined $\log\widehat{OR}$ in the ML-DF model. ML-DF results in Table 3 are based on datasets without such empty cells in the validation data. Empty cells also result in nonsensical ML estimate of $\log OR$ using numerical optimizer in the nondifferential case. After examining results, we decide to remove such simulations from calculating ML-NDF results when $\log OR$ is inestimable in numeric optimization or the boundary of 95% confidence interval of $\log OR$ is beyond positive or negative 100. In general, the problem of nearly or exactly empty cells does limit the utility of ML procedures, particularly given that rare exposures and small validation sample-sizes are common in epidemiological settings. In contrast, the performance of the Bayesian procedures evidenced in Table 3 based on 2000 simulations seems quite reasonable, with dramatic MSE reductions for the Bayes-DF inferences compared to ML-DF. The smoothing which results from combining prior distributions on sensitivity and specificity with empty or near-empty validation-data cells appears to yield much more satisfactory inferences. Results for SIMEX estimators in the low exposure setting are not shown, but again the overall performance is worse than Bayes and ML procedures, and the use of “plugged-in” sensitivity and specificity estimates leads to the “empty-cell” concerns as with ML methods in calculating the observed misclassification matrix when $\lambda = 1$, and its exponential of a negative size when $\lambda < 1$.

Results for the nearly nondifferential Bayesian (Bayes-NNDF) analysis, in both low and high exposure prevalence settings, appear in Table 4. For the sake of comparison, results are also given here for a two-stage non-Bayesian procedure that we refer to as *test-then-estimate* (TTE). The first TTE step applied a likelihood ratio test to the validation data, with the null hypothesis that the binary exposure is nondifferentially misclassified. Then as the second step ML-DF or ML-NDF point and interval estimates are reported, depending

on whether the null is rejected or not in the first step. For some datasets one or more empty validation cell a_{ij} results in zero- or one-valued estimate for sensitivity, specificity or exposure prevalence hence yields nonsensical likelihood ratios, so that TTE estimates and inferential results are reported for only a subset of the simulated datasets. The number of discarded datasets is higher in Table 4 than in Tables 2 and 3, for more simulated datasets have one or more empty cell than those having empty (a_{i1}, a_{i2}) pair(s) simultaneously. Again empty pair (a_{i1}, a_{i2}) causes undefined $\log\widehat{OR}$ in QI method and results for a subset are presented. As before, the Bayes-NNDF results are reported for all 2000 datasets.

In terms of both point and interval estimator performance, Bayes-NNDF is seen to be moderately better than TTE (in terms of MSE, sample variance and coverage) in the high exposure prevalence setting, and very substantially better than TTE in the low prevalence setting. Comparing Table 4 to previous tables, Bayes-NNDF is seen to offer satisfactory average performance across scenarios, particularly in relation to either Bayes-NDF or Bayes-DF applied in a “wrong” scenario, although we notice that, point estimate of $\log OR$ from Bayes-NNDF is more biased and leads to increase of MSE when misclassification is much differential.

4 Example: Maternal use of antibiotics during pregnancy and sudden infant death syndrome

We consider a case-control study on sudden infant death syndrome (SIDS) [16] to further illustrate how Bayes, ML and SIMEX adjustments for misclassification work in practice. During investigation of a potential impact of maternal use of antibiotics during pregnancy on the occurrence of SIDS, surrogate exposure X was obtained from an interview question (yes=1, no=0). Information on antibiotic use from medical records, taken to be the actual exposure status T , was extracted for a subset of study participants. The data are shown in Table 5. Ignoring misclassification, the $X - Y$ log odds ratio is estimated as 0.352 with 95%

confidence interval (0.101, 0.603).

The same prior distributions used in the simulation studies of Section 3 are employed here for drawing Bayesian inferences, except that a noninformative prior for $\text{logit}(r_i)$ is used ($\mu = -1.946$, $\sigma = 100$). Study results after the various adjustments for misclassification are presented in Table 6. Estimated X-Y and T-Y logOR s from validation data are also included for model comparison. Point and interval estimates of logOR via Bayes and ML methods are similar. Parameters are estimated with slightly more certainty under the nondifferential assumption than under the differential assumption, which is consistent with simulation findings. Compared with Bayesian and ML estimates, the quality indices (QI) point estimate is smaller and has a larger SE, which is again consistent with the simulation results. Its confidence interval covering zero suggests no evidence of T-Y association, in concordance with the Bayes and ML methods under differential misclassification. Given moderate size of the validation data, a 47% increase of X-Y log odds ratio from the actual T-Y logOR on validation data suggests deviation from completely nondifferential misclassification, and the unadjusted $\widehat{\text{logOR}}$ on whole data could be falsely large. The DF, nearly NDF and QI methods demonstrate capability to correct for such bias.

Note that a considerably stronger exposure-disease association is estimated under the nondifferential misclassification assumption than under the differential misclassification assumption, with ‘significance’ (i.e., interval estimate excluding zero) in the former case but not the latter. Moreover, the validation data evidence concerning differentiability is equivocal (likelihood ratio test P-value of 0.096 for the null hypothesis of nondifferential misclassification). Therefore, the Bayes-NNDF analysis may be viewed as an appropriate compromise between the nondifferential and differential analyses, with a tempered point estimate (relative to NDF) but still significant interval estimate.

As simulation results suggest a quadratic extrapolation function together with asymptotic variance estimator performs better than alternatives, we report this SIMEX estimate in the table. In line with the Bayes and ML results, adding further misclassification pushes

estimates toward the null in the nondifferential case but away from the null in the differential case. In the nondifferential case, both choices of extrapolation function appear to fit the simulated data well. Extrapolating back to the no misclassification setting, however, produces adjusted estimates which are much more extreme than those obtained by either Bayes or ML method.

5 Example: HSV-2 and invasive cervical cancer

The second example describes a case-control study consisting of 732 subjects of cervical cancer and 1312 community or hospital controls with negative cervical cancer diagnosis [17]. Researchers were interested in assessing the impact of herpes simplex virus type 2 (HSV-2, a binary variable) in the development of invasive cervical cancer. The exposure status was detected by the western blot assay, which produced error-prone measurements. A refined, more accurate procedure was performed on a randomly selected sample of study subjects (selected without regard to their disease status), in order to assess the misclassification rates. The data are displayed in Table 7. It is noticeable from the validation and main data that the exposure prevalence of HSV-2 is high in both cases and controls. Carroll et al. observed from the validation sample that the misclassification differs between cases and controls (Fisher’s exact two-sided test implied a greater sensitivity for the cases, $p=0.049$), and proposed a pseudo-likelihood model to adjust for the differential measurement error [18].

Ignoring measurement error arising from the inaccurate western blot procedure, the naive log odds ratio is estimated as 0.453 (standard error = 0.093), with 95% confidence interval (0.271, 0.635), indicating HSV-2 is positively correlated with the occurrence of invasive cervical cancer. We conduct Bayesian adjustment under three misclassification situations (NDF, NNDF and DF), again using the prior distributions for *logit* transformed sensitivities and specificities described in Section 3. For *logit*(r_i), a flat prior with large variance is used here to generate posterior inference ($\mu=-1.946$, $\sigma=100$). Similar results are observed when

same hyperparameters for $\text{logit}(r_i)$ stated in Section 3 are used ($\mu=-1.946$, $\sigma=0.993$).

Table 8 presents results of the various analyses. For all three methods (Bayes, ML and SIMEX under the more appropriate quadratic extrapolation), moving from the nondifferential assumption to the differential assumption moves the point estimate of the exposure-disease association toward the null, and causes the left endpoint of the interval estimate to move from positive to negative, i.e., “significance” is lost. The QI method posts no assumption on misclassification between two groups, with odds ratios estimated separately from cases and controls, hence can be treated as an inherent differential adjustment. The QI estimate is closest to zero association compared with other DF methods. It is interesting to observe that only $\log\widehat{OR}$ s from QI and SIMEX models are smaller than the unadjusted X-T log odds ratio of on the whole data assuming differential misclassification. An interesting point about this example is that with a small validation sample the Bayes and ML differential analysis can yield an adjustment in a different direction than the QI analysis. That is, the validation data are such that the T-Y and X-Y marginals suggest, albeit with much uncertainty, that differential misclassification is inducing a stronger association between X and Y than between T and Y. However, looking at the three-way T-X-Y relations in the validation data (i.e., estimating case-specific and control-specific sensitivity and specificity) suggests, also with considerable uncertainty, that misclassification has the attenuating effect of yielding a weaker association between X and Y than between T and Y. While this observation is curious, it should be tempered by the fact that the Bayes and ML interval estimates are quite compatible with the QI interval estimate.

As Carroll, Gail, and Lubin [18] pointed out, there is moderate evidence to show measurement error is differential across cases and controls. Sensitivities estimated from validation data alone are 0.78 for cases and 0.5 for controls. Nevertheless, if both the complete and incomplete data are considered, a likelihood ratio test for the nondifferentiability of misclassification with 2 degrees of freedom, generates a p-value at 0.073, indicating lack of evidence to reject the null at 5% significance level. The same test based merely on the validation data re-

ports a consistent result (p-value= 0.084). Hence, it seems more appropriate to interpret the differentiability of measurement as borderline. One advantage of Bayesian adjustment emerges in this context, as it can incorporate the “in-between” scenario of nearly nondifferential misclassification via an appropriate prior distribution. As expected, the NNDF analysis yields a posterior mean and SD falling in between those arising from the NDF and DF assumptions. The resulting interval estimate is wholly positive, providing evidence for a positive exposure-disease association without concern about imposing an overly-strong assumption of nondifferential misclassification.

