Parainfluenza-3 and bovine respiratory syncytial virus: intraherd correlation adjusted for sensitivity and specificity

Parainfluenza-3 y virus respiratorio bovino sincitial: correlación intrahato ajustado por la sensibilidad y especificidad

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ABSTRACT

Objective. The purpose of this study was to compare the intra-class correlation coefficients (ICC) and design effects (D) estimates adjusted or unadjusted for sensibility (Se) and specificity (Sp) of the diagnostic tests using a Bayesian procedure. Materials and methods. Sera from 232 animals from 44 randomly selected herds, to detect antibodies against parainfluenza-3 virus (PIV3) from non-vaccinated dual-purpose cattle from Colima Mexico, were used. Only 176 animals from 33 herds were used to evaluate the presence of the bovine respiratory syncytial virus (BRSV). Results. The ICC and D values adjusted and unadjusted for PIV3 were 0.33, 2.73, 0.32, and 2.71, respectively. For BRSV the values were 0.31, 2.64, 0.28 and 2.49. Conclusions. The adjusted or unadjusted ICC and D estimates were similar because of the high Se and Sp of the diagnostic tests and the relatively high prevalence of the diseases here studied.

Key words: Bovine respiratory syncytial virus, design effect, intraclass correlation, parainfluenza-3 virus, (Source: CAB).

RESUMEN

Objetivo. El propósito de este estudio fue comparar los valores de los coeficientes de correlación intraclase (ICC) y efecto de diseño (D) ajustados y no ajustados por la sensibilidad (Se) y especificidad (Es) de la prueba de diagnóstico usando procedimientos bayesianos. Materiales y métodos. Se utilizaron los sueros sanguíneos de 232 animales de 44 hatos seleccionados al azar, para detectar anticuerpos contra el virus de la parainfluenza-3 (VPI3) en bovinos de doble propósito, no vacunados en Colima, México. Se usaron 176 animales de 33 hatos para evaluar la presencia del virus respiratorio sincitial bovino (VRSB) Resultados. Los valores ajustados y no ajustados de ICC y D para PIV3 fueron 0.33, 2.73, 0.32 y 2.71, respectivamente. Para VRSB los valores fueron 0.31, 2.64, 0.28 y 2.49. Conclusiones. Los estimadores de ICC y D fueron similares debido a la alta Se y Es de las pruebas de diagnóstico y a la relativa alta prevalencia de las enfermedades estudiadas.

Palabras clave: Correlación intraclase, efecto de diseño, virus parainfluenza-3, virus respiratorio sincitial bovino (Fuente: CAB).
INTRODUCTION

The responses to the animal infection within herds (clusters) are normally correlated due to similarity caused by environmental and geographical factors, by characteristics linked to the choice of herd, or by management practices within herds. The intraclass correlation coefficient (ICC) is a measure of correlation between the individuals from a cluster (herd) that is used in the design and statistical analysis of epidemiologic studies (1). As an analytical tool, ICC has been used to adjust for infection clustering as part of a variance inflation factor (VIF) or design effect (D) for prevalence estimations (2).

The square root of the D measures, the increase in the standard error of the estimated prevalence due to the sampling procedure and it is used in calculating the correct sample size in cluster studies (3,4). It can be also used to adjust the significant levels of risk factors and confidence intervals in cross sectional cluster studies (4, 5). Knowing D for a specific disease may sample size calculation relatively easily. According to Solís et al (3), sample size for a cluster study can be calculated as D*n; where n is the sample size for a simple random survey (n=Z^2pq/L^2); p is the expected prevalence; Z is the table value for a standardized normal distribution with a desired confidence level and L is the desired precision or sampling error.

Statistical procedures that account for infection clustering but do not rely on ICC were also used (6) including Bayesian hierarchical models presented in Branscum et al (1,7). If the infectious status of each sampled animal was known, positive or negative ICC data could be estimated in different ways (8). However, in many epidemiological studies, researchers have only apparent infectious status data which can be used to estimate ICC, so ICC could be biased. Field studies are typically implemented by sampling animals from multiple herds and testing the sampled animals with an imperfect diagnostic test. Using data obtained from these studies, makes estimators calculated as suggested by Ridout et al (8) biased unless they are adjusted for imperfect test accuracy (1).

The previous authors proposed a Bayesian approach to estimate the ICC, which incorporates imperfect sensitivity (Se) and specificity (Sp), in a simulation study. In that study, they showed that ICC values obtained using analysis of variance are two- to three-fold lower than those using Bayesian approaches, and adjusting herd prevalence for Se and Sp in the diagnostic test. However, Branscum et al (1) did not compare the ICC values obtained using the adjusted and unadjusted data of the Bayesian approach.

The objective of this study was to determine the effect of imperfect sensitivity and specificity on ICC and D estimates using Bayesian procedures.

