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MEASUREMENT ERROR AND CORRECTIONS FOR
ATTENUATION IN META-ANALYSES

by

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Institute of Statistics Mimeograph Series No. 2222

May 1992

NORTH CAROLINA STATE UNIVERSITY
Raleigh, North Carolina

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META-ANALYSES**

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ABSTRACT

MacMahon, et al. (1990) present a meta-analysis of the effect of blood pressure on coronary heart disease, as well as new methods for estimation in measurement error models for the case when a replicate or second measurement is made of the fallible predictor. The correction for attenuation used by these authors is compared to others already existing in the literature, as well as to a new instrumental variable method. The assumptions justifying the various methods are examined and their efficiencies are studied via simulation. Compared to the methods we discuss, that of MacMahon, et al. (1990) may have substantial bias in some circumstances because it does not take into account: (i) possible correlations among the predictors within a study; (ii) possible bias in the second measurement; or (iii) possibly differing marginal distributions of the predictors and/or measurement errors across studies.

1. INTRODUCTION

In this paper we discuss some problems that arise in regression meta-analysis when the predictor of interest is measured with nonnegligible error. Motivating our study is a recent paper by MacMahon, et al. (1990) presenting a meta-analysis of the effect of blood pressure on stroke and coronary heart disease. The methodology employed by MacMahon, et al. (1990) has been publicized in the press (Palca, 1990a,b), and it is likely to promote the use of measurement-error-model meta-analysis in future studies.

Underlying the method proposed by MacMahon, et al. (1990) is a correction for attenuation due to measurement error that employs both baseline and post-baseline measurements of blood pressure from one of the studies; see also Ederer (1972). We compare their correction for attenuation to some of the standard methods in the statistical literature for adjusting regression coefficients for the effects of measurement error. The assumptions justifying the various methods are examined in light of the type of violations that are likely to arise in a meta-analysis, and the estimated corrections for attenuation are studied via a small simulation experiment.

A necessary feature of a meta-analysis is that the individual studies share sufficient common ground to warrant the combination of information from the different data sets. With respect to measurement error modelling this imposes some restrictions on the relationship between the so-called true predictors and the surrogate predictors actually measured, especially when replicate measurements and/or validation data are limited. Implicit in the MacMahon, et al. (1990) method is the assumption that a single correction for attenuation is adequate for all studies in the meta-analysis. We show that this is not appropriate in general and describe the estimation of study-specific corrections.

In § 2 we establish notation and discuss the effects of measurement error on regression meta-analyses. In § 3 we present and compare some corrections for attenuation due to measurement error. In § 4 we discuss the method employed by MacMahon, et al. (1990) in light of the results from § 2-3. We conclude in § 5.

2. GENERAL THEORY

2.1. Study Data, Models and Meta-Analysis in the Absence of Measurement Error

Let Y denote the response variate and X the risk factor that is to be investigated in the meta-analysis. We suppose the availability of data from \mathcal{K} studies, with Y and X common to all studies. In addition, data from the k^{th} study includes a vector Z_k of study-specific covariates.

Justification for the meta-analysis depends on the assumption that the effect of X on Y is the same in all study populations after appropriate covariate adjustments. Thus as a model for the expected value of Y in the k^{th} population we have

$$E(Y | X, Z_k) = f(\alpha_k + \beta X + \gamma_k^T Z_k), \quad k = 1, \dots, \mathcal{K}. \quad (2.1)$$

The form of f is application dependent. For example, for linear regression f is the identity function, for binary regression f is commonly the logistic distribution function.

In some models, *e.g.*, logistic regression, (2.1) completely specifies the model once the type of variation, (Bernoulli, exponential, Poisson, *etc.*) in Y is specified. In other models there are additional variance parameters to model the residual variation which may be homoscedastic in the simplest cases or involve the mean function in addition to other variance parameters. Thus we allow for the possibility that accompanying (2.1) there is a variance-function model of the form

$$V(Y | X, Z_k) = \sigma_k^2 v(\alpha_k + \beta X + \gamma_k^T Z_k, \tau_k), \quad k = 1, \dots, \mathcal{K}. \quad (2.2)$$

Together (2.1) and (2.2) comprise a generalized quasiliikelihood/variance-function model with linear effects, and includes, for example, generalized linear models and heteroscedastic nonlinear models; see Carroll & Ruppert (1988) for details.

Let $\hat{\beta}_k$, $k = 1, \dots, \mathcal{K}$ denote the estimates of β obtained by fitting the model to the k^{th} study data. The combined-study estimate of β is obtained as a weighted average

$$\hat{\beta} = \sum_{k=1}^{\mathcal{K}} \hat{w}_k \hat{\beta}_k, \quad (\hat{w}_1 + \dots + \hat{w}_k = 1).$$

In the absence of special considerations, the natural choice of weights has $\hat{w}_k \propto s_k^{-2}$ where s_k is the standard error of $\hat{\beta}_k$.

2.2. The Effect of Measurement Error

Now suppose that the risk factor X is not accurately measured. Let W denote the measured risk factor and assume the representation $W = X + U$ where U is a measurement error independent of Y , X and the other covariates. Furthermore assume that in the k^{th} study $\text{Var}(U) = \sigma_U^2$, independent of k .

In this paper we focus primarily on the simple additive measurement error model just described, although we point out some problems that may arise when the model is inadequate. The acceptability of the model for the epidemiologic application motivating the present paper is difficult to assess with the data available to us. As a referee noted, it fails to account for improvements and standardization in blood pressure measurement methods that have occurred since the start of the Framingham study. Thus a more realistic model might have heteroscedastic measurement errors with variances decreasing over time. We discuss this problem later in the paper (§ 4.3) in an example.

When the simple additive error model is known to be inadequate it may be necessary to allow for more complex dependencies between the proxy, W , and X and possibly the other covariates as well; see Fuller (1987) and Carroll & Stefanski (1990).