As a final point, we note that the Bayesian parameter estimates are consistent with the results given by Skrondal and Rabe-Hesketh [8] for these data, using generalized latent variable modeling techniques.

6 Discussion

Mismeasurement of exposure is an issue of broad concern in epidemiological studies, and there is a substantial literature on adjusting inferences on exposure-disease relationships in light of such mismeasurement. Bayesian methods, likelihood methods, and SIMEX methods are three general tools for implementing such adjustments. At least in the context of misclassified binary exposure, this paper has illustrated several positive attributes of the Bayesian approach. First, Bayesian methods can provide more reasonable and stable inferences when the resulting data are sparse, which is of particular relevance to small validation datasets in rare exposure contexts. Second, the infusion of prior information offered by the Bayesian approach can be used to good effect. Rather than committing to nondifferential or ‘fully’ differential assumptions concerning the exposure misclassification, a prior can be constructed to represent a ‘nearly nondifferential’ assumption. That is, the analyst can assert that substantial deviations from nondifferentiability are unlikely. This would seem to be a particularly useful device when the data themselves do not clearly support or refute nondifferentiability,

as occurred in both our real-data examples.

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Table 1: Validation data and main data

	<i>Validation Data</i>				<i>Main Data</i>			
	Y=1		Y=0		Y=1		Y=0	
T	X=1	X=0	X=1	X=0	X=1	X=0	X=1	X=0
T=1	a_{11}	a_{12}	a_{01}	a_{02}	b_{11}	b_{12}	b_{01}	b_{02}
T=0	a_{13}	a_{14}	a_{03}	a_{04}	b_{13}	b_{14}	b_{03}	b_{04}
N	$a_{11} + a_{13}$	$a_{12} + a_{14}$	$a_{01} + a_{03}$	$a_{02} + a_{04}$	a_{15}	a_{16}	a_{05}	a_{06}

Table 2: Comparative performance models on simulated datasets of size 1760 ($N_{rep}=2000$) given high exposure prevalences and $OR = 1.25$

	ML		Bayes		QI	SIMEX-NDF				SIMEX-DF				
	NDF	ML DF	NDF	Bayes DF		Quadratic ASY.	JCK.	Loglinear ASY.	JCK.	Quadratic ASY.	JCK.	Loglinear ASY.	JCK.	
Scenario 1	MSE	0.023	0.073	0.024	0.052	0.082	0.023	0.023	0.024	0.024	0.108	0.108	0.213	0.213
	Bias	0.007	0.009	0.013	0.005	0.008	-0.005	-0.005	0.004	0.004	-0.007	-0.007	0.084	0.084
	Variance	0.023	0.073	0.024	0.052	0.082	0.023	0.023	0.024	0.024	0.108	0.108	0.206	0.206
	Coverage	0.957	0.961	0.952	0.979	0.962	0.895	0.871	0.856	0.86	0.539	0.509	0.599	0.744
Width	0.598	1.06	0.62	1.007	1.138	0.497	0.463	0.498	0.463	0.498	0.463	0.471	0.463	
Scenario 2	MSE	0.05	0.087	0.051	0.065	0.1	0.049	0.049	0.059	0.059	0.134	0.134	0.164	0.164
	Bias	0.147	0	0.15	0.011	0.003	0.151	0.151	0.171	0.171	0.022	0.022	0.158	0.158
	Variance	0.029	0.088	0.028	0.065	0.1	0.027	0.027	0.03	0.03	0.133	0.133	0.139	0.139
	Coverage	0.872	0.945	0.853	0.96	0.945	0.758	0.693	0.717	0.648	0.533	0.488	0.65	0.726
Width	0.651	1.118	0.664	1.05	1.215	0.525	0.478	0.544	0.478	0.524	0.475	0.423	0.475	
Scenario 3	MSE	0.124	0.099	0.119	0.076	0.117	0.133	0.133	0.174	0.174	0.159	0.159	0.424	0.424
	Bias	0.302	0.012	0.297	0.028	0.011	0.32	0.32	0.366	0.366	0.066	0.066	0.259	0.259
	Variance	0.033	0.099	0.031	0.075	0.117	0.031	0.031	0.04	0.04	0.155	0.155	0.357	0.357
	Coverage	0.638	0.939	0.615	0.948	0.944	0.419	0.35	0.378	0.277	0.517	0.463	0.608	0.664
Width	0.711	1.163	0.711	1.087	1.282	0.557	0.494	0.599	0.494	0.555	0.487	0.472	0.487	
Scenario 4	MSE	0.245	0.098	0.231	0.076	0.122	0.295	0.295	0.418	0.418	0.194	0.194	0.298	0.298
	Bias	0.456	-0.004	0.443	0.019	-0.004	0.508	0.508	0.599	0.599	0.124	0.124	0.336	0.336
	Variance	0.038	0.098	0.035	0.076	0.122	0.037	0.037	0.059	0.059	0.179	0.179	0.185	0.185
	Coverage	0.329	0.944	0.316	0.953	0.944	0.121	0.084	0.095	0.052	0.48	0.423	0.387	0.526
Width	0.761	1.195	0.75	1.116	1.34	0.587	0.506	0.661	0.506	0.586	0.495	0.442	0.495	

Table 3: Comparative performance of ML, QI ($N'_{rep} < 2000$) and Bayes, SIMEX ($N_{rep} = 2000$) models on simulated datasets of size 1760, given low exposure prevalences and $OR = 1.25$

	ML			ML			Bayes			SIMEX-NDF			SIMEX-DF				
	NDF	N'_{rep}	DF	DF	N'_{rep}	NDF	DF	QI	N'_{rep}	Quadratic ASY.	JCK.	Loglinear ASY.	JCK.	Quadratic ASY.	JCK.	Loglinear ASY.	JCK.
Scenario 1	MSE	0.276	1898	0.574	1889	0.152	0.205	0.59	1889								
	Bias	0.008		-0.039		-0.036	-0.081	-0.032									
	Variance	0.276		0.572		0.151	0.198	0.589									
	Coverage	0.984		0.983		0.962	0.984	0.979									
	Width	2.154		3.092		1.74	2.252	3.165									
Scenario 2	MSE	1.145	1901	0.565	1884	0.621	0.208	0.576	1884								
	Bias	0.93		-0.029		0.696	-0.011	-0.033									
	Variance	0.279		0.564		0.137	0.208	0.575									
	Coverage	0.651		0.987		0.597	0.986	0.982									
	Width	2.242		3.124		1.722	2.282	3.202									
Scenario 3	MSE	2.916	1908	0.599	1874	1.735	0.221	0.619	1874								
	Bias	1.617		-0.017		1.269	0.011	-0.028									
	Variance	0.302		0.599		0.125	0.221	0.619									
	Coverage	0.057		0.984		0.076	0.984	0.982									
	Width	2.272		3.176		1.651	2.314	3.258									
Scenario 4	MSE	4.815	1928	0.617	1897	3.073	0.22	0.643	1897								
	Bias	2.129		-0.023		1.725	0.026	-0.021									
	Variance	0.282		0.616		0.099	0.22	0.643									
	Coverage	0.001		0.984		0.001	0.989	0.981									
	Width	2.278		3.177		1.591	2.323	3.27									

Table 4: Comparative performance of ML-TTE ($N'_{rep} < 2000$) and Bayes-NNDF ($N_{rep} = 2000$) on simulated datasets of size 1760, $OR = 1.25$

		High prevalences			Low prevalences		
		Bayes-NNDF	ML-TTE	N'_{rep}	Bayes-NNDF	ML-TTE	N'_{rep}
Scenario 1	MSE	0.025	0.027	1943	0.148	0.19	481
	Bias	0.01	0.01		-0.056	-0.011	
	Variance	0.025	0.027		0.145	0.19	
	Coverage	0.977	0.954		0.988	0.985	
	Width	0.728	0.617		1.971	1.877	
Scenario 2	MSE	0.042	0.059	1960	0.286	0.841	526
	Bias	0.108	0.116		0.364	0.723	
	Variance	0.031	0.045		0.154	0.319	
	Coverage	0.942	0.872		0.933	0.656	
	Width	0.787	0.705		2.038	2.013	
Scenario 3	MSE	0.077	0.117	1955	0.602	1.636	563
	Bias	0.205	0.178		0.654	0.892	
	Variance	0.035	0.085		0.175	0.841	
	Coverage	0.875	0.732		0.788	0.334	
	Width	0.849	0.851		2.104	2.184	
Scenario 4	MSE	0.121	0.16	1971	0.93	2.001	589
	Bias	0.284	0.166		0.864	0.799	
	Variance	0.041	0.132		0.184	1.365	
	Coverage	0.784	0.67		0.652	0.52	
	Width	0.904	0.999		2.162	2.378	

Table 5: Validation study and main study of SIDS

	<i>Validation Data</i>				<i>Main Data</i>			
	Y=1		Y=0		Y=1		Y=0	
T	X=1	X=0	X=1	X=0	X=1	X=0	X=1	X=0
T=1	29	17	21	16	b_{11}	b_{12}	b_{01}	b_{02}
T=0	22	143	12	168	b_{13}	b_{14}	b_{03}	b_{04}
N	51	160	33	184	122	442	101	479