MATERIAL AND METHODS

Data source and study design. A two-stage cross-sectional study was carried out from November 2007 to March 2008 in Colima, Mexico. Information on 232 animals from 44 randomly selected farms, was used (9) to detect antibodies against parainfluenza-3 virus (PIV3) from non-vaccinated dual-purpose cattle from Colima, Mexico. However, because of economic constraints, blood samples of only 33 of the 44 farms were evaluated for BRSV. Herd sizes varied from 12 to 350 animals. The number of animals sampled within each herd varied between 4 and 8, and only animals over 6 months of age were sampled, to avoid the detection of maternal antibodies. The detection of serum antibodies for the viral diseases PIV3 and BRSV was carried out using commercial indirect ELISA kits (SVANOVA Biotech, Uppsala, Sweden). The individual and herd prevalence for PIV3 and BRSV have been reported in a previous study (9) and are shown in Table 1. The sensitivity (Se) and specificity (Sp) of the diagnostic tests for those infections were 100 and 86%, and 95 and 92% (10), respectively. The optical density (OD) was measured at 450 nm with a Titertek Multiscan Spectrophotometer (Flow Laboratories, Irvine, UK). The sample OD-corrected values < 0.20 were considered negative as indicated by the ELISA kit.

Intraherd correlation coefficient and design effect estimates. Binomial data (Yij) were modeled as beta-binomial assuming independent beta prior distributions for Se, Sp, µ (mean prevalence distribution) and γ (variability of prevalence) modeled using a gamma prior distribution. The Bayesian model was:

\[ Y_{ij} \mid p_i \sim Bernoulli(p_i) \]
\[ p_i = n_i Se + (1-n_i)(1-Sp) \]
\[ n_i = n_i^* \text{ with probability } 1-T \]
\[ n_i = 0 \text{ with probability } T \]
\[ n_i^* \mid \mu_i, \gamma \sim Beta(\mu_i, \gamma(1-\mu)) \]
\[ \mu \sim Beta(a_\mu, \beta_\mu) \]
\[ \gamma \sim Gamma(\alpha_\gamma, \beta_\gamma) \]
\[ Se \sim Beta(a_{Se}, \beta_{Se}) \]
\[ Sp \sim Beta(a_{Sp}, \beta_{Sp}) \]
Where \( p_i \) is the apparent prevalence for the \( i \)th herd, \( \pi_i \) is the infection prevalence of the \( i \)th herd and \( \tau \) is the proportion of infected herds sampled, which was set to the herd prevalence of the sample.

The ICC based on the above model was calculated as suggested by Branscum et al (1):

\[
ICC = \frac{1 - \mu + \mu(y + 1)(1 - \tau)}{1 - \tau y}
\]

D was estimated as (3):

\[
D = 1 + (b-1) \times ICC
\]

Where \( b \) is the average number of animals sampled per herd.

The model used the WinBUGS program (11). In this study a total of 20,000 samples of possible ICC and D values were generated and the results of the first 500 rounds were deleted. The parameters of the prior beta distributions (\( \alpha_\mu \) and \( \beta_\mu \)) were obtained with the Betabuster software (12). On the other hand, the parameters of the prior gamma distribution (\( \alpha_\gamma \) and \( \beta_\gamma \)) were obtained according to Hanson et al (13), using a program developed in the Gauss software for Windows. The parameters used in this paper are shown in table 1.

### RESULTS

The ICC and D values, adjusted for Se and Sp of their respective diagnostic tests, and their 95% credible intervals for PIV3 and BRSV are presented in table 2. ICC values for PIV3 were 0.33 (adjusted) and 0.32 (unadjusted) and their D values were 2.73 (adjusted) and 2.71 (unadjusted). For BRSV the ICC values were 0.31 (adjusted) and 0.28 (unadjusted) and the D values 2.64 (adjusted) and 2.49 (unadjusted). Credible intervals are provided in table 2.

#### Table 1

<table>
<thead>
<tr>
<th>Infection</th>
<th>IP</th>
<th>HP</th>
<th>SST</th>
<th>BDP</th>
<th>GDP</th>
</tr>
</thead>
<tbody>
<tr>
<td>Parainfluenza-3</td>
<td>60.8</td>
<td>78.7</td>
<td>100,86</td>
<td>33.7</td>
<td>21.6</td>
</tr>
<tr>
<td>Bovine respiratory syncytial</td>
<td>52.2</td>
<td>93.2</td>
<td>95,92</td>
<td>21.1</td>
<td>18.73</td>
</tr>
</tbody>
</table>


#### Table 2

<table>
<thead>
<tr>
<th>Infection</th>
<th>ICC</th>
<th>95% CI</th>
<th>Design effect</th>
<th>95% CI</th>
</tr>
</thead>
<tbody>
<tr>
<td>Parainfluenza type 3**</td>
<td>0.33</td>
<td>0.24, 0.44</td>
<td>2.73</td>
<td>2.30, 3.31</td>
</tr>
<tr>
<td>Parainfluenza type 3**</td>
<td>0.32</td>
<td>0.25, 0.42</td>
<td>2.71</td>
<td>2.33, 3.22</td>
</tr>
<tr>
<td>Bovine respiratory syncytial</td>
<td>0.31</td>
<td>0.19, 0.46</td>
<td>2.64</td>
<td>2.04, 3.45</td>
</tr>
<tr>
<td>Bovine respiratory syncytial</td>
<td>0.28</td>
<td>0.19, 0.40</td>
<td>2.49</td>
<td>2.00, 3.11</td>
</tr>
</tbody>
</table>

ICC: Intraherd correlation coefficient; CI: Credible interval.