Now suppose that the models (2.1) and (2.2) are fit to the data with W substituted for X . The study-specific and combined-study estimators so obtained are denoted $\tilde{\beta}_k$, $k = 1, \dots, \mathcal{K}$ and $\tilde{\beta}$ respectively. In general neither $\tilde{\beta}_k$, $k = 1, \dots, \mathcal{K}$ nor $\tilde{\beta}$ are consistent for β ; see Stefanski (1985).

In the case that (2.1) and (2.2) define common homoscedastic linear models it is well known that $\tilde{\beta}_k$ consistently estimates $\lambda_k^{-1}\beta$ where

$$\lambda_k = \frac{\sigma_{W,k}^2}{\sigma_{W,k}^2 - \sigma_U^2}, \quad (2.3)$$

and $\sigma_{W,K}^2$ is the limiting value, assumed to exist, of $s_{W,k}^2 =$ the mean square error from the linear regression of W on Z_k in the k^{th} study data. Since $\sigma_{W,k}^2 = \sigma_{X,k}^2 + \sigma_U^2$ where $\sigma_{X,k}^2$ is the residual variation from the linear regression of X on Z_k , the attenuation factor for the k^{th} study is

$$\lambda_k^{-1} = \frac{\sigma_{X,k}^2}{\sigma_{X,k}^2 + \sigma_U^2}. \quad (2.4)$$

It is evident from (2.4) that attenuation generally varies across study groups, and that collinearity between X and Z_k accentuates attenuation due to measurement error. Note that $\sigma_{W,k}^2 = \text{Var}(W)$ and $\sigma_{X,k}^2 = \text{Var}(X)$ only in the absence of covariates or when X and Z_k are uncorrelated.

It follows that for linear models the combined-study estimator $\tilde{\beta}$ converges in probability to $\lambda^{-1}\beta$ where

$$\lambda^{-1} = \sum_{k=1}^{\mathcal{K}} w_k \lambda_k^{-1},$$

where w_k are the limiting values of weights \tilde{w}_k .

Assuming that data are available for consistent estimation of λ_k , by say $\tilde{\lambda}_k$, $k = 1, \dots, \mathcal{K}$, study-specific estimators of β corrected for attenuation are obtained as $\tilde{\beta}_{k,CA} = \tilde{\lambda}_k \tilde{\beta}_k$, $k = 1, \dots, \mathcal{K}$. A combined-study estimator can be obtained by either taking a weighted average of $\{\tilde{\beta}_{k,CA}\}$ or as $\tilde{\lambda} \tilde{\beta}$ where $\tilde{\lambda}$ is a consistent estimator of λ . The latter estimator is just a particular weighted average of $\{\tilde{\beta}_{k,CA}\}$.

The best way to combine the study-specific estimators $\tilde{\lambda}_k$ and $\tilde{\beta}_k$ to produce an efficient and consistent estimator of β will depend of the specific nature of the $\tilde{\lambda}_k$ s, *i.e.*, exactly how these corrections are obtained. This is an important problem but one that we will not discuss in detail in this paper.

Starting with (2.1) and (2.2) and the simple measurement error model $W = X + U$ the following approximations to $E(Y|W)$ and $\text{Var}(Y|W)$ can be derived:

$$E(Y|W, Z_k) \approx f\{\alpha_k + \beta E(X | W, Z_k) + \gamma_k^T Z_k\} \approx f(\alpha_k^* + \lambda_k^{-1} \beta W + \gamma_k^{*T} Z_k); \quad (2.5)$$

$$V(Y|W, Z_k) \approx \sigma_k^2 v\{\alpha_k + \beta E(X | W, Z_k) + \gamma_k^T Z_k, \tau_k\} \approx \sigma_k^2 v(\alpha_k^* + \lambda_k^{-1} \beta W + \gamma_k^{*T} Z_k, \tau_k). \quad (2.6)$$

The first approximation in (2.5) follows via a Taylor series expansion of (2.1) in X around $E(X | W, Z_k)$. The second approximation is obtained by replacing $E(X | W, Z_k)$ with the best linear approximation to this regression function. The approximations in (2.6) are similarly obtained. Estimation based on substituting $E(X | W, Z_k)$ for X is discussed by Carroll & Stefanski (1990), Gleser (1990), Fuller (1987), Pierce, et al. (1991), Prentice (1982), Rosner, et al. (1989, 1990) and Rudemo, et al. (1989), among others.

Equations (2.5) and (2.6) show that if the models (2.1) and (2.2) are fit to the data with W substituted for X , then the asymptotic bias in $\tilde{\beta}_k$ will be approximately the same as if linear models had been fit to the data. That is, $\tilde{\beta}_k$ converges in probability to some value, say β_k^\dagger , that is approximately equal to $\lambda_k^{-1}\beta$. It follows that whenever the approximations in (2.5) and (2.6) are justified, the k^{th} study estimator can be corrected for attenuation just as in the case of linear models, *viz.*, $\tilde{\beta}_{k,CA} = \tilde{\lambda}_k\tilde{\beta}_k$. Similarly a combined-study estimator corrected for attenuation is obtained as a weighted average of $\tilde{\beta}_{k,CA}$, $k = 1, \dots, \mathcal{K}$.

When the variation in $\tilde{\lambda}_k$ is small relative to that in $\tilde{\beta}_k$, $k = 1, \dots, \mathcal{K}$, $\text{Var}(\tilde{\beta}) \approx \sum_{k=1}^{\mathcal{K}} \tilde{w}_k^2 \tilde{\lambda}_k^2 \tilde{s}_k^2$ where \tilde{s}_k is the usual standard error of $\tilde{\beta}_k$. In this case the best choice of weights has approximately $\tilde{w}_k \propto (\tilde{\lambda}_k \tilde{s}_k)^{-2}$ for which the standard error of $\tilde{\beta}$ is approximately $\{\sum_{k=1}^{\mathcal{K}} (\tilde{\lambda}_k \tilde{s}_k)^{-2}\}^{-1/2}$. If the variation in $\tilde{\lambda}_k$ is not negligible compared to that of $\tilde{\beta}_k$, $k = 1, \dots, \mathcal{K}$, then calculation of $\text{Var}(\tilde{\beta})$ is more difficult and will depend on the particular form of the estimators.