Table 6: $\log\widehat{OR}$, SE and 95% intervals for $\log OR$ in SIDS study based on a flat prior

	$\log(\widehat{OR})$	SE	95% intervals
Unadjusted-whole data	0.352	0.128	(0.101, 0.603)
Unadjusted-validation data	0.575	0.248	(0.089, 1.020)
Adjusted-validation data	0.305	0.246	(-0.177, 0.786)
Bayes-NDF	0.396	0.193	(0.016, 0.770)
Bayes-NNDF	0.329	0.200	(-0.060, 0.723)
Bayes-DF	0.208	0.219	(-0.227, 0.637)
ML-NDF	0.398	0.191	(0.024, 0.772)
ML-DF	0.193	0.221	(-0.241, 0.626)
QI	0.082	0.246	(-0.400, 0.563)
SIMEX-NDF <i>quadratic, asymptotic</i>	0.663	0.227	(0.219, 1.108)
SIMEX-DF <i>quadratic, asymptotic</i>	-0.010	0.221	(-0.443, 0.424)

Table 7: Validation data and main data for cervical cancer study

	<i>Validation Data</i>				<i>Main Data</i>			
	Y=1		Y=0		Y=1		Y=0	
T	X=1	X=0	X=1	X=0	X=1	X=0	X=1	X=0
T=1	18	5	16	16	b_{11}	b_{12}	b_{01}	b_{02}
T=0	3	13	11	33	b_{13}	b_{14}	b_{03}	b_{04}
N	21	18	27	49	375	318	525	701

Table 8: $\log\widehat{OR}$, SE and 95% intervals for $\log OR$ in cervical cancer study based on a flat prior for $\text{logit}(r_i)$

	$\log(\widehat{OR})$	SE	95% interval
Unadjusted-whole data	0.453	0.093	(0.271, 0.635)
Unadjusted-validation data	0.750	0.401	(-0.035, 1.536)
Adjusted-validation data	0.681	0.400	(-0.103, 1.465)
Bayes-NDF	0.921	0.223	(0.520, 1.404)
Bayes- NNDF	0.809	0.266	(0.320, 1.356)
Bayes-DF	0.583	0.324	(-0.033, 1.262)
ML-NDF	0.958	0.237	(0.494, 1.422)
ML-DF	0.608	0.350	(-0.079, 1.295)
QI	0.384	0.411	(-0.421, 1.189)
SIMEX-NDF			
<i>quadratic, asymptotic</i>	0.903	0.184	(0.542, 1.264)
SIMEX-DF			
<i>quadratic, asymptotic</i>	0.146	0.172	(-0.191, 0.482)

Table 9: Comparative performance models on simulated datasets of size 1760 ($N_{rep}=2000$) given high exposure prevalences and $OR = 1.8$

	ML		Bayes		Bayes		QI	SIMEX-NDF				SIMEX-DF			
	NDF	DF	NDF	DF	NDF	DF		Quadratic ASY.	JCK.	Loglinear ASY.	JCK.	Quadratic ASY.	JCK.	Loglinear ASY.	JCK.
Scenario 1	MSE	0.023	0.07	0.023	0.05	0.081	0.023	0.023	0.024	0.024	0.024	0.111	0.111	1.768	1.768
	Bias	0.005	0.007	0.017	-0.001	0.002	-0.022	-0.022	-0.005	-0.005	-0.005	-0.031	-0.031	0.16	0.16
	Variance	0.023	0.07	0.023	0.05	0.081	0.022	0.022	0.024	0.024	0.024	0.111	0.111	1.743	1.743
	Coverage	0.955	0.95	0.954	0.97	0.945	0.883	0.858	0.881	0.85	0.85	0.551	0.523	0.488	0.532
Width	0.603	1.027	0.618	0.97	1.105	0.484	0.453	0.494	0.453	0.453	0.485	0.451	0.59	0.451	
Scenario 2	MSE	0.041	0.074	0.041	0.054	0.089	0.035	0.035	0.048	0.048	0.125	0.125	1.151	1.151	
	Bias	0.118	0.009	0.122	0.008	0.011	0.099	0.099	0.133	0.133	-0.005	-0.005	0.159	0.159	
	Variance	0.027	0.073	0.026	0.054	0.089	0.026	0.026	0.031	0.031	0.125	0.125	1.126	1.126	
	Coverage	0.927	0.954	0.896	0.968	0.951	0.835	0.79	0.787	0.723	0.536	0.498	0.534	0.598	
Width	0.657	1.074	0.661	1.006	1.169	0.513	0.469	0.535	0.469	0.51	0.465	0.514	0.465		
Scenario 3	MSE	0.088	0.082	0.083	0.062	0.1	0.086	0.086	0.135	0.135	0.146	0.146	1.698	1.698	
	Bias	0.236	0	0.229	0.004	-0.006	0.231	0.231	0.298	0.298	-0.015	-0.015	0.165	0.165	
	Variance	0.033	0.082	0.03	0.062	0.1	0.032	0.032	0.046	0.046	0.146	0.146	1.671	1.671	
	Coverage	0.783	0.951	0.747	0.962	0.945	0.597	0.538	0.514	0.42	0.544	0.49	0.617	0.692	
Width	0.717	1.116	0.707	1.041	1.236	0.548	0.486	0.591	0.486	0.543	0.48	0.525	0.48		
Scenario 4	MSE	0.159	0.088	0.143	0.068	0.11	0.168	0.168	0.293	0.293	0.166	0.166	1.398	1.398	
	Bias	0.349	-0.008	0.331	-0.001	-0.013	0.361	0.361	0.477	0.477	0.009	0.009	0.227	0.227	
	Variance	0.037	0.088	0.034	0.068	0.11	0.037	0.037	0.065	0.065	0.166	0.166	1.347	1.347	
	Coverage	0.603	0.949	0.571	0.956	0.951	0.364	0.302	0.255	0.178	0.519	0.45	0.628	0.69	
Width	0.769	1.147	0.748	1.07	1.292	0.578	0.498	0.648	0.498	0.574	0.488	0.551	0.488		

Table 10: Comparative performance models on simulated datasets of size 1760 ($N_{rep}=2000$) given high exposure prevalences and $OR = 2.5$

	ML		Bayes		QI	SIMEX-NDF				SIMEX-DF				
	NDF	DF	NDF	DF		Quadratic ASY.	JCK.	Loglinear ASY.	JCK.	Quadratic ASY.	JCK.	Loglinear ASY.	JCK.	
Scenario 1	MSE	0.028	0.072	0.027	0.052	0.081	0.026	0.026	0.028	0.028	0.107	0.107	3.205	3.205
	Bias	0.01	0.013	0.025	-0.001	0.008	-0.026	-0.026	-0.001	-0.001	-0.035	-0.035	0.27	0.27
	Variance	0.028	0.072	0.026	0.052	0.081	0.025	0.025	0.028	0.028	0.106	0.106	3.134	3.134
	Coverage	0.949	0.94	0.946	0.96	0.948	0.86	0.821	0.853	0.807	0.518	0.486	0.365	0.369
	Width	0.627	1.012	0.633	0.953	1.086	0.476	0.443	0.485	0.443	0.477	0.444	0.575	0.444
Scenario 2	MSE	0.041	0.076	0.038	0.055	0.088	0.032	0.032	0.049	0.049	0.12	0.12	1.927	1.927
	Bias	0.098	0.009	0.1	-0.002	0.007	0.064	0.064	0.112	0.112	-0.037	-0.037	0.19	0.19
	Variance	0.031	0.076	0.028	0.055	0.088	0.028	0.028	0.036	0.036	0.119	0.119	1.892	1.892
	Coverage	0.943	0.944	0.921	0.953	0.946	0.858	0.808	0.789	0.724	0.539	0.501	0.448	0.458
	Width	0.685	1.059	0.678	0.989	1.153	0.507	0.465	0.528	0.465	0.503	0.46	0.545	0.46
Scenario 3	MSE	0.066	0.08	0.058	0.06	0.097	0.055	0.055	0.101	0.101	0.146	0.146	2.768	2.768
	Bias	0.178	-0.001	0.165	-0.011	-0.003	0.149	0.149	0.231	0.231	-0.054	-0.054	0.2	0.2
	Variance	0.035	0.08	0.031	0.059	0.097	0.033	0.033	0.048	0.048	0.143	0.143	2.729	2.729
	Coverage	0.895	0.958	0.871	0.965	0.952	0.753	0.683	0.627	0.527	0.524	0.478	0.446	0.474
	Width	0.741	1.097	0.721	1.02	1.215	0.54	0.481	0.581	0.481	0.535	0.473	0.591	0.473
Scenario 4	MSE	0.117	0.082	0.097	0.062	0.105	0.1	0.1	0.224	0.224	0.165	0.165	2.081	2.081
	Bias	0.279	0.016	0.253	0.004	0.01	0.253	0.253	0.394	0.394	-0.054	-0.054	0.168	0.168
	Variance	0.038	0.081	0.033	0.062	0.105	0.037	0.037	0.069	0.069	0.163	0.163	2.054	2.054
	Coverage	0.779	0.947	0.756	0.957	0.939	0.587	0.505	0.407	0.299	0.511	0.441	0.477	0.512
	Width	0.799	1.127	0.766	1.049	1.272	0.574	0.496	0.643	0.496	0.569	0.484	0.577	0.484

