**Title:**

**Characterizing the relationship between porcine reproductive and respiratory syndrome virus prevalence at the piglet and litter level in a farrowing room using computer models**

**Authorship**

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**Ethical statement**

The authors confirm that they have complied with the ethical guidelines of this journal. No animal use protocol was required as this study was done using computer models and retrospective data

**Summary**

The estimated prevalence is an important assumption in calculating the minimum number of units to be sampled for PRRSV surveillance/monitoring programs; the minimum number of litters to be sampled (sample size) for family oral fluids (FOFs) in a farrowing room depends on the estimated number of positive litters in a farrowing room, the diagnostic performance of the test, and a chosen confidence level. The relationship between the estimated PRRSV prevalence at the individual piglet level, the estimated PRRSV prevalence at the litter level, and the apparent PRRSV prevalence from FOF sampling in a farrowing room has not been previously characterized.

Using Monte Carlo simulations and data from a previous study, the relationship between the proportion of PRRSV-positive (viremic) pigs, the proportion of PRRSV-positive litters, and the likely proportion of litters to be positive by a FOF RT-rtPCR test in a farrowing room was characterized; taking into account the degree of clustering of viremic pigs within farrowing rooms.

The results of this study provide guidelines on how practitioners can ascertain the number of FOF samples needed to match serum samples at an assumed piglet-level prevalence. The results from this study also provide a framework for swine practitioners to estimate the likely proportion of viremic pigs in their barns, given the PRRSV-RT-rtPCR positivity rate of FOF samples submitted from a farrowing room.

**Keywords:** prevalence, simulations, FOF, clustering, PRRSV, piglets

**1. Introduction**

Porcine reproductive and respiratory syndrome virus (PRRSV) poses a significant challenge to the global swine industry (Holtkamp et al., 2013; Calderón Díaz et al., 2020).

Monitoring/surveillance remains an integral component of PRRSV control and elimination programs, and ascertaining the PRRSV status of pig populations around the time of weaning is crucial to guide decisions on health interventions and pig flow (Holtkamp et al., 2011). Efficient PRRSV surveillance/monitoring programs allow for the early detection of infection and helps evaluate changes in PRRSV prevalence over time; aiding swine producers and veterinarians alike to forestall the spread of PRRSV (Mccaw, 2000; Silva et al., 2017), and evaluate progress made with instituted PRRSV management programs (Linhares et al., 2014; Holtkamp et al., 2021).

Different sample types are routinely submitted to Veterinary diagnostic laboratories in the United States for PRRSV RT-rtPCR tests; these would include samples taken from individual pigs such as serum, swabs, semen, and post-mortem tissues; or aggregate samples taken from multiple pigs such as processing fluids and oral fluids (Trevisan et al., 2019). These samples are either submitted and tested individually or in pools.

The number of samples submitted for disease pathogen investigation is crucial to the success of a surveillance/monitoring exercise. Guided by epidemiological/statistical assumptions, the sample size should have enough power to detect at least one positive unit if the herd is truly positive for the pathogen of interest (Cameron et al., 2020; Stevenson, 2021).

Estimated prevalence at the individual pig level is one of the key variables used in calculating sample size to demonstrate the presence of a pathogen in a herd (Fosgate, 2009; Stevenson, 2021).

The diagnostic sample of choice for PRRSV surveillance in sow herds is serum from weaning-age pigs (Holtkamp et al., 2011). Although the serum sample is the sample of choice for PRRSV surveillance, it requires more skill, more manpower, is less animal welfare friendly, and is often impractical for frequent PRRSV surveillance in large herds (Turlewicz-Podbielska et al., 2020) compared to population-based sampling options. For these reasons, since 2018, validated aggregate samples have been the most frequently submitted samples for PRRSV surveillance in the US (Trevisan et al., 2019).

Almeida (Nunes de Almeida, 2020) demonstrated that, especially at low prevalence, family oral fluids (FOFs) are a more convenient and cost-efficient alternative to serum sampling for PRRSV surveillance in weaning-age pigs. A FOF sample is an aggregate sample obtained when oral fluids are wrung off a rope chewed by a sow and her suckling piglets (Almeida et al., 2020). A challenge with interpreting a positive result from testing FOF and other aggregate sample types is that one only knows that at least one animal that contributed to the sample matrix is pathogen-positive but cannot ascertain the exact number of pathogen-positive animals. Consequently, little to nothing is known about the number of positive pigs in a sampled room, given the proportion of PRRSV-positive aggregate samples, such as FOF, obtained from that room.