3. ESTIMATING CORRECTIONS FOR ATTENUATION

3.1. Introduction

The composition of the data, especially with regard to replicate measurements, validation data and the presence of instrumental variables, generally dictates the appropriate method of estimating λ_k , $k = 1, \dots, \mathcal{K}$. We will discuss only methods that are applicable to the epidemiologic application motivating the paper, (MacMahon *et al.*, 1990).

In that application all studies measured diastolic blood pressure (DBP) at baseline; this is W in the established notation. In addition, one study, Framingham, measured DBP at two-years and four-years post baseline. As in MacMahon, et al. (1990) we will use only the four-year post-baseline measurement. So now in addition to having available (Y, W, Z_k) in all k studies, there is available in one study, taken as the first, another variate, call it T . In the application under consideration T is the post-baseline measurement of DBP; more generally we regard T as a *second measurement* of X .

We make a distinction between a *second measurement* and a *replicate measurement*. The former implies only that T and X are correlated. The latter embodies the usual statistical notion of replicates; W and T are replicate measurements of X when $W = X + U_1$, $T = X + U_2$ and U_1

and U_2 are independent and identically distributed. The distinction is useful for it dictates when T should be employed as an instrumental variable. See Fuller (1987, p. 52) for a completely general definition of an instrumental variable. We note here that the essential requirements are that: i) T is correlated with X ; ii) T is uncorrelated with $W - X$; and iii) T is uncorrelated with $Y - \text{pr}(Y = 1 | X)$. The importance and use of instrumental variables will be evident in § 3.3 .

The data from the first study contain most of the information for estimating σ_U^2 . These data are used either to provide a direct estimate of σ_U^2 or an indirect estimate by first estimating λ_1 directly and then obtaining an estimate of σ_U^2 via (2.3), viz.,

$$\hat{\sigma}_U^2 = \frac{\hat{\lambda}_1 - 1}{\hat{\lambda}_1} s_{W,1}^2. \quad (3.1)$$

In either case λ_k , $k = 2, \dots, K$ are estimated by

$$\tilde{\lambda}_k = \frac{s_{W,k}^2}{s_{W,k}^2 - \hat{\sigma}_U^2}.$$

3.2. When W and T are Replicates

For the model with $W = X + U_1$ and $T = X + U_2$ where U_1 and U_2 are independent, identically distributed and independent of Z_1 , the coefficient of W in the linear regression of T on (W, Z_1) , denoted $\beta_{T|W,Z_1}$, is equal to λ_1^{-1} . Thus an estimator of λ_1 is

$$\bar{\lambda}_1 = \frac{1}{\hat{\beta}_{T|W,Z_1}}. \quad (3.2)$$

Although statistical justification for this estimator is not strong, it has much intuitive appeal. In particular, it makes clear the connection between attenuation due to measurement error and the more widely understood phenomenon of regression to the mean. Apart from differences due to grouping and the inclusion of covariates, $\bar{\lambda}_1$ corresponds to the estimator employed by MacMahon, et al. (1990), see § 4 for additional discussion.

Objections to this estimator arise because it is not symmetric in T and W , while the statistical model is. Under the replication model is it natural to estimate σ_U^2 by

$$\hat{\sigma}_U^2 = (1/2) \text{ sample variance of the } (W_i - T_i), \quad (3.3)$$

$\sigma_{W,k}^2$ by $s_{W,k}^2 =$ the mean square error from the linear regression of W_k on Z_k , and then λ_k by

$$\tilde{\lambda}_k = \frac{s_{W,k}^2}{s_{W,k}^2 - \hat{\sigma}_U^2}, \quad k = 1, \dots, \mathcal{K}. \quad (3.4)$$

The variance component estimate in (3.3) differs from the usual ANOVA within-subjects variance component estimate, $(2n)^{-1} \sum_{i=1}^n (W_i - T_i)^2$. Both are unbiased and they are asymptotically equivalent, and thus equally efficient in large samples. Our preference for (3.3) over the usual ANOVA variance components estimate is based on model robustness considerations. Departures from the replicate-measurements model in the direction of the instrumental variable model (3.3) have a greater effect on the ANOVA variance component estimate than they do on (3.3).

Even the latter procedure is not completely satisfying, for in the replication model the best measurement of X in the first study is $W^* = (T + W)/2$ and logic dictates first regressing Y_1 on W^* and Z_1 obtaining $\hat{\beta}_1^*$ with corresponding attenuation factor

$$\lambda_1^{*-1} = \frac{\sigma_{X,1}^2}{\sigma_{X,1}^2 + \sigma_U^2/2}.$$

The measurement error variance, σ_U^2 , again is estimated as in (3.3), and the mean square error from the regression of W^* on Z_1 , denote it $s_{W,1}^{*2}$, consistently estimates $\sigma_{X,1}^2 + \sigma_U^2/2$. Thus

$$\tilde{\lambda}_1^* = \frac{s_{W,1}^{*2}}{s_{W,1}^{*2} - \hat{\sigma}_U^2/2} \quad (3.5)$$

is a consistent estimator of λ_1^* .

The latter procedure makes more efficient use of the data but may be objected to on the grounds that it treats the first study differently from the rest. Greater similarity among studies, in this case having the study-specific analyses depend on baseline data only, makes it easier to present and defend the meta-analysis via nontechnical arguments.

Treating baseline and post-baseline measurements equally may be objectionable on statistical grounds as well. It would not be surprising to find differences in cohort distributions of DBP measurements taken four years apart, especially among study participants. In general, when the assumption that W and T are replicates is untenable, T should be employed as an instrumental variable.