Table 11: Comparative performance of ML, QI ($N'_{rep} < 2000$) and Bayes, SIMEX ($N_{rep} = 2000$) models on simulated datasets of size 1760, given low exposure prevalences and $OR = 1.8$

	ML			ML		Bayes		SIMEX-NDF			SIMEX-DF				
	NDF	N'_{rep}	DF	DF	N'_{rep}	NDF	DF	QI	N'_{rep}	Quadratic ASY.	Loglinear ASY.	JCK.	Quadratic ASY.	Loglinear ASY.	JCK.
Scenario 1	MSE	0.231	1931	0.498	1921	0.136	0.224	0.51	1921						
	Bias	0.024		-0.018		-0.084	-0.178	-0.027							
	Variance	0.23		0.498		0.129	0.192	0.509							
	Coverage	0.961		0.974		0.956	0.97	0.97							
	Width	1.957		2.802		1.602	2.116	2.874							
Scenario 2	MSE	0.945	1912	0.541	1910	0.444	0.207	0.544	1910						
	Bias	0.826		0.032		0.569	-0.082	0.02							
	Variance	0.263		0.541		0.119	0.2	0.544							
	Coverage	0.81		0.974		0.699	0.982	0.968							
	Width	2.176		2.883		1.629	2.16	2.966							
Scenario 3	MSE	2.173	1925	0.53	1921	1.206	0.216	0.554	1921						
	Bias	1.381		-0.021		1.049	-0.087	-0.022							
	Variance	0.267		0.53		0.105	0.209	0.553							
	Coverage	0.14		0.982		0.123	0.986	0.977							
	Width	2.206		2.885		1.584	2.173	2.972							
Scenario 4	MSE	3.756	1921	0.488	1912	2.236	0.209	0.519	1912						
	Bias	1.864		-0.001		1.463	-0.072	-0.009							
	Variance	0.282		0.488		0.096	0.204	0.519							
	Coverage	0.004		0.984		0.004	0.986	0.977							
	Width	2.275		2.921		1.572	2.202	3.019							

Table 12: Comparative performance of ML, QI ($N'_{rep} < 2000$) and Bayes, SIMEX ($N_{rep} = 2000$) models on simulated datasets of size 1760, given low exposure prevalences and $OR = 2.5$

	ML			ML		Bayes		QI	N'_{rep}	SIMEX-NDF			SIMEX-DF		
	NDF	N'_{rep}	DF	DF	N'_{rep}	NDF	DF			Quadratic ASY. JCK. ASY. JCK.	Loglinear JCK. ASY. JCK.	Quadratic ASY. JCK. ASY. JCK.	Loglinear JCK. ASY. JCK.		
Scenario 1	MSE	0.226	1925	0.438	1921	0.123	0.209	0.452	1921						
	Bias	0.045		0.037		-0.088	-0.188	0.045							
	Variance	0.224		0.437		0.115	0.174	0.45							
	Coverage	0.949		0.971		0.958	0.971	0.965							
	Width	1.929		2.628		1.52	2.006	2.699							
Scenario 2	MSE	0.733	1905	0.428	1904	0.305	0.207	0.449	1904						
	Bias	0.7		0.012		0.448	-0.161	0.001							
	Variance	0.242		0.428		0.105	0.181	0.449							
	Coverage	0.917		0.965		0.833	0.969	0.962							
	Width	2.12		2.687		1.562	2.049	2.765							
Scenario 3	MSE	1.729	1940	0.466	1939	0.853	0.215	0.494	1939						
	Bias	1.218		0.01		0.875	-0.15	0							
	Variance	0.246		0.466		0.089	0.193	0.495							
	Coverage	0.319		0.965		0.247	0.966	0.953							
	Width	2.211		2.719		1.555	2.069	2.808							
Scenario 4	MMSE	3.002	1934	0.467	1931	1.622	0.211	0.491	1931						
	Bias	1.652		0.029		1.239	-0.131	0.023							
	Variance	0.273		0.466		0.086	0.194	0.491							
	Coverage	0.015		0.974		0.015	0.977	0.97							
	Width	2.295		2.774		1.56	2.099	2.876							

Table 13: Comparative performance of ML-TTE ($N'_{rep} < 2000$) and Bayes-NNDF ($N_{rep} = 2000$) on simulated datasets of size 1760, $OR = 1.8$

		High prevalences			Low prevalences		
		Bayes-NNDF	ML-TTE	N'_{rep}	Bayes-NNDF	ML-TTE	N'_{rep}
Scenario 1	MSE	0.025	0.028	1950	0.147	0.16	599
	Bias	0.011	0.002		-0.126	-0.036	
	Variance	0.024	0.028		0.131	0.159	
	Coverage	0.979	0.949		0.974	0.955	
	Width	0.71	0.623		1.824	1.762	
Scenario 2	MSE	0.035	0.048	1969	0.217	0.62	653
	Bias	0.089	0.089		0.285	0.579	
	Variance	0.027	0.04		0.136	0.286	
	Coverage	0.962	0.923		0.957	0.828	
	Width	0.76	0.702		1.897	1.968	
Scenario 3	MSE	0.057	0.09	1966	0.435	1.232	734
	Bias	0.157	0.142		0.538	0.741	
	Variance	0.032	0.07		0.146	0.684	
	Coverage	0.915	0.83		0.827	0.38	
	Width	0.816	0.83		1.938	2.056	
Scenario 4	MSE	0.083	0.119	1959	0.724	1.537	741
	Bias	0.216	0.116		0.753	0.59	
	Variance	0.037	0.106		0.158	1.19	
	Coverage	0.867	0.801		0.68	0.529	
	Width	0.865	0.972		2.002	2.309	

Table 14: Comparative performance of ML-TTE ($N'_{rep} < 2000$) and Bayes-NNDF ($N_{rep} = 2000$) on simulated datasets of size 1760, $OR = 2.5$

		High prevalences			Low prevalences		
		Bayes-NNDF	ML-TTE	N'_{rep}	Bayes-NNDF	ML-TTE	N'_{rep}
Scenario 1	MSE	0.028	0.032	1941	0.134	0.16	703
	Bias	0.017	0.005		-0.132	-0.046	
	Variance	0.027	0.032		0.117	0.158	
	Coverage	0.969	0.943		0.97	0.943	
	Width	0.71	0.646		1.715	1.7	
Scenario 2	MSE	0.035	0.046	1959	0.157	0.446	753
	Bias	0.072	0.076		0.201	0.433	
	Variance	0.029	0.041		0.116	0.259	
	Coverage	0.968	0.939		0.979	0.935	
	Width	0.76	0.723		1.778	1.878	
Scenario 3	MSE	0.044	0.07	1959	0.325	1.035	800
	Bias	0.112	0.108		0.451	0.628	
	Variance	0.032	0.058		0.122	0.642	
	Coverage	0.955	0.914		0.882	0.493	
	Width	0.808	0.835		1.82	2.06	
Scenario 4	MSE	0.064	0.095	1960	0.563	1.2	771
	Bias	0.171	0.103		0.655	0.481	
	Variance	0.034	0.084		0.134	0.97	
	Coverage	0.918	0.869		0.722	0.503	
	Width	0.858	0.976		1.878	2.219	

Table 15: Comparative performance models on simulated datasets of size 960 ($N_{rep}=2000$) given high exposure prevalences and $OR = 1.8$

	ML		Bayes		QI	SIMEX-NDF				SIMEX-DF			
	NDF	DF	NDF	DF		Quadratic ASY.	JCK.	Loglinear ASY.	JCK.	Quadratic ASY.	JCK.	Loglinear ASY.	JCK.
Scenario 1	MSE	0.038	0.075	0.038	0.057	0.083	0.041	0.041	0.043	0.043	0.041	0.041	0.043
	Bias	0.01	0.011	0.015	0.001	0.008	-0.015	-0.015	0.002	0.002	0.008	-0.015	0.002
	Variance	0.038	0.075	0.038	0.057	0.083	0.041	0.041	0.043	0.043	0.041	0.041	0.043
	Coverage	0.952	0.951	0.95	0.965	0.952	0.9	0.878	0.892	0.863	0.952	0.9	0.878
	Width	0.765	1.061	0.777	1.015	1.13	0.655	0.611	0.668	0.611	1.13	0.655	0.611
Scenario 2	MSE	0.055	0.084	0.052	0.065	0.095	0.058	0.058	0.076	0.076	0.058	0.058	0.076
	Bias	0.099	0.004	0.095	0.001	-0.002	0.098	0.098	0.136	0.136	0.098	0.098	0.136
	Variance	0.045	0.084	0.043	0.065	0.095	0.049	0.049	0.058	0.058	0.049	0.049	0.058
	Coverage	0.927	0.951	0.92	0.963	0.952	0.854	0.824	0.83	0.77	0.952	0.854	0.83
	Width	0.813	1.1	0.813	1.043	1.185	0.694	0.635	0.723	0.635	1.185	0.694	0.635
Scenario 3	MSE	0.087	0.088	0.078	0.069	0.101	0.109	0.109	0.162	0.162	0.109	0.109	0.162
	Bias	0.197	0.004	0.182	0.006	0.006	0.23	0.23	0.299	0.299	0.006	0.23	0.299
	Variance	0.048	0.088	0.045	0.069	0.101	0.056	0.056	0.073	0.073	0.101	0.056	0.073
	Coverage	0.891	0.948	0.884	0.958	0.951	0.728	0.654	0.653	0.543	0.951	0.728	0.653
	Width	0.868	1.142	0.856	1.077	1.249	0.742	0.659	0.798	0.659	1.249	0.742	0.659
Scenario 4	MSE	0.137	0.092	0.119	0.074	0.113	0.198	0.198	0.344	0.344	0.113	0.198	0.344
	Bias	0.285	0.002	0.26	0.003	0	0.366	0.366	0.489	0.489	0	0.366	0.489
	Variance	0.056	0.092	0.051	0.074	0.113	0.064	0.064	0.104	0.104	0.113	0.064	0.104
	Coverage	0.796	0.952	0.795	0.96	0.953	0.553	0.468	0.458	0.323	0.953	0.553	0.468
	Width	0.917	1.169	0.895	1.101	1.303	0.788	0.677	0.887	0.677	1.303	0.788	0.677