The individual pig is the unit for which sample size is calculated when non-aggregate samples are collected, while the litter is the unit for which sample size is calculated when an aggregate sample such as FOF is to be collected (Rotolo et al., 2017; Osemeke et al., 2022) it will be helpful to swine practitioners in making sampling decisions if they understood how the proportion of PRRSV-positive (viremic) piglets related with the proportion of PRRSV-positive litters, as both parameters are needed assumptions in estimating sample sizes.

To the best of our knowledge, the relationship between the piglet-level and litter-level prevalence in swine farrowing rooms has not been previously characterized. This study builds upon a previous study that assessed the probability of a positive FOF sample given the number of viremic PRRSV RT-rtPCR positive piglets within a litter (Almeida, Zhang, Lopez et al., 2021). We used an *in-silico* study to assess and model the relationship between the piglet-level prevalence (PP), true litter-level PRRSV prevalence (TLP) and apparent litter-level PRRSV prevalence (ALP) in a farrowing room. It is expected that the results from this study will not only provide insights to swine practitioners as to the likely relationship between PP, TLP, and ALP, but also provide a template for determining the piglet-level prevalence of PRRSV when monitored using aggregate samples.

**2. Methods**

**2.1 PRRSV detection in pig litters using FOF**

Based on a dataset from Almeida, Zhang, Zimmerman et al. (2021) 199 litters had all piglets sampled for PRRSV RNA detection by RT-rtPCR (reverse transcription real-time polymerase chain reaction); also, each litter (*i*=1,...,199) was sampled using FOF. The litters were sampled from 11 farrowing rooms across six different swine breeding farms (*j*=1,...,6).

The effect of the proportion of PRRSV-viremic piglets () in a litter on the detection of a positive litter using FOF () was assessed with a generalized linear mixed model employing a logit link function and a 'residual' Bernoulli distribution. In addition, the linear predictor comprised random effects for farms according to:

, (1)

where is the intercept of the model, is the random error assumed , and is the random effect accounting for the farm-effect in the model, assumed , where all are independent. Approximate maximum likelihood inference was based upon Laplacian integration, as implemented in R (R Core Team, 2018) in routine *glmer* from library *lme4* (Bates et al., 2015).

**2.2 Stochastic model**

The number of positive piglets in the *i-*thlitter ) is considered a random variable, and assuming that each piglet's status (positive/negative) is a Bernoulli trial, with a fixed *p* probability, arises from a binomial process. Consider a room with *n* litters with different sizes () drawn from a discrete empirical distribution, and total number of piglets in the room . In a simplistic scenario, the allocation of positive piglets in each litter ) would follow the relative size of the litter in the room. However, given that we are modeling an infectious disease, there might be situations where the total number of positive animals (*N*) may be "clustered" in a few litters (Carpenter, 2001; Kostoulas et al., 2013).

Accounting for this, the number of positive animals in each *i* litter () is calculated as a special case of the multinomial distribution, sampling recursively from binomial distributions using a clustering factor:

, (2)

where *j* stands for the successive allocation of positive animals within each litter, and is the probability of success in this binomial process defined as:

. (3)

The notation *c* represents a clustering factor. Thus when , the positive piglets will be totally clustered in the smallest number of litters as possible. On the other hand, when , piglets will be spread according to the relative size (number of piglets) of each litter regarding the room size.