*Number of herds (n=33; average number of animals sampled per herd (b=5.33); Total number of animals sampled (N=176);

**Number of herds (n=44; average number of animals sampled per herd (b=5.27); Total number of animals sampled (N=232).

*Adjusted.

Unadjusted for sensitivity and specificity of the diagnostic test.

#### DISCUSSION

The results for the unadjusted and adjusted ICC and D closely agree for both PIV3 and BRSV data because the Se and Sp of the ELISA tests were high. A simulated scenario considering Se and Sp of 70% provided ICC values of 0.38 and 0.35 for PIV3 and BRSV and D values of 3.03 and 2.83, respectively, which produced a slight increase in the parametrical estimates. Another factor that may affect the results here found is the prevalence of the diseases. Branscum et al (1) used a prevalence of 26% for ovine progressive pneumonia, whereas in this study the prevalence for PIV3 and BRSV was much higher (60.8 and 52.2%, respectively) and closer to 50%.

Branscum et al (1) showed that treating diagnostic test results as indices of true infection status and variance analysis procedures can result in two- to three-fold underestimation of the true ICC. They concluded that their Bayesian model for ICC estimation, which accounted for diagnostic test imperfection and uncertainty in the true values of Se and Sp, provided reasonable estimates of the ICC for simulated data. However, they compared ICC estimates using analysis of variance and Bayesian procedures. They also observed that their ICC estimates for data from a study on ovine progressive pneumonia gave similar results by using analysis of variance and Bayesian procedures, because of the high Se and Sp in the diagnostic test. In the present study the adjusted and unadjusted estimates were both obtained using Bayesian procedures.

Estimations of ICC or D are needed in order to perform sample size calculation when the unit of interest (animal) is clustered within herds. One program (among others) that uses D to calculate sample size is OpenEpi (14). ICC or D also could...
be used for adjustment of standard errors, and prevalence and odd ratios confidence intervals in cluster sampling studies, where computer programs are not available. However, because of the variability in herd susceptibility to diseases, and environmental and management differences between regions, it may be inappropriate to uniformly assign a particular value of the ICC to a specific disease process, even when the ICC has been estimated through a well-designed epidemiologic study.

ICC and D values for PIV3 and BRSV (adjusted for Se and Sp) found in this study (0.33 and 2.73, and 0.31 and 2.64, respectively) were different from those reported (4) in beef cattle in Yucatan (0.19 and 3.44 for PIV3, and respectively), which used an analysis of variance methodology. The prevalence and average number of animals per herd in the Solis-Calderon et al (4) study were 90.8% and 14 animals, and 85.6% and 14 animals, respectively. Even though the ICC values here obtained were larger, the D values were smaller, because of the small number of animals (mean=5.3 animals) sampled in each herd. McDermott and Schukken (15) showed that cluster size has the single most important effect on the VIF or D. The ICC values for PIV3 and BRSV obtained in this study indicate that nearly 30% of the variation of the disease is explained by in-between herd variation and 70% by animal or individual risk factors within herds.

The D values for PIV3 and BRSV indicate that under the conditions of this study, sample size should be 2.73 and 2.64 times greater than the sample size needed for a simple random sampling survey, in order to achieve the same prevalence precision. Unfortunately, ICC and D are unknown at the beginning of the study, so it is desirable that cluster-sampling studies report both the ICC and D values to design better survey studies and obtain unbiased prevalence confidence intervals.

The differences here reported and those by Solís-Calderón et al (4) suggest that each region requires the use of appropriate ICC and D values, since the incidence of a disease depends on environmental factors in the region and specific practices within the herd, as well as the number of animals sampled per herd. However, ICC and D values for a given disease could be used from one region to another, when management and environmental conditions are similar and where the infection is endemic and in reasonable equilibrium (1).

In conclusion under the conditions of this study the ICC and D values for PIV3 and BRSV indicate that similar sample sizes are required for both diseases to achieve the required precision in prevalence estimates. The report of ICC and D values could help plan and design epidemiological studies and to adjust parameter estimates of cluster studies.

Acknowledgement

Authors thanks Dr Jorge Sosa Argaez for calculating the gamma distribution parameters used in this study.

REFERENCES