Table 16: Comparative performance models on simulated datasets of size 960 ($N_{rep}=2000$) given high exposure prevalences and $OR = 1.25$

	ML		Bayes		QI	SIMEX-NDF				SIMEX-DF				
	NDF	ML DF	NDF	Bayes DF		Quadratic ASY.	JCK.	Loglinear ASY.	JCK.	Quadratic ASY.	JCK.	Loglinear ASY.	JCK.	
Scenario 1	MSE	0.04	0.078	0.041	0.06	0.087	0.044	0.044	0.045	0.045	0.11	0.11	0.126	0.126
	Bias	0.006	0	0.008	-0.002	-0.001	-0.005	-0.005	0.008	0.008	-0.014	-0.014	0.073	0.073
	Variance	0.04	0.078	0.041	0.06	0.087	0.044	0.044	0.045	0.045	0.11	0.11	0.121	0.121
	Coverage	0.954	0.958	0.951	0.969	0.955	0.893	0.867	0.86	0.862	0.7	0.666	0.72	0.807
	Width	0.776	1.094	0.795	1.051	1.163	0.673	0.626	0.665	0.626	0.674	0.626	0.578	0.626
Scenario 2	MSE	0.059	0.09	0.058	0.071	0.104	0.07	0.07	0.083	0.083	0.134	0.134	0.322	0.322
	Bias	0.12	-0.009	0.118	0.001	-0.01	0.151	0.151	0.174	0.174	0.007	0.007	0.158	0.158
	Variance	0.045	0.09	0.044	0.071	0.104	0.047	0.047	0.053	0.053	0.134	0.134	0.298	0.298
	Coverage	0.93	0.951	0.924	0.965	0.946	0.813	0.762	0.789	0.737	0.667	0.632	0.752	0.805
	Width	0.829	1.148	0.837	1.09	1.234	0.713	0.649	0.731	0.649	0.711	0.646	0.646	0.646
Scenario 3	MSE	0.111	0.095	0.104	0.076	0.11	0.158	0.158	0.208	0.208	0.155	0.155	0.306	0.306
	Bias	0.237	-0.003	0.228	0.011	-0.002	0.313	0.313	0.365	0.365	0.061	0.061	0.232	0.232
	Variance	0.055	0.095	0.052	0.076	0.11	0.06	0.06	0.075	0.075	0.152	0.152	0.252	0.252
	Coverage	0.826	0.949	0.821	0.961	0.952	0.608	0.541	0.576	0.476	0.66	0.609	0.706	0.761
	Width	0.881	1.186	0.877	1.12	1.293	0.754	0.669	0.818	0.669	0.752	0.66	0.598	0.66
Scenario 4	MSE	0.195	0.097	0.179	0.079	0.119	0.323	0.323	0.463	0.463	0.186	0.186	0.315	0.315
	Bias	0.375	0.014	0.358	0.033	0.017	0.511	0.511	0.608	0.608	0.144	0.144	0.339	0.339
	Variance	0.055	0.097	0.051	0.078	0.119	0.063	0.063	0.094	0.094	0.166	0.166	0.2	0.2
	Coverage	0.655	0.952	0.659	0.955	0.955	0.313	0.248	0.282	0.181	0.645	0.564	0.538	0.64
	Width	0.93	1.214	0.915	1.144	1.345	0.795	0.686	0.891	0.686	0.794	0.668	0.614	0.668

Table 17: Comparative performance models on simulated datasets of size 960 ($N_{rep}=2000$) given high exposure prevalences and $OR = 2.5$

	ML		Bayes		QI	SIMEX-NDF				SIMEX-DF			
	NDF	DF	NDF	DF		Quadratic ASY.	JCK.	Loglinear ASY.	JCK.	Quadratic ASY.	JCK.	Loglinear ASY.	JCK.
Scenario 1	MSE	0.039	0.075	0.037	0.057	0.084	0.04	0.04	0.044	0.044			
	Bias	0.008	0.01	0.013	-0.008	0.006	-0.028	-0.028	0	0			
	Variance	0.039	0.075	0.037	0.057	0.084	0.04	0.04	0.044	0.044			
	Coverage	0.956	0.947	0.957	0.961	0.943	0.885	0.866	0.887	0.85			
	Width	0.773	1.047	0.778	0.998	1.114	0.645	0.603	0.658	0.603			
Scenario 2	MSE	0.053	0.084	0.049	0.065	0.097	0.051	0.051	0.068	0.068			
	Bias	0.079	0.012	0.07	-0.004	0.011	0.053	0.053	0.104	0.104			
	Variance	0.047	0.084	0.044	0.065	0.097	0.048	0.048	0.057	0.057			
	Coverage	0.943	0.945	0.94	0.959	0.945	0.888	0.847	0.853	0.792			
	Width	0.826	1.088	0.818	1.028	1.174	0.689	0.631	0.719	0.631			
Scenario 3	MSE	0.07	0.084	0.06	0.066	0.099	0.075	0.075	0.128	0.128			
	Bias	0.147	0.004	0.126	-0.01	0.003	0.145	0.145	0.23	0.23			
	Variance	0.049	0.084	0.044	0.066	0.099	0.054	0.054	0.075	0.075			
	Coverage	0.926	0.95	0.921	0.963	0.951	0.825	0.773	0.747	0.647			
	Width	0.876	1.121	0.856	1.055	1.228	0.732	0.653	0.788	0.653			
Scenario 4	MSE	0.11	0.091	0.089	0.072	0.111	0.131	0.131	0.268	0.268			
	Bias	0.23	0.009	0.196	-0.004	0.008	0.255	0.255	0.402	0.402			
	Variance	0.057	0.091	0.05	0.072	0.111	0.066	0.066	0.107	0.107			
	Coverage	0.862	0.95	0.866	0.957	0.95	0.717	0.637	0.579	0.446			
	Width	0.925	1.149	0.894	1.08	1.281	0.778	0.674	0.871	0.674			

Table 18: Comparative performance of ML, QI ($N'_{rep} < 2000$) and Bayes, SIMEX ($N_{rep} = 2000$) models on simulated datasets of size 960, given low exposure prevalences and $OR = 1.8$

	ML			ML		Bayes		SIMEX-NDF			SIMEX-DF			
	NDF	N'_{rep}	DF	N'_{rep}	NDF	DF	QI	N'_{rep}	Quadratic ASY.	Loglinear JCK.	Quadratic ASY.	Loglinear JCK.	Quadratic ASY.	Loglinear JCK.
Scenario 1	MSE	0.317	1917	0.51	1916	0.181	0.233	0.527	1916					
	Bias	0.02		-0.009		-0.108	-0.164	-0.01						
	Variance	0.317		0.51		0.17	0.206	0.527						
	Coverage	0.977		0.974		0.964	0.971	0.97						
	Width	2.292		2.842		1.854	2.152	2.906						
Scenario 2	MSE	0.712	1910	0.489	1911	0.28	0.203	0.5	1911					
	Bias	0.623		0.037		0.353	-0.091	0.033						
	Variance	0.325		0.488		0.156	0.195	0.5						
	Coverage	0.936		0.982		0.915	0.984	0.98						
	Width	2.46		2.897		1.873	2.187	2.968						
Scenario 3	MSE	1.489	1921	0.546	1918	0.663	0.228	0.565	1918					
	Bias	1.073		-0.025		0.715	-0.104	-0.032						
	Variance	0.338		0.546		0.151	0.217	0.564						
	Coverage	0.603		0.971		0.64	0.98	0.969						
	Width	2.429		2.917		1.845	2.205	2.998						
Scenario 4	MSE	2.582	1918	0.561	1917	1.28	0.23	0.582	1917					
	Bias	1.492		-0.017		1.066	-0.085	-0.026						
	Variance	0.357		0.561		0.145	0.223	0.581						
	Coverage	0.203		0.973		0.283	0.974	0.97						
	Width	2.424		2.931		1.805	2.215	3.021						

Table 19: Comparative performance of ML, QI ($N'_{rep} < 2000$) and Bayes, SIMEX ($N_{rep} = 2000$) models on simulated datasets of size 960, given low exposure prevalences and $OR = 1.25$