To obtain the baseline clustering factor *c*, the observed distribution of the within litter prevalence reported in Almeida, Zhang, Zimmerman et al. (2021) across seven rooms, each room with *n* litters was used. The lost function was the minimization of the mean squared errors of the predicted (eq 2) *vs* observed distribution of the within litter prevalence . The objective function can be used to calculate a parameter estimate . Each room was randomly chosen, 10000 times obtaining the parameters ,, and . For each sampled room, 1000 acquisition points in the parameter space of *c* were sampled from a uniform distribution , obtaining a distribution to optimize *.*

2.2.2 *Apparent prevalence at the litter level*

The simulated proportions of positive piglets per litter obtained from 2.2 were used as input for the logistic model fit in 2.1, calculating the probability of detection of each simulated litter using FOF sampling. A random variable (*S*) was modeled describing the most probable number of positive litters detected in a routine FOF sampling in a farrowing room. Assuming the probability of each litter being detected by FOF ( see eq. 1) are independent of each other, and the positive/negative status of a litter , *S* equals . The expected apparent litter prevalence was obtained as . was generated a total of 2,000 times to improve the accuracy of the Monte Carlo estimation, and the mean was obtained and stored for that iterated room.

The parameters and distributions used in the simulations are described in table 1. In this simulation, 5000 stochastic iterations were performed, each one representing a different room, propagating the between litter variability observable in different farrowing rooms.

Table 1 – Descriptions of baseline model parameters used to compare the true and apparent liter prevalence of PRRSV.

|  |  |  |  |
| --- | --- | --- | --- |
| **Parameter/variable** | **Distribution/function** | **Description** | **Source** |
|  | Fixed= (range of values from 1% to 100%) | Probability of a piglet being positive in a room (prevalence) | Almeida |
|  |  | Total number of positive animals in the room | Calculation |
|  |  | Total number of piglets in the room | Calculation |
|  | empirical {(), ()} \* | Number of piglets in the *i-*thlitter | Almeida |
|  | Fixed=56 | Number of crates in a room | Authors’ opinion |
|  |  | Number of positive piglets in *i-*thlitter | Calculation |
|  |  | Probability of success in this binomial process (i.e., allocation of positive piglets in a litter) for the in *i-*thlitter | Calculation |
|  | Fixed=0.61 | Clustering factor | Optimized based on Almeida |

\* empirical {(3, 4, 5, 6, 7, 8, 9, 10 ,11, 12, 13, 14, 15, 25), (0.0092, 0.0092, 0.0046, 0.0046, 0.0553, 0.0691, 0.0922, 0.1014 0.1982, 0.2074, 0.1244, 0.0783, 0.0415, 0.0046)}

**2.3 Sensitivity analysis**

To assess the effect of the clustering factor (*c*) and the room size (*n*) on the estimated relationship between pig-level-prevalence and litter-level prevalence we selected five values for *c* (0.05, 0.34, 0.63, 0.83, 1) and five values for *n* (10, 33, 56, 79, 102) combining them as a factorial design for the sensitivity analysis, totaling 25 different scenarios.

**3. Results**

**3.1 PRRSV detection in pig litters using FOF**

The probability of PRRSV RNA detection in FOF by RT-rtPCR improved with an increasing proportion of PRRSV-positive piglets within a litter, staying above 95% from about 35% WLP (Fig 1, Table S2)

Chart, line chart

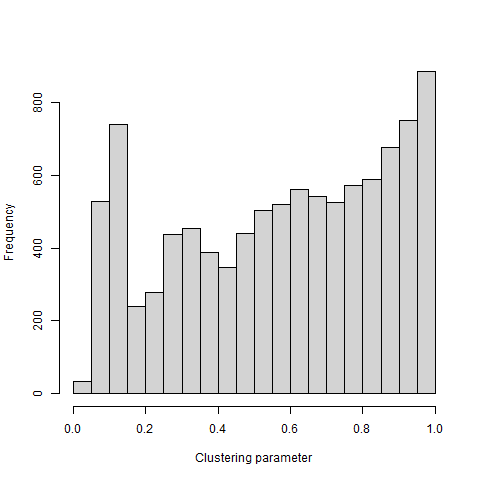
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*Figure 1: A jitter plot of the Probability of PRRSV RNA detection in FOF by the proportion of positive pigs within litters (within litter prevalence). 95% prediction intervals are represented by the grey region around the regression line.*

**3.2 Stochastic model**

**3.2.1 Observed distribution of clustering in sampled farms**

The clustering distribution across all sampled rooms had a minimum value of 0.00136, a median of 0.61, a mean of 0.57, and a maximum value of 1. The distributions of the clustering parameter across all sampled rooms are represented in figure 2.