3.3. T as an Instrumental Variable

Approximate instrumental variable estimation in generalized quasiliikelihood/variance-function models will be studied in detail elsewhere. Following is a brief outline of that theory and a discussion of its application in meta-analysis of measurement error models.

If (2.1) and (2.2) are expanded around $E(X | T, Z_k)$ instead of $E(X | W, Z_k)$, then analogous to (2.5) and (2.6) we obtain, after replacing $E(X | T, Z_k)$ with its best linear approximant, the approximations

$$E(Y|T, Z_k) \approx f\{\alpha_k + \beta E(X | T, Z_k) + \gamma_k^T Z_k\} \approx f(\alpha_k^* + \delta_1 \beta T + \gamma_k^{*T} Z_k);$$

$$V(Y|T, Z_k) \approx \sigma_k^2 v\{\alpha_k + \beta E(X | T, Z_k) + \gamma_k^T Z_k, \tau_k\} \approx \sigma_k^2 v(\alpha_k^* + \delta_1 \beta T + \gamma_k^{*T} Z_k, \tau_k),$$

where δ_1 is the coefficient of T in the best linear approximation to the regression of X on T and Z_1 .

Thus if the model (2.1) and (2.2) is fit to the data with T replacing X , the estimated coefficient of T , denoted $\hat{\beta}_{Y|T, Z_1}$, is approximately consistent for $\delta_1 \beta$. In other words, $\hat{\beta}_{Y|T, Z_1}$ converges in probability to some constant that is approximately equal to $\delta_1 \beta$.

Now for the additive model $W = X + U_1$, it is easy to establish that δ_1 is also the coefficient of T in the best linear approximation to the regression of W on T and Z_1 , and thus can be consistently estimated by $\hat{\beta}_{W|T, Z_1}$ = the estimated coefficient of T in the least squares regression of W on T and Z_1 .

This leads to the approximate instrumental variable estimator

$$\tilde{\beta}_{1,IV} = \frac{\hat{\beta}_{Y|T, Z_1}}{\hat{\beta}_{W|T, Z_1}},$$

from which is derived the estimator of λ_1

$$\tilde{\lambda}_{1,IV} = \frac{\tilde{\beta}_{1,IV}}{\tilde{\beta}_1}. \quad (3.6)$$

3.4. Comparing the Estimators of Attenuation

The four estimators $\bar{\lambda}_1$, $\tilde{\lambda}_1$, $\tilde{\lambda}_1^*$ and $\tilde{\lambda}_{1,IV}$, are all consistent for λ_1 when W and T are replicate measurements of X . We now examine the effect of departures from the replicate measurement error model on the four estimators.

We assume that $W = X + U_1$ as before, but that

$$T = \xi + \eta X + U_2^* \quad (3.7)$$

where U_2^* is a random error independent of U_1 and Z_1 , but not necessarily having the same distribution as U_1 . The model for T can be motivated as follows.

Let X and X_* denote the ‘true’ DBP at baseline and four years post-baseline respectively, of a randomly selected patient. If there is no change in true DBPs over the study period then $X = X_*$. If there is change, then it is reasonable to assume that X and X_* are jointly normal, in which case the regression of X_* on X is linear. Provided T is an unbiased measurement of the true DBP at follow-up, then $T = E(X_* | X) + U_2^*$ where $E(X_* | X) = \xi + \eta X$ and U_2^* is the sum of the measurement error and the residual error $X_* - E(X_* | X)$.

For this model $\eta = \rho \sigma_{X_*} \sigma_X^{-1}$, where $\rho = \text{corr}(X, X_*)$. Note that $\eta < 1$ unless $\text{Var}(X_*) \geq \rho^{-2} \text{Var}(X)$. So unless the variation in true DBPs increases over time, we expect $\eta < 1$.

Now consider $\bar{\lambda}_1$ defined in (3.2). For the model described above, $\beta_{T|W, Z_1}$ = the coefficient of W in the linear regression of T on (W, Z_1) , is $\eta \lambda_1^{-1}$, and thus $\bar{\lambda}_1$ is a consistent estimator of λ_1 / η . The correction for attenuation is overestimated in the common situation that $\eta < 1$ and underestimated when $\eta > 1$.

The second correction for attenuation, $\tilde{\lambda}_1$ given in (3.4) depends on the post-baseline measurements only via (3.3). If the T_i in (3.3) follow the model in (3.7) then $\hat{\sigma}_T^2$ is a consistent estimator of

$$\frac{1}{2} \left\{ \sigma_{U_1}^2 + \sigma_{U_2^*}^2 + (\eta - 1)^2 \sigma_X^2 \right\}.$$

This is greater than $\sigma_{U_1}^2$, and thus results in over correction for attenuation, when

$$\sigma_{U_2^*}^2 + (\eta - 1)^2 \sigma_X^2 > \sigma_{U_1}^2. \quad (3.8)$$

For the model described above, U_1 is the measurement error at baseline and U_2^* is the sum of the measurement error at post-baseline and the residual error $X_* - E(X_* | X)$. Thus under constant measurement error variance, $\sigma_{U_2^*}^2 > \sigma_{U_1}^2$ and the inequality in (3.8) holds.

The third correction for attenuation, $\tilde{\lambda}_1^*$ in (3.5), depends on the post-baseline measurements via $\hat{\sigma}_T^2$ as well as through $s_{W,1}^{*2}$. This makes it difficult to assess the effect of departures from

the replicate measurements model. However, the calculations are manageable in the absence of covariates. In this case $\tilde{\lambda}_1^*$ approaches asymptotically

$$g(\eta) = \frac{(\eta + 1)^2 \sigma_X^2 / 4 + (\sigma_{U_1}^2 + \sigma_{U_2}^2) / 4}{\eta \sigma_X^2}.$$

Since $g(1) > \lambda_1^*$ when $\sigma_{U_2}^2 > \sigma_{U_1}^2$, and $\partial g(\eta) / \partial \eta < 0$ whenever $\eta^2 \leq 1$, the net effect of departures from the replicate measurements model in the direction of (3.7) in the likely situation that η is less than 1 but positive, is to inflate the correction for attenuation, λ_1^* . Regardless of the presence or absence of covariates, the bias in $\tilde{\lambda}_1^*$ affects the corrections for attenuation in the other studies only through $\hat{\sigma}_U^2$, which is generally overestimated under (3.7). Thus the corrections for attenuation in studies $k = 2, \dots, \mathcal{K}$, are positively biased as well.