	ML			ML			Bayes			SIMEX-NDF			SIMEX-DF		
	NDF	N'_{rep}	DF	NDF	N'_{rep}	DF	NDF	DF	DF	Quadratic	Loglinear	Quadratic	Loglinear	Quadratic	Loglinear
Scenario 1	MSE	0.363	1883	0.558	1881	0.212	0.182	0.212	0.577	1881					
	Bias	-0.006		-0.023		-0.083	-0.054	-0.083	-0.022						
	Variance	0.363		0.558		0.206	0.18	0.206	0.577						
	Coverage	0.984		0.98		0.985	0.976	0.985	0.979						
	Width	2.5		3.087		2.278	1.983	2.278	3.153						
Scenario 2	MSE	0.841	1893	0.579	1887	0.212	0.376	0.212	0.596	1887					
	Bias	0.68		0.022		-0.005	0.447	-0.005	0.025						
	Variance	0.379		0.579		0.212	0.176	0.212	0.596						
	Coverage	0.881		0.986		0.987	0.883	0.987	0.986						
	Width	2.609		3.144		2.306	1.997	2.306	3.213						
Scenario 3	MSE	1.896	1911	0.625	1900	0.227	0.939	0.227	0.63	1900					
	Bias	1.215		-0.052		-0.011	0.87	-0.011	-0.06						
	Variance	0.42		0.622		0.227	0.182	0.227	0.626						
	Coverage	0.48		0.984		0.984	0.558	0.984	0.983						
	Width	2.596		3.189		2.325	1.966	2.325	3.269						
Scenario 4	MSE	3.262	1900	0.622	1877	0.228	1.744	0.228	0.638	1877					
	Bias	1.692		-0.006		0.028	1.258	0.028	-0.007						
	Variance	0.398		0.622		0.227	0.162	0.227	0.638						
	Coverage	0.131		0.986		0.984	0.226	0.984	0.984						
	Width	2.533		3.199		2.345	1.915	2.345	3.284						

Table 20: Comparative performance of ML, QI ($N'_{rep} < 2000$) and Bayes, SIMEX ($N_{rep} = 2000$) models on simulated datasets of size 960, given low exposure prevalences and $OR = 2.5$

	ML			ML			Bayes			SIMEX-NDF			SIMEX-DF			
	NDF	N'_{rep}	DF	DF	N'_{rep}		NDF	DF	QI	N'_{rep}	Quadratic	Loglinear	Quadratic	Loglinear	Quadratic	Loglinear
Scenario 1	MSE	0.297	1926	0.443	1926		0.164	0.213	0.449	1926						
	Bias	0.07		0.058			-0.113	-0.187	0.06							
	Variance	0.292		0.44			0.151	0.178	0.446							
	Coverage	0.966		0.971			0.962	0.97	0.966							
	Width	2.206		2.694			1.752	2.049	2.761							
Scenario 2	MSE	0.606	1933	0.453	1933		0.206	0.219	0.48	1933						
	Bias	0.544		0.026			0.254	-0.172	0.022							
	Variance	0.311		0.453			0.142	0.189	0.479							
	Coverage	0.962		0.963			0.949	0.967	0.966							
	Width	2.315		2.704			1.765	2.07	2.775							
Scenario 3	MSE	1.295	1920	0.481	1920		0.509	0.217	0.506	1920						
	Bias	0.982		0.056			0.607	-0.127	0.048							
	Variance	0.33		0.478			0.141	0.201	0.504							
	Coverage	0.735		0.974			0.741	0.972	0.97							
	Width	2.388		2.754			1.763	2.093	2.841							
Scenario 4	MSE	1.944	1930	0.46	1929		0.879	0.232	0.49	1929						
	Bias	1.277		-0.041			0.865	-0.173	-0.036							
	Variance	0.313		0.458			0.131	0.202	0.489							
	Coverage	0.367		0.967			0.431	0.969	0.967							
	Width	2.359		2.733			1.736	2.105	2.822							

Table 21: Comparative performance of ML-TTE ($N'_{rep} < 2000$) and Bayes-NNDF ($N_{rep} = 2000$) on simulated datasets of size 960, $OR = 1.8$

		High prevalences			Low prevalences		
		Bayes-NNDF	ML-TTE	N'_{rep}	Bayes-NNDF	ML-TTE	N'_{rep}
Scenario 1	MSE	0.039	0.043	1963	0.184	0.231	599
	Bias	0.011	0.008		-0.13	-0.084	
	Variance	0.038	0.043		0.168	0.224	
	Coverage	0.964	0.946		0.973	0.977	
	Width	0.831	0.779		1.96	2.071	
Scenario 2	MSE	0.048	0.06	1960	0.194	0.439	641
	Bias	0.07	0.076		0.181	0.384	
	Variance	0.044	0.054		0.161	0.292	
	Coverage	0.949	0.924		0.975	0.947	
	Width	0.872	0.843		2.013	2.176	
Scenario 3	MSE	0.063	0.087	1961	0.313	0.807	702
	Bias	0.129	0.112		0.368	0.493	
	Variance	0.046	0.074		0.178	0.565	
	Coverage	0.942	0.905		0.928	0.738	
	Width	0.921	0.944		2.054	2.283	
Scenario 4	MSE	0.083	0.11	1960	0.482	0.997	758
	Bias	0.175	0.106		0.537	0.366	
	Variance	0.053	0.098		0.193	0.864	
	Coverage	0.911	0.877		0.859	0.656	
	Width	0.967	1.048		2.096	2.401	

Table 22: Comparative performance of ML-TTE ($N'_{rep} < 2000$) and Bayes-NNDF ($N_{rep} = 2000$) on simulated datasets of size 960, $OR = 2.5$

		High prevalences			Low prevalences		
		Bayes-NNDF	ML-TTE	N'_{rep}	Bayes-NNDF	ML-TTE	N'_{rep}
Scenario 1	MSE	0.041	0.044	1937	0.178	0.232	473
	Bias	0.006	0.004		-0.065	-0.044	
	Variance	0.041	0.044		0.174	0.231	
	Coverage	0.967	0.948		0.984	0.994	
	Width	0.856	0.791		2.089	2.238	
Scenario 2	MSE	0.053	0.069	1963	0.248	0.57	538
	Bias	0.085	0.087		0.263	0.517	
	Variance	0.046	0.061		0.179	0.303	
	Coverage	0.957	0.925		0.959	0.862	
	Width	0.908	0.869		2.143	2.325	
Scenario 3	MSE	0.079	0.104	1962	0.423	1.049	583
	Bias	0.16	0.133		0.471	0.645	
	Variance	0.054	0.087		0.201	0.634	
	Coverage	0.912	0.866		0.9	0.643	
	Width	0.956	0.971		2.197	2.441	
Scenario 4	MSE	0.114	0.135	1966	0.626	1.295	594
	Bias	0.245	0.135		0.647	0.528	
	Variance	0.054	0.117		0.207	1.019	
	Coverage	0.858	0.825		0.819	0.603	
	Width	1.003	1.086		2.253	2.511	

Table 23: Comparative performance of ML-TTE ($N'_{rep} < 2000$) and Bayes-NNDF ($N_{rep} = 2000$) on simulated datasets of size 960, $OR = 1.25$

		High prevalences			Low prevalences		
		Bayes-NNDF	ML-TTE	N'_{rep}	Bayes-NNDF	ML-TTE	N'_{rep}
Scenario 1	MSE	0.038	0.043	1950	0.167	0.2	677
	Bias	0.007	0.003		-0.139	-0.089	
	Variance	0.038	0.043		0.148	0.192	
	Coverage	0.966	0.949		0.97	0.96	
	Width	0.825	0.786		1.853	1.954	
Scenario 2	MSE	0.047	0.058	1955	0.158	0.34	736
	Bias	0.05	0.061	1953	0.097	0.261	
	Variance	0.045	0.054		0.149	0.273	
	Coverage	0.958	0.939		0.983	0.963	
	Width	0.868	0.851		1.889	2.063	
Scenario 3	MSE	0.053	0.073			0.262	0.648
	Bias	0.087	0.081	0.317		0.429	
	Variance	0.045	0.067	0.162		0.465	
	Coverage	0.953	0.935	0.942		0.84	
	Width	0.91	0.94	1.932		2.155	
Scenario 4	MSE	0.07	0.095	1968	0.347	0.8	839
	Bias	0.135	0.086		0.423	0.254	
	Variance	0.052	0.088		0.168	0.737	
	Coverage	0.932	0.91		0.904	0.731	
	Width	0.953	1.036		1.97	2.284	

Table 24: Comparative performance models on simulated datasets of size 500 ($N_{rep}=2000$) given high exposure prevalences and $OR = 1.8$