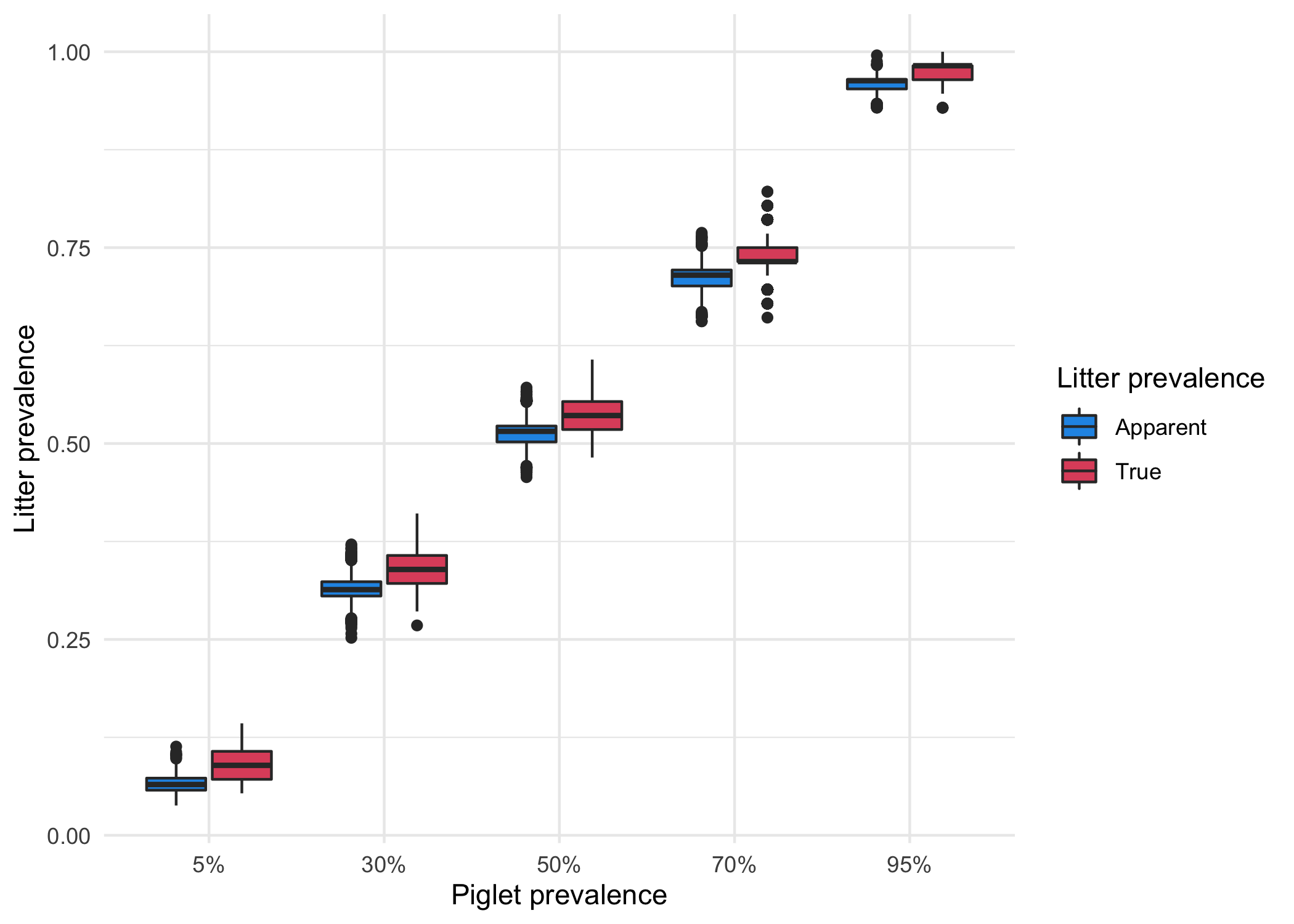
*Figure 2: A histogram showing the distribution of the clustering parameter across all sampled rooms from Almeida's study.*

**3.2.2 The relationship between piglet-level prevalence and litter-level prevalence**

Figure 3 and Table 2 show changes in median TLP and median ALP with increasing proportion of PRRSV-positive pigs in a 56-crate farrowing room considering a clustering factor of 0.61. When 1% of the piglets in the room are PRRSV-positive, about 5.36% of the 56 crates (~ 3 crates) are expected to have at least 1 PRRSV-positive piglet, and 2.06 % of the 56 crates (~1 crate) is expected to give a positive FOF.