The instrumental variable estimator, $\tilde{\lambda}_{1,IV}$, depends only on a nonzero correlation between T and X and linearity in the regressions of W and X on (T, Z_k) , $k = 1, \dots, \mathcal{K}$, and is therefore robust to departures from the replicate-measurements model in the direction of (3.7).

Summarizing these results we have: (i) all of the corrections for attenuation are consistent when the replicate-measurements model holds; (ii) only the instrumental-variable correction for attenuation is consistent under the more general second-measurement error model (3.7); and (iii) excluding the instrumental variable correction, the general effect of departures from the replicate-measurements model is to inflate the corrections for attenuation.

The consistency robustness of the instrumental-variable method to departures from the replicate-measurements model is obtained at the expense of greater finite-sample variability in the estimated corrections for attenuation. Table 1 displays the results of a simulation study designed to compare the four estimators $\bar{\lambda}_1$, $\tilde{\lambda}_1$, $\tilde{\lambda}_1^*$ and $\tilde{\lambda}_{1,IV}$ and the corresponding estimators of β . The four methods are designated RM (regression to the mean), MM and MM* (method of moments) and IV (instrumental variable) respectively.

The model for the simulation study was:

$$Y | X, Z \sim N(\alpha + \beta X + \gamma^T Z, 1), \quad \alpha = 0, \beta = 1, \gamma = (0, 0)^T,$$

$$(X, Z)^T \sim N(0_{3 \times 1}, I_3), \quad W | X \sim N(X, \sigma_U^2),$$

$$T | X \sim N(\xi + \eta X, 1 - \rho^2 + \sigma_U^2), \quad \xi = 0, \eta = \rho.$$

The measurement error variance and ρ were investigated at two levels $\sigma_U^2 = 0.25, 1.00$ and $\rho^2 = 1.00, 0.90$. Sample size was set at $n = 100$, and 500 independent data sets were generated at each factor level combination. Note that for methods RM, MM and IV, $\lambda = 1.25, 2.00$ when $\sigma_U^2 = 0.25, 1.00$ respectively, while for method MM*, $\lambda = 1.125, 1.50$ when $\sigma_U^2 = 0.25, 1.00$.

Monte Carlo means and mean squared errors are reported for the estimates of β and λ . The instrumental variable estimators of λ and β have generally smaller biases than the other estimators in the study. Furthermore the biases for the cases of $\rho^2 = 1.00$ and 0.90 are comparable, confirming the consistency robustness noted above. In terms of mean squared error $\tilde{\beta}_{IV}$ performs well also. However, the mean squared error of $\tilde{\lambda}_{IV}$ is consistently larger than the mean squared errors of the other three estimators of λ indicating greater sampling variability in this estimator.

4. DISCUSSION OF THE META-ANALYSIS OF CHD AND DBP

4.1. The Corrections in the Framingham Data

MacMahon, et al. (1990) describe analysis of the Framingham study. We give here analyses which are meant to illustrate the issues discussed in the previous sections. The Framingham data consist of measurements at different examinations spaced two years apart, so that, for example, Exam 5 takes place 4 years after Exam 3. In what follows, we define X to be the true diastolic blood pressure (DBP).

We consider two possibilities for the definitions of W , T and Y :

- (*Follow-up Instrument*) $W = \text{DBP at Exam 3}$, $T = \text{DBP at Exam 5}$, $Y = \text{coronary heart disease (CHD) incidence in a 10 year follow-up from Exam 3}$;
- (*Precursor Instrument*) $W_* = \text{DBP at Exam 5}$, $T_* = \text{DBP at Exam 3}$, $Y = \text{CHD incidence in a 10 year follow-up from Exam 5}$.

The first situation (follow-up instrument) corresponds more closely to that described by MacMahon, et al. (1990), in the sense that they defined T as the four-year follow-up measure of blood pressure. The other predictors Z used here are age (at Exam 3), serum cholesterol (at Exam 3) and cigarettes/day (at Exam 1). It might be preferable to use Exam 5 instead of Exam 3 for age and serum cholesterol in the second situation, but we have not done so in order to correspond as closely as possible to the analysis of MacMahon, et al. (1990), and because the results for DBP will likely

not change significantly upon the redefinition, due to the lack of correlation between observed DBP and Z , see below.

In order to keep the discussion as simple as possible, we will ignore the possibility that the last two components of Z are measured with error, noting only that the regression method (3.2), the moments method (3.4) and the instrumental variables method (3.6) are all easily extended to handle more than one covariate measured with error.

The use of T_* as a precursor instrument (the second situation defined above) most closely fits into the framework for instrumental variables discussed in the previous section. The use of T as a follow-up instrument (the first situation defined above) in this problem may pose problems. Due to censoring by death, T is an unusual "second" measurement, i.e., it is not possible to measure T in those subjects who die before Exam 5. For the instrumental variable method to make sense in this context, it seems necessary that the logistic regression of Y on (X, Z) be approximately the same in the population starting from Exam 3 as it is in the censored population who survived to Exam 5. This may be a problematic assumption in this example, although we are unable to reach definitive conclusions.

We now describe results of the analyses for all estimators under the two definitions of W , T and Y described at the start of this section.