	ML		Bayes		QI	SIMEX-NDF				SIMEX-DF			
	NDF	DF	NDF	DF		Quadratic ASY.	JCK.	Loglinear ASY.	JCK.	Quadratic ASY.	JCK.	Loglinear ASY.	JCK.
Scenario 1	MSE	0.072	0.125	0.07	0.089	0.14	0.078	0.078	0.083	0.083	0.078	0.078	0.083
	Bias	0.009	0.014	0.011	-0.003	0.006	-0.024	-0.024	-0.004	-0.004	-0.024	-0.024	-0.004
	Variance	0.071	0.125	0.07	0.089	0.14	0.078	0.078	0.083	0.083	0.078	0.078	0.083
	Coverage	0.954	0.951	0.952	0.973	0.954	0.898	0.87	0.89	0.864	0.898	0.87	0.89
	Width	1.044	1.363	1.062	1.295	1.451	0.908	0.847	0.921	0.847	0.908	0.847	0.921
Scenario 2	MSE	0.095	0.141	0.087	0.101	0.158	0.102	0.102	0.131	0.131	0.102	0.102	0.131
	Bias	0.103	0.014	0.091	0.003	0.01	0.101	0.101	0.146	0.146	0.101	0.101	0.146
	Variance	0.085	0.141	0.078	0.101	0.158	0.092	0.092	0.11	0.11	0.092	0.092	0.11
	Coverage	0.949	0.942	0.941	0.964	0.947	0.884	0.841	0.856	0.795	0.884	0.841	0.856
	Width	1.115	1.423	1.11	1.33	1.533	0.969	0.883	1.006	0.883	0.969	0.883	1.006
Scenario 3	MSE	0.124	0.148	0.104	0.107	0.167	0.157	0.157	0.24	0.24	0.157	0.157	0.24
	Bias	0.185	0.006	0.158	0.001	0.003	0.233	0.233	0.313	0.313	0.233	0.233	0.313
	Variance	0.089	0.148	0.079	0.107	0.167	0.102	0.102	0.143	0.143	0.102	0.102	0.143
	Coverage	0.931	0.948	0.928	0.963	0.951	0.828	0.766	0.786	0.67	0.828	0.766	0.786
	Width	1.174	1.46	1.147	1.355	1.592	1.03	0.911	1.115	0.911	1.03	0.911	1.115
Scenario 4	MSE	0.174	0.157	0.14	0.115	0.184	0.251	0.251	0.453	0.453	0.251	0.251	0.453
	Bias	0.266	0.005	0.225	0.002	0.002	0.362	0.362	0.499	0.499	0.362	0.362	0.499
	Variance	0.103	0.157	0.089	0.115	0.184	0.12	0.12	0.204	0.204	0.12	0.12	0.204
	Coverage	0.892	0.942	0.896	0.957	0.949	0.711	0.628	0.63	0.51	0.711	0.628	0.63
	Width	1.233	1.495	1.189	1.384	1.66	1.09	0.937	1.228	0.937	1.09	0.937	1.228

Table 25: Comparative performance models on simulated datasets of size 500 ($N_{rep}=2000$) given high exposure prevalences and $OR = 1.25$

	ML NDF	ML DF	Bayes		QI	SIMEX-NDF			SIMEX-DF			
			NDF	DF		Quadratic ASY.	JCK.	Loglinear ASY.	JCK.	Quadratic ASY.	JCK.	Loglinear ASY.
Scenario 1	MSE	0.073	0.138	0.074	0.099	0.152						
	Bias	0.006	0.009	0.007	0.001	0.001						
	Variance	0.073	0.138	0.074	0.099	0.152						
	Coverage	0.956	0.949	0.954	0.972	0.945						
	Width	1.067	1.41	1.099	1.345	1.501						
Scenario 2	MSE	0.102	0.145	0.097	0.106	0.163						
	Bias	0.123	0.009	0.116	0.018	0.005						
	Variance	0.087	0.145	0.083	0.106	0.163						
	Coverage	0.94	0.958	0.932	0.968	0.959						
	Width	1.139	1.479	1.146	1.386	1.588						
Scenario 3	MSE	0.146	0.16	0.131	0.116	0.185						
	Bias	0.231	-0.003	0.214	0.02	-0.005						
	Variance	0.093	0.16	0.085	0.116	0.185						
	Coverage	0.899	0.949	0.893	0.96	0.947						
	Width	1.196	1.518	1.182	1.41	1.652						
Scenario 4	MSE	0.217	0.169	0.186	0.126	0.197						
	Bias	0.329	-0.008	0.302	0.02	-0.015						
	Variance	0.109	0.169	0.094	0.126	0.197						
	Coverage	0.824	0.956	0.826	0.959	0.956						
	Width	1.257	1.551	1.222	1.438	1.715						

Table 26: Comparative performance models on simulated datasets of size 500 ($N_{rep}=2000$) given high exposure prevalences and $OR = 2.5$

	ML		Bayes		QI	SIMEX-NDF			SIMEX-DF		
	NDF	ML DF	NDF	Bayes DF		Quadratic ASY.	Loglinear ASY.	JCK.	Quadratic ASY.	Loglinear ASY.	JCK.
Scenario 1	MSE	0.077	0.131	0.072	0.092	0.145					
	Bias	0.021	0.017	0.019	-0.012	0.016					
	Variance	0.077	0.13	0.071	0.092	0.145					
	Coverage	0.95	0.943	0.949	0.962	0.941					
Width	1.052	1.344	1.055	1.268	1.43						
Scenario 2	MSE	0.087	0.136	0.075	0.095	0.151					
	Bias	0.084	0.016	0.065	-0.011	0.015					
	Variance	0.08	0.136	0.071	0.095	0.151					
	Coverage	0.956	0.95	0.952	0.964	0.945					
Width	1.112	1.392	1.095	1.297	1.495						
Scenario 3	MSE	0.118	0.148	0.095	0.106	0.174					
	Bias	0.152	0.015	0.113	-0.014	0.005					
	Variance	0.095	0.147	0.082	0.106	0.174					
	Coverage	0.94	0.945	0.943	0.959	0.942					
Width	1.182	1.44	1.142	1.33	1.573						
Scenario 4	MSE	0.152	0.154	0.113	0.111	0.19					
	Bias	0.222	0.019	0.165	-0.011	0.012					
	Variance	0.103	0.154	0.085	0.111	0.19					
	Coverage	0.919	0.943	0.923	0.957	0.938					
Width	1.24	1.471	1.18	1.355	1.635						

Table 27: Comparative performance of ML, QI ($N'_{rep} < 2000$) and Bayes, SIMEX ($N_{rep} = 2000$) models on simulated datasets of size 500, given low exposure prevalences and $OR = 1.8$

	ML			ML			Bayes			SIMEX-NDF			SIMEX-DF		
	NDF	N'_{rep}	DF	DF	N'_{rep}	NDF	DF	QI	N'_{rep}	Quadratic ASY.	Loglinear JCK.	Loglinear JCK.	Quadratic ASY.	Loglinear JCK.	Loglinear JCK.
Scenario 1	MSE	0.429	1695	0.631	1679	0.257	0.277	0.666	1679						
	Bias	-0.07		-0.071		-0.191	-0.226	-0.071							
	Variance	0.424		0.626		0.22	0.226	0.661							
	Coverage	0.975		0.978		0.971	0.983	0.979							
	Width	2.978		3.47		2.268	2.5	3.582							
Scenario 2	MSE	0.588	1726	0.645	1719	0.222	0.255	0.699	1719						
	Bias	0.406		-0.124		0.151	-0.176	-0.137							
	Variance	0.424		0.63		0.2	0.224	0.68							
	Coverage	0.987		0.977		0.981	0.985	0.977							
	Width	3.108		3.549		2.275	2.521	3.666							
Scenario 3	MSE	1.136	1705	0.628	1690	0.418	0.255	0.671	1690						
	Bias	0.846		-0.131		0.464	-0.14	-0.137							
	Variance	0.421		0.612		0.203	0.235	0.652							
	Coverage	0.918		0.978		0.91	0.986	0.978							
	Width	3.105		3.554		2.269	2.545	3.683							
Scenario 4	MSE	1.87	1751	0.646	1724	0.706	0.246	0.68	1724						
	Bias	1.201		-0.115		0.721	-0.134	-0.128							
	Variance	0.427		0.633		0.187	0.228	0.664							
	Coverage	0.726		0.987		0.776	0.988	0.982							
	Width	3.117		3.594		2.244	2.56	3.731							

Table 28: Comparative performance of ML, QI ($N'_{rep} < 2000$) and Bayes, SIMEX ($N_{rep} = 2000$) models on simulated datasets of size 500, given low exposure prevalences and $OR = 1.25$

	ML			ML			Bayes			SIMEX-NDF			SIMEX-DF		
	NDF	N'_{rep}	DF	NDF	N'_{rep}	DF	NDF	DF	DF	Quadratic ASY. JCK. ASY. JCK.	Loglinear ASY. JCK. ASY. JCK.	Quadratic ASY. JCK. ASY. JCK.	Loglinear ASY. JCK. ASY. JCK.		
Scenario 1	MSE	0.47	1616	0.646	1597	0.23	0.23	0.23	0.23	0.676	1597				
	Bias	-0.032		-0.032		-0.091	-0.1	-0.019		-0.019					
	Variance	0.47		0.645		0.221	0.22	0.676		0.676					
	Coverage	0.991		0.989		0.983	0.989	0.986		0.986					
	Width	3.256		3.732		2.388	2.608	3.854		3.854					
Scenario 2	MSE	0.732	1603	0.641	1585	0.306	0.227	0.681	1585						
	Bias	0.514		-0.061		0.302	-0.026	-0.065		-0.065					
	Variance	0.469		0.638		0.215	0.227	0.677		0.677					
	Coverage	0.971		0.988		0.96	0.993	0.987		0.987					
	Width	3.373		3.802		2.405	2.637	3.92		3.92					
Scenario 3	MSE	1.44	1653	0.685	1616	0.595	0.238	0.742	1616						
	Bias	0.965		-0.091		0.61	-0.013	-0.1		-0.1					
	Variance	0.51		0.677		0.223	0.238	0.733		0.733					
	Coverage	0.843		0.99		0.857	0.991	0.986		0.986					
	Width	3.364		3.839		2.397	2.658	3.963		3.963					
Scenario 4	MSE	2.371	1668	0.621	1608	1.038	0.227	0.674	1608						
	Bias	1.364		-0.099		0.905	0.006	-0.105		-0.105					
	Variance	0.511		0.612		0.219	0.227	0.663		0.663					
	Coverage	0.576		0.993		0.679	0.995	0.99		0.99					
	Width	3.248		3.835		2.374	2.675	3.971		3.971					