*Table 2: Relationship between the proportion of positive piglets in a 56-crate farrowing room and the True and Apparent (by FOF) proportion of positive litters assuming a clustering level of 0.61.*

|  |  |  |
| --- | --- | --- |
| **Proportion of PRRSV-positive piglets**  **(%)** | **True proportion of PRRSV-positive litters (Upper and lower 95% quantiles)**  **(%)** | **Apparent proportion of PRRSV-positive litters by FOF (Upper and lower 95% quantiles)**  **(%)** |
| 1 | 5.36(1.79,7.14) | 2.06(1.07,3.53) |
| 5 | 8.93(7.14,12.50) | 6.48(5.30,8.58) |
| 10 | 14.29(10.71,17.86) | 11.25(9.31,13.92) |
| 15 | 19.64(16.07,23.21) | 16.35(14.47,19.21) |
| 20 | 23.21(21.43,26.79) | 21.60(18.73,24.19) |
| 25 | 28.57(25.00,32.14) | 26.66(23.50,29.31) |
| 30 | 33.93(30.36,37.50) | 31.35(28.77,34.33) |
| 35 | 39.29(35.71,42.86) | 36.16(33.49,39.44) |
| 40 | 44.64(41.07,48.21) | 41.30(38.05,44.71) |
| 45 | 48.21(44.64,53.57) | 46.54(43.10,49.68) |
| 50 | 53.57(50.00,57.14) | 51.56(48.34,54.58) |



*Figure 3: Distribution of True- and Apparent litter prevalence in a 56-crate room given different piglet-level prevalence scenarios and a clustering factor of 0.63.*

**3.3 Sensitivity analysis**

The sensitivity analysis was done to evaluate the effect of variations in clustering level and room size on the proposed relationship between piglet level prevalence and litter prevalence. The ALP was relatively more stable to changes in clustering and the number of crates compared to TLP. Generally, TLP and ALP increasingly converged to similar values with increasing clustering and increasing room size. Clustering changes appeared to have a more significant effect on ALP and TLP than changes in the number of crates in the room.

**Diagram, line chart

Description automatically generated**

*Figure 4: Graphical representation of changes in the relationship between the proportion of PRRSV-positive pigs and the proportion of PRRSV-positive litters (True and Apparent) with changes in clustering of PRRSV within room, and number of litters within rooms.*

**4. Discussion**

The use of mathematical models to describe disease dynamics in swine populations is not new. A few examples include the use of mathematical models to characterize and describe PRRSV transmission dynamics (Nodelijk et al., 2000; Evans et al., 2010; Amirpour Haredasht et al., 2017; Rotolo et al., 2017; Suksamran et al., 2017; Phoo-ngurn et al., 2019) and in the evaluation of PRRSV control strategies (Jeong et al., 2014; Arruda et al., 2016).

Earlier studies described the non-homogenous distribution of PRRSV in pig barns (M. N. Almeida et al., 2021; Rotolo et al., 2017). The non-homogenous aerial distribution of an infectious pathogen however is not limited to PRRSV alone (Carpenter, 2001; Kostoulas et al., 2013), and may be explained by PRRSV being highly infectious but not necessarily highly contagious (Pileri & Mateu, 2016), or by the mere fact that pigs in conventional US barns do not interact randomly with each other and are more likely to have direct contact with pigs within the same crate or with their closest neighbors (Murato et al., 2020).

Some popular statistical methods used in veterinary epidemiology for detecting and evaluating spatial (areal) clustering include Moran’s *I*, ohno, black-white, Geary’s *c*, and *I* pop (Carpenter, 2001), however, the use of the recursive binomial model in this study offered the authors a method to not only measure clustering, but to also propagate clustering in simulated data. The use of binomial models to detect and simulate clustering is also not new (Nauta, 2005; Li et al., 2018).