Definition 1. (*Follow-up Instrument*): For the moments and regression methods, we followed the same procedure as MacMahon, et al., namely deleting those who developed CHD or died in the four years post baseline. In the Framingham data, we found W had sample variance 140.80, while $s_{W,1}^2 = 138.443$. This means, in effect, that the predictors Z are uncorrelated with W . The method of moments correction is $\bar{\lambda}_1 = 1.69$ since the estimate of σ_T^2 equals 56.39. If one replaces $s_{W,1}^2$ by the sample variance of W , then the moments correction becomes 1.67. In this particular case, with this particular set of covariates, ignoring the covariates has little effect.

The coefficient for W in regressing T on (Z, W) is .5872, leading to a regression correction of $\bar{\lambda}_1 = 1.70$, essentially the same as the moments correction.

The sample mean and variance of W are 82.16 and 140.80, respectively, while the sample mean and variance of T are 83.51 and 136.93. Although the sample means are quite close in numerical

value, the difference is statistically significant. With $\sigma_U^2 = 56.39$ and $n = 1605$, the number of subjects who were alive and disease free at Exams 3 and 5, the test statistic for equality of means is $t = (83.51 - 82.16)/\sqrt{2\hat{\sigma}_U^2/n} = 5.09$, approximately 5 standard errors from zero under the replication model. This does not prove that the instrumental variable model is appropriate, but it does suggest the possibility of problems with the replicate measurements model.

For the instrumental variable method, using only those who were alive and free of CHD at Exam 5, we find that $\hat{\beta}_{W|T,Z} = .5958$. Among those alive and free of CHD at Exam 3, $\hat{\beta}_{Y|W,Z} = .02439$. Among those alive at Exam 5, $\hat{\beta}_{Y|T,Z} = .01988$. This yields a correction of $\tilde{\lambda}_{1,IV} = 1.37$.

Definition 2. (Precursor Instrument): The moments correction does not change in this situation. The regression correction does change, but only in the third significant digit. For the instrumental variables method, $\hat{\beta}_{Y|W^*,Z} = .01985$, $\hat{\beta}_{Y|T^*,Z} = .01580$, and hence $\tilde{\lambda}_{1,IV} = 1.34$.

To summarize the results of this purely illustrative exercise, all methods yield similar corrections using either a precursor or a follow-up instrument. The moments and regression method corrections are essentially the same (≈ 1.70), while the instrumental variable method suggests a smaller change (≈ 1.35). Under an assumption of normality, the standard error of the moments correction estimate is approximately 0.043 under the replication model; see Appendix A. We do not have full access to the data and hence cannot verify the normality assumption. However, it does appear that the methods suggest corrections which are nontrivially different.

In order to understand whether the differences in the corrections, and hence in the parameter estimates, can be explained by a departure from the replicate measurements model, or might be due to some other systematic bias in one of the methods, we performed a small simulation study. We assumed that there were no additional variables Z , and we observe the response Y , the surrogate W and the second measurement T . The parameters in the simulation were chosen to correspond roughly to those reported in the Framingham data. The sample size was $n = 1605$, of whom in the Framingham data, 157 CHD cases developed within 10 years of Exam #3. It was noted above that $\text{var}(X) \approx 80$, $\text{var}(U) \approx 60$, $\lambda \approx 1.7$ and hence $\beta \approx .0244 \times 1.7$. We standardized so that X was normally distributed with mean 0.0 and variance 1.0. With this normalization, our simulations were constructed under the distributional assumptions that $W|X \sim N(X, \sigma_U^2)$ and

$T|X \sim N(\xi + \eta X, 1 - \rho^2 + \sigma_U^2)$, with $\xi = 0$, $\eta = \rho$, $\sigma_U^2 = .75$, $\rho^2 = .9, 1.0$. The binary responses followed a logistic linear model with intercept -2.25 and slope 0.371. We also performed, but do not report on here, simulations with for slope parameters 0.1855 and 0.742. The results of the simulation are given in Table 2, which is based on 250 repetitions of the experiment.

In reading Table 2, one should note that there is no “correct” value of λ , because in logistic regression the corrections are only approximate. However, some conclusions are possible for the set of circumstances defined by the simulation. With regard to estimation of λ , Table 2 is consistent with the the theoretical results that all methods are approximately unbiased in the replicate measurements model, ($\rho^2 = 1$), but that the method of moments and the regression to the mean methods over estimate λ relative to the instrumental variable method when $\rho^2 < 1$. The same conclusions apply to estimation of β . The means of the instrumental variable estimator are similar at $\rho^2 = 1.00$ and 0.90, whereas the other two procedures exhibit a positive bias at $\rho^2 = 0.90$.

There is a large difference in the variability of the estimates of λ and β . The instrumental variable estimate of λ is approximately nine times more variable than the other two estimators, whereas for estimation of β the instrumental variable estimator has comparable variability. We conjecture that the explanation is due to a negative correlation between the instrumental variable estimator of λ and the naive estimator of β .

The magnitudes of the differences of the estimates in our simulation study are not as great as the differences observed in the Framingham estimates. Thus, the differences observed between the estimators from the Framingham data cannot be explained simply as bias induced by departures from the replicate measurements model or by any other systematic bias of any of the methods under a normal distribution for errors and predictors. However, the differences in the estimates of λ can be explained by the increased variability of the IV estimator, although other explanations such as outliers cannot be ruled out.

Although estimation of λ is of some independent interest, β is the primary parameter of interest. Thus the various methods should ultimately be evaluated in terms of the β estimates they produce. Translated to the standardized scale used in the simulations, we found that the moments and instrumental variable estimates differed by $\tilde{\beta}_1 - \tilde{\beta}_{1,IV} \approx 0.076 (= (1.70 - 1.35) \times .0244 \times \sqrt{80})$.

Recall that the estimate of β not corrected for attenuation is .0244, and $\sqrt{80}$ is approximately the standard deviation of X in the Framingham data. In our simulations, the standard deviation of the difference was 0.106 and 0.101 for $\rho^2 = 0.90$ and 1.00, respectively. This suggests that the various point estimates of β in the Framingham data are not statistically significantly different. In addition, these calculations point out the importance of attaching standard errors to parameter estimates; we have not done so here because of lack of access to the data.