Table 29: Comparative performance of ML, QI ($N'_{rep} < 2000$) and Bayes, SIMEX ($N_{rep} = 2000$) models on simulated datasets of size 500, given low exposure prevalences and $OR = 2.5$

	ML			ML			Bayes			SIMEX-NDF			SIMEX-DF							
	NDF	N'_{rep}	DF	NDF	N'_{rep}	DF	NDF	DF	QI	N'_{rep}	Quadratic		Loglinear		Quadratic		Loglinear			
											ASY.	JCK.	ASY.	JCK.	ASY.	JCK.	ASY.	JCK.	ASY.	JCK.
Scenario 1	MSE	0.388	1768	0.566	1752	0.253	0.301	0.589	1752	0.023	0.023	0.024	0.024	0.024	0.024	0.023	0.023	0.024	0.024	
	Bias	-0.045		-0.076		-0.237	-0.301	-0.087		-0.005	-0.005	0.004	0.004	0.004	0.004	-0.005	-0.005	0.004	0.004	
	Variance	0.387		0.561		0.197	0.211	0.582		0.023	0.023	0.024	0.024	0.024	0.023	0.023	0.024	0.024	0.024	
	Coverage	0.961		0.969		0.958	0.962	0.968		0.895	0.871	0.856	0.86	0.86	0.895	0.871	0.856	0.86	0.86	0.86
	Width	2.901		3.275		2.145	2.386	3.381		0.497	0.463	0.498	0.463	0.463	0.497	0.463	0.498	0.498	0.463	0.463
Scenario 2	MSE	0.538	1732	0.586	1729	0.2	0.287	0.631	1729	0.049	0.049	0.059	0.059	0.059	0.049	0.049	0.059	0.059	0.059	
	Bias	0.391		-0.104		0.081	-0.248	-0.105		0.151	0.151	0.171	0.171	0.171	0.151	0.151	0.171	0.171	0.171	
	Variance	0.385		0.575		0.193	0.225	0.62		0.027	0.027	0.03	0.03	0.03	0.027	0.027	0.03	0.03	0.03	
	Coverage	0.99		0.965		0.979	0.972	0.966		0.758	0.693	0.717	0.648	0.648	0.758	0.693	0.717	0.648	0.648	0.648
	Width	2.977		3.34		2.167	2.423	3.456		0.525	0.478	0.544	0.478	0.478	0.525	0.478	0.544	0.478	0.478	0.478
Scenario 3	MSE	0.99	1727	0.612	1723	0.324	0.285	0.65	1723	0.133	0.133	0.174	0.174	0.174	0.133	0.133	0.174	0.174	0.174	
	Bias	0.763		-0.102		0.358	-0.22	-0.113		0.32	0.32	0.366	0.366	0.366	0.32	0.32	0.366	0.366	0.366	
	Variance	0.407		0.602		0.196	0.237	0.638		0.031	0.031	0.04	0.04	0.04	0.031	0.031	0.04	0.04	0.04	
	Coverage	0.951		0.97		0.943	0.972	0.973		0.419	0.35	0.378	0.277	0.277	0.419	0.35	0.378	0.277	0.277	
	Width	2.986		3.375		2.159	2.447	3.505		0.557	0.494	0.599	0.494	0.494	0.557	0.494	0.599	0.494	0.494	
Scenario 4	MSE	1.512	1735	0.61	1735	0.523	0.275	0.654	1735	0.295	0.295	0.418	0.418	0.418	0.295	0.295	0.418	0.418	0.418	
	Bias	1.059		-0.114		0.584	-0.21	-0.114		0.508	0.508	0.599	0.599	0.599	0.508	0.508	0.599	0.599	0.599	
	Variance	0.391		0.597		0.182	0.231	0.641		0.037	0.037	0.059	0.059	0.059	0.037	0.037	0.059	0.059	0.059	
	Coverage	0.829		0.973		0.833	0.974	0.969		0.121	0.084	0.095	0.052	0.052	0.121	0.084	0.095	0.052	0.052	
	Width	2.982		3.375		2.14	2.456	3.513		0.587	0.506	0.661	0.506	0.506	0.587	0.506	0.661	0.506	0.506	

Table 30: Comparative performance of ML-TTE ($N'_{rep} < 2000$) and Bayes-NNDF ($N_{rep} = 2000$) on simulated datasets of size 500, $OR = 1.8$

		High prevalences			Low prevalences		
		Bayes-NNDF	ML-TTE	N'_{rep}	Bayes-NNDF	ML-TTE	N'_{rep}
Scenario 1	MSE	0.07	0.075	1696	0.252	0.318	252
	Bias	0.008	0.008	-0.201	-0.203		
	Variance	0.07	0.074	0.212	0.278		
	Coverage	0.963	0.95	0.977	0.964		
	Width	1.102	1.062	2.328	2.518		
Scenario 2	MSE	0.084	0.101	1785	0.2	0.308	291
	Bias	0.073	0.084	0.057	0.195		
	Variance	0.079	0.094	0.197	0.271		
	Coverage	0.952	0.947	0.99	0.986		
	Width	1.153	1.139	2.357	2.617		
Scenario 3	MSE	0.095	0.125	1799	0.285	0.62	320
	Bias	0.123	0.129	0.273	0.463		
	Variance	0.08	0.108	0.211	0.406		
	Coverage	0.946	0.932	0.971	0.928		
	Width	1.193	1.219	2.393	2.694		
Scenario 4	MSE	0.12	0.161	1776	0.382	0.829	332
	Bias	0.172	0.137	0.42	0.443		
	Variance	0.09	0.142	0.205	0.634		
	Coverage	0.933	0.913	0.942	0.849		
	Width	1.24	1.316	2.426	2.688		

Table 31: Comparative performance of ML-TTE ($N'_{rep} < 2000$) and Bayes-NNDF ($N_{rep} = 2000$) on simulated datasets of size 500, $OR = 2.5$

		High prevalences			Low prevalences		
		Bayes-NNDF	ML-TTE	N'_{rep}	Bayes-NNDF	ML-TTE	N'_{rep}
Scenario 1	MSE	0.074	0.074	1664	0.219	0.284	186
	Bias	0.005	0.011		-0.093	-0.156	
	Variance	0.074	0.074		0.211	0.261	
	Coverage	0.962	0.96		0.988	0.978	
	Width	1.143	1.087		2.446	2.647	
Scenario 2	MSE	0.092	0.105	1770	0.25	0.401	201
	Bias	0.095	0.101		0.204	0.341	
	Variance	0.083	0.095		0.208	0.286	
	Coverage	0.952	0.941		0.985	0.965	
	Width	1.197	1.166		2.486	2.708	
Scenario 3	MSE	0.116	0.145	1785	0.386	0.864	246
	Bias	0.169	0.161		0.403	0.601	
	Variance	0.087	0.119		0.224	0.505	
	Coverage	0.93	0.909		0.948	0.858	
	Width	1.238	1.245		2.521	2.743	
Scenario 4	MSE	0.151	0.198	1777	0.552	1.16	255
	Bias	0.231	0.165		0.572	0.648	
	Variance	0.098	0.171		0.225	0.743	
	Coverage	0.9	0.876		0.898	0.718	
	Width	1.284	1.357		2.565	2.898	

Table 32: Comparative performance of ML-TTE ($N'_{rep} < 2000$) and Bayes-NNDF ($N_{rep} = 2000$) on simulated datasets of size 500, $OR = 1.25$

		High prevalences			Low prevalences		
		Bayes-NNDF	ML-TTE	N'_{rep}	Bayes-NNDF	ML-TTE	N'_{rep}
Scenario 1	MSE	0.072	0.079	1673	0.256	0.285	302
	Bias	0.013	0.01		-0.254	-0.185	
	Variance	0.072	0.079		0.192	0.252	
	Coverage	0.957	0.95		0.959	0.947	
	Width	1.09	1.067		2.204	2.409	
Scenario 2	MSE	0.074	0.09	1754	0.195	0.279	356
	Bias	0.05	0.069		-0.01	0.105	
	Variance	0.072	0.085		0.195	0.269	
	Coverage	0.959	0.956		0.987	0.992	
	Width	1.131	1.13		2.248	2.444	
Scenario 3	MSE	0.09	0.127	1810	0.241	0.495	411
	Bias	0.086	0.101		0.186	0.368	
	Variance	0.083	0.117		0.207	0.36	
	Coverage	0.958	0.936		0.973	0.966	
	Width	1.18	1.221		2.275	2.53	
Scenario 4	MSE	0.102	0.148	1797	0.308	0.69	407
	Bias	0.125	0.115		0.325	0.29	
	Variance	0.086	0.135		0.203	0.607	
	Coverage	0.946	0.926		0.96	0.921	
	Width	1.222	1.312		2.3	2.609	