The restricted movement of pigs in conventional US swine barns and the non-homogenous distribution of viremic animals have been historically recognized to make conventional sample size assumptions (to detect a disease pathogen) not an exact fit; some previously proposed solutions include replacing simple random sampling with fixed spatial sampling (Rotolo et al., 2017), risk-based sampling (Almeida, Zhang, Zimmerman et al., 2021), or stratified sampling (Almeida, Zhang, Lopez et al., 2021). This study may well be one more step on adjusting conventional sampling schemes to better fit peculiarities with typical modern swine barns and with the ecology of PRRSV.

Clustering estimates the degree of homogeneity (or more aptly put; heterogeneity) of PRRSV between litters in a farrowing room; it may be overreaching to deterministically model a one-size-fits-all clustering for PRRSV. This is because the spread of PRRSV between litters within a farrowing room depends on a variety of factors, examples of which would include; 1) Management practices such as cross-fostering, and vaccination (Mccaw, 2000; Pileri and Mateu, 2016) 2) PRRSV strain (there is evidence of differences in characteristics such as virulence and spread between PRRSV strains) (Cho et al., 2007; Pileri and Mateu, 2016; Arruda et al., 2019; Ogno et al., 2019) 3) Barn structure (Rotolo et al., 2017) 4)Time since outbreak (Rotolo et al., 2017) 5) Secondary infections which may increase pig susceptibility to PRRSV, encourage huddling or increase the production of infectious respiratory fluids.

The uncertainty in definitively ascertaining clustering level however does not undermine the importance of these results or pose a challenge to its utilization, on the contrary, considering/estimating clustering adds some precision to the estimated prevalence guiding sample size calculations for disease pathogen surveillance (An example is given in *Table S3*).

The main goal of this study is to estimate the most likely relationship between the pig level prevalence and apparent litter prevalence by FOF, considering the pen-level sensitivity and specificity of this sample type. As observed from Figure 3, ALP is not as sensitive as TLP to variations in clustering parameter; we are therefore confident of the estimates on Table 2. One can also decide the number of crates or litters to randomly sample for FOF to detect PRRSV given an assumed piglet level prevalence. For example, assuming at least 10% piglet-level prevalence, serum sampling requires that about 30 pigs are sampled to be 95% confident of detecting at least one positive animal (Cannon and Roe, 1982; Holtkamp et al., 2011). From the table, 10% pig-level prevalence corresponds to about 11% ALP or about 7 litters in a 56-crate room likely to give a positive FOF test. This number can be used to calculate an appropriate sample size for FOF to detect at least 1 positive litter; Table 2 is, therefore, useful in estimating the litter prevalence from an assumed piglet-level prevalence.

Our approach to calculating ALP implicitly considers the diagnostic performance of FOF sampling; simply put, for a given piglet-level prevalence, the difference between the ALP and TLP is due to the diagnostic performance of FOF (the probability of RT-rtPCR testing of FOF samples to correctly assign PRRSV statuses to each tested litter). This implies that ALP can be used directly to estimate FOF sample size and the only diagnostic performance we may need to consider is that of the RT-rtPCR test kit.

Another key application of the proposed tables is to help the swine practitioner estimate piglet-level prevalence given the results of FOF testing. Given that a representative number of litters were sampled (sample size to estimate prevalence), the proportion of positive FOF results on RT-qPCR tests (apparent litter prevalence by FOF) can be used to deduce the likely proportion of viremic pigs (piglet-level prevalence). In Table S3, there were scenarios where the ALP was greater than the TLP, for example, the 100% clustering scenarios in the 56 and 102 crate rooms. This is because from the reference study, FOF from one of the sampled litters tested PRRSV-positive by Rt-rtPCR when there was no PRRSV-positive piglet (WLP = 0); consequently, in the predictive model used for the stochastic simulations, the probability of a positive FOF given that WLP is 0 greater than 0 (Table S2). A 100% clustering in the stochastic model restricts the distribution of PRRSV-positive pigs to the fewest number of litters possible (hereafter called PRRSV-positive litters), with a consequent maximization of the number of litters without any PRRSV-positive piglets (hereafter called PRRSV-negative litters). The probability of PRRSV-detection in FOF samples obtained from these PRRSV positive litters is almost always 100% owing to relatively high number of PRRSV-positive piglets “concentrated" within each of these litters. This should ordinarily put TLP and ALP at par, but ALP is further increased by the probability of a positive FOF RT-rtPCR result from the many (relatively) PRRSV-negative litters. The RT-rtPCR detection of PRRSV RNA in FOF from a PRRSV-negative litter may be an issue of test specificity, or may be explained by the sow that contributed to the FOF sample being PRRSV-positive and shedding (WLP does not consider the PRRSV status of the sow).