We note that the moments and regression estimators are special cases of a class of estimators studied by Carroll and Stefanski (1990). Asymptotic standard errors for the moments and regression estimators can be derived from the general formulas in this paper.

4.2. The Method of MacMahon, et al. (1990) applied to Framingham Data

The method of MacMahon, et al. (1990) is closely related to the regression method (3.2) and is described as follows. Let C_1 and C_2 be extreme intervals, i.e., $C_1 = \{W \leq \text{lower bound}\}$ and $C_2 = \{W \geq \text{upper bound}\}$, and for $p = 1, 2$, define

$$\hat{C}_{p1} = \frac{n_1^{-1} \sum_{i=1}^{n_1} W_i I(W_i \in C_p)}{n_1^{-1} \sum_{i=1}^{n_1} I(W_i \in C_p)},$$

$$\hat{C}_{p2} = \frac{n_1^{-1} \sum_{i=1}^{n_1} T_i I(W_i \in C_p)}{n_1^{-1} \sum_{i=1}^{n_1} I(W_i \in C_p)}$$

Then their correction for attenuation estimate is

$$\hat{\lambda}_{M,1} = \frac{\hat{C}_{21} - \hat{C}_{11}}{\hat{C}_{22} - \hat{C}_{12}}. \quad (4.1)$$

Upon replacing the numerators and denominators in \hat{C}_{p1} and \hat{C}_{p2} ($p = 1, 2$), by their expectations, it is easily shown that that when T and W are replicates as defined in § 3.2, and X is independent of Z , then $\hat{\lambda}_{M,1}$ consistently estimates λ_1 , although we can show theoretically that it is generally inefficient. However, violation of these assumptions can result in inconsistent estimators.

For example, suppose that we continue to assume normality and continue to assume that T is a replicate, but now allow X and Z to be correlated. Then (4.1) estimates not λ_1 but instead estimates $\text{Var}(W) / \{\text{Var}(W) - \sigma_U^2\}$, leading to a correction which is generally too *small* since $\text{Var}(W) > \sigma_W^2$.

On the other hand, according to our analysis, the effect of T not being a replicate is that the correction (4.1) is generally too *large*, see § 2.

In the Framingham data, using T as a follow-up instrument, W and Z are essentially uncorrelated, so that the correction (4.1) yields a reasonable estimate of a 60% correction in the ordinary logistic regression coefficients.

In summary, the correction (4.1) proposed by MacMahon, et al. is inefficient. Its appropriateness rests fundamentally on the two assumptions that T is a replicate and that X is unrelated to all the other covariates, assumptions which appear to be reasonable in the Framingham data. Significant violation of these assumptions means that this correction should not be used.

4.3. Implications for Meta-Analyses

We highlight the issues that arise in meta-analyses, using as a second study the MRFIT data, see Kannel, et al. (1986). MacMahon, et al. (1990) compute their correction (4.1) in the Framingham data, and then apply it to all the other data sets in their meta-analysis, including the MRFIT data. As illustrated by (2.3), this is not always appropriate since the correction for attenuation need not be constant across studies even when the measurement error variance is. Since we do not have access to the MRFIT data, we can only illustrate the potential pitfalls of applying the correction from the first study (Framingham) to a second study (MRFIT) in a meta-analysis.

The following numbers are purely illustrative. We will ignore the covariates Z as the evidence suggests that they are at best weakly predictive of W , i.e., we will assume that X is independent of Z and that T is a replicate. Then, for Framingham men, the sample variance of W is 140.80, $\hat{\sigma}_U^2 = 56.39$ from (3.3) and hence $\tilde{\lambda}_1 = 1.67$. For the MRFIT data, Kannel, et al. state that the sample variance of W is 110.25, so that $\tilde{\lambda}_2 = 2.04$. In other words, the fact that the predictor W is much less variable in the MRFIT study means that its regression coefficient should have been corrected by the factor 2.04, not 1.66 as suggested by MacMahon, et al., a 23% difference.

Of course these conclusions are highly dependent on the assumption of constant measurement error variance across studies. We note that the MRFIT measurements were taken in 1973-75, whereas the Framingham measurements date to 1955. Thus one might expect greater reliability in the MRFIT measurements. If instead we assume that the variances of X in the Framingham and MRFIT study populations are equal, then the difference between the respective variances of W (140.80 - 110.25), is an estimate of the difference in the measurement error variances. That is, with

$\hat{\sigma}_U^2 = 56.39$ for the Framingham study, the estimate of the measurement error variance for MRFIT is $\hat{\sigma}_U^2 = 56.39 - (140.80 - 110.25) = 25.84$. This results in an estimated correction for attenuation of $\hat{\lambda}_2 = 110.25/84.41 = 1.31$, a 21% decrease relative to the correction reported by MacMahon, et al. (1990).

As this example makes clear, assumptions of constancy across studies, either with respect to the population variances of X or with respect to the measurement error variances, can have a substantial impact on conclusions.

5. CONCLUSIONS

We have considered corrections for attenuation in semilinear regression models. There is a tendency for users of measurement error methods to base corrections for attenuation on the variance of the fallible covariate W . We have noted that the correction depends on the size of the measurement error σ_U^2 and on the size of the regression MSE, σ_W^2 , from regressing the surrogate W on all the other predictors Z , see (2.3). Failing to take into account the presence of other covariates can lead to an undercorrection of the regression coefficients. These comments are relevant to single-study measurement error analyses as well as to meta-analyses.

In correcting the results of a single study, we have distinguished between *replicate measurements* and *second measurements* of a fallible covariate. For replicates, the regression to the mean (RM) method (3.2), the method of moments (MM) methods (3.4)–(3.5) and the instrumental variable (IV) method (3.6) all yield consistent estimates of the attenuation.

For those cases that the second measurement is biased for the fallible covariate, only the IV method consistently estimates the attenuation. We have shown that in this case, the usual effect of using RM or MM is to *overestimate* the correction for attenuation.

We have also noted that the correction for attenuation proposed by MacMahon, et al. (1990) yields a consistent estimate only if the second measurement is a replicate *and* the true and surrogate predictors X and W are unrelated to all the other covariates Z . Failure of these assumptions generally causes under- and over-corrections, respectively.

We have also discussed the use of corrections for attenuation in meta-analysis. We have stressed in (2.3)–(2.4) that even when the measurement error variance is constant across studies, *the proper*

correction for attenuation may vary from study to study, depending on the marginal distribution of the predictors through the mean squared error from the linear regression of W on Z . Also, as the example in § 4.3 demonstrates, a single correction for attenuation is not possible when the measurement error variances differ across studies. Thus, in general, study-specific corrections are necessary, and using the same correction for attenuation across studies generally leads to biased estimates. Furthermore, neither the direction nor the magnitude of the bias is predictable.

ACKNOWLEDGEMENT

Carroll's research was supported by a grant from the National Institutes of Health. We are indebted to Paul Sorlie for computing the Framingham numbers.

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Appendix 1. CALCULATION OF STANDARD ERROR FOR MOMENTS ESTIMATE

Let $Y_1 = 2^{-1/2}(W - T)$ and $Y_2 = W$. Let s_{11} and s_{22} be the sample covariance matrices for the Y_1 and Y_2 observations. Then

$$\begin{aligned}\sigma_{11} &= \sigma_U^2; \\ \sigma_{22} &= \sigma_W^2,\end{aligned}$$

and

$$\sigma_{12} = \sigma_{21} = \text{cov}(Y_1, Y_2) = 2^{-1/2} \sigma_U^2.$$

According to Fuller's Appendix to Chapter 1, this means that

$$\begin{aligned}\text{cov}(s_{11}, s_{11}) &= 2\sigma_{11}^2; \\ \text{cov}(s_{22}, s_{22}) &= 2\sigma_{22}^2; \\ \text{cov}(s_{11}, s_{22}) &= 2\sigma_{12}^2 = \sigma_U^4.\end{aligned}$$

If

$$\hat{\lambda} = \frac{\hat{\sigma}_W^2}{\hat{\sigma}_W^2 - \hat{\sigma}_U^2},$$

then

$$\hat{\lambda} - \lambda \approx (1 - \lambda) \frac{\hat{\sigma}_W^2 - \sigma_W^2}{\sigma_X^2} + \lambda \frac{\hat{\sigma}_U^2 - \sigma_U^2}{\sigma_X^2}.$$

Hence,

$$n\text{var}(\hat{\lambda} - \lambda) \approx 2 \frac{(1 - \lambda)^2 \sigma_W^4 + \lambda^2 \sigma_U^4 + \lambda(1 - \lambda) \sigma_U^4}{\sigma_X^4}.$$

Because $\sigma_U^2/\sigma_X^2 = \lambda - 1$ and $\lambda = \sigma_W^2/\sigma_X^2$, we thus have

$$\text{var}(\hat{\lambda} - \lambda) \approx 2\lambda^2(\lambda - 1)^2 \{2 + (1 - \lambda)/\lambda\}/n.$$

With $\lambda = 1.6$ as in the Framingham study, and $n = 1605$, the standard error for $\hat{\lambda}$ is approximately 0.043.

TABLE 1: COMPARISON OF ESTIMATORS

This simulation is described in § 3.4. Here “RM” denotes the regression to the mean method correction (3.2), “MM” and “MM*” denote the method of moments corrections (3.4) and (3.5), and “IV” is the instrumental variable correction (3.6). Table entries are Monte Carlo means (upper entry) and mean squared errors (lower entry).

		$\sigma_U^2 = 0.25$				$\sigma_U^2 = 1.00$			
		RM	MM	MM*	IV	RM	MM	MM*	IV
$\rho^2 = 1.00$	β	1.005 .018	1.009 .018	1.003 .015	1.001 .017	1.033 .060	1.053 .074	1.027 .041	1.024 .050
	λ	1.253 .010	1.255 .004	1.127 .001	1.250 .013	2.062 .216	2.089 .188	1.529 .037	2.071 .344
$\rho^2 = 0.90$	β	1.062 .026	1.065 .026	1.032 .018	1.002 .019	1.097 .091	1.120 .111	1.061 .055	1.027 .057
	λ	1.323 .021	1.325 .012	1.161 .003	1.250 .016	2.190 .347	2.223 .317	1.592 .062	2.079 .375

TABLE 2: COMPARISON OF ESTIMATORS IN LOGISTIC REGRESSION

This simulation is described in § 4.1. Here “RM” denotes the regression to the mean method correction (3.2), “MM” denotes the method of moments correction (3.4), and “IV” is the instrumental variable correction (3.6). The term “MAD” refers to the median absolute deviation from the median, “MSE” is mean squared error, “Mean AE” is the mean absolute error and “Median AE” is the median absolute error. The correct value of $\beta = .371$.

Results for λ				
	$\rho^2 = 1.00$	MM	RM	IV
Mean		1.756	1.755	1.780
Median		1.754	1.756	1.670
S.D.		.063	.065	.543
MAD		.045	.046	.323
	$\rho^2 = 0.90$	MM	RM	IV
Mean		1.851	1.846	1.775
Median		1.846	1.839	1.668
S.D.		.070	.071	.678
MAD		.049	.045	.290
Results for $\beta = .371$				
	$\rho^2 = 1.00$	MM	RM	IV
Mean		.386	.386	.376
Median		.389	.389	.377
MSE		.014	.014	.013
Mean AE		.094	.093	.090
Median AE		.081	.079	.079
	$\rho^2 = 0.90$	MM	RM	IV
Mean		.409	.408	.374
Median		.406	.409	.369
MSE		.015	.015	.013
Mean AE		.099	.097	.090
Median AE		.082	.076	.075