The referenced study (Almeida, Zhang, Lopez, et al., 2021) was not designed to answer the question of spatial distribution of viremic piglets within farrowing rooms, as such, in some sampled rooms, not every litter was sampled; consequently, the observed clustering values for those rooms may be inaccurate. To be able to deduce the number of viremic piglets from FOF positivity rate using the provided tables, it is important that one should have sampled the minimum number of litters needed to estimate prevalence.

**5. Conclusion**

This study explored the use of mathematical models to characterize the relationship between PP, TLP, and ALP in a farrowing room, this is a handy PRRSV surveillance tool for weaning age pigs in typical swine breeding herds in the United States. Prevalence comparisons here provided are intended to help guide sample choice and sample size for PRRSV surveillance in weaning age pigs. This study also demonstrate the effect of a clustering parameter on the characterized relationship between the mentioned proportions; like other sampling assumptions, clustering could be considered when estimating sample size. Further similar studies on other aggregate sample types, for other subpopulations and perhaps, for other pathogens will be helpful in guiding practitioners on how they can be up-to-speed with best practice surveillance as sampling methods evolve.

**6. Conflict of interest statement**

The authors declare that they have no conflicts of interest.

**7. Availability of data**

The simulations used to generate the figures and tables provided in this manuscript are available in the repository: https://github.com/onyechux/Prevalence-simulations

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|  |

**SUPPLEMENTARY MATERIALS**

*Table S1: A general description of the stochastic model with pictorial illustrations*



|  |  |  |  |
| --- | --- | --- | --- |
|  | **Goal** | **Example** | **Example in pictures** |
| 1 | **To create a farrowing room with *n* litters** by generating *n* random numbers from a discrete empirical distribution corresponding to the average litter size we observed from actual field data | From this empirical distribution, a possible set of random numbers generated will be 8, 9, 9, 7, 9, 8. These numbers are shown in the picture on the right. | A picture containing shape  Description automatically generated |
| 2. | **To create a disease prevalence scenario within this created room**.  Suppose we wanted a 10% prevalence. We will want 10% of the total pigs in the simulated room to be positive. | When we add all the litters (generated numbers), we have 52 pigs (8+9+9+7+9+8 = 50).  10% of 50 is 5, meaning five pigs will be PRRSV positive | A picture containing shape  Description automatically generated  Icon  Description automatically generated |
| 3 | **To distribute the diseased animals between pens in a manner typical of PRRSV**. PRRSV has been repeatedly reported to be heterogeneously distributed (clustered) within a farrowing room. A clustering factor in the recursive binomial model, which could range between 0 (homogenous distribution) and 1 (complete clustering), was used to assign positive pigs to litters.  The clustering factor determines the True Litter Prevalence (TLP), defined as the number of litters with at least one positive pig. | The first image shows what would be obtainable if there was no clustering (Clustering = 0).  **TLP** = 5/6  The next image shows what is expected in complete clustering (clustering = 1).  **TLP** = 1/6. | A picture containing text  Description automatically generated  A picture containing text  Description automatically generated |
| 4 | **To determine the expected number of positive FOF from this room if all the litters are tested**.  A predictive model is fitted using data from a previous study (Almeida, Zhang, Lopez et al., 2021). This model gives the probability of a positive FOF (pFOF) test given the proportion of positive pigs within a litter or within-litter prevalence (WLP).  After these probabilities (pFOFs) are generated, the Expected number of positive FOF tests for that room is determined using a Monte Carlo process. The Apparent litter prevalence by FOF (ALP) is then the expected number of positive FOF divided by the total number of litters or tests | The graph shows the relationship between pFOF and  WLP.  For the clustering = 0, the WLP and pFOF for each litter are calculated.  The expected number of positive FOF for that room is 0.2998 (<1).  **ALP** = <1/6  For the clustering = 1.  The expected number of positive FOF for that room is 1.014 (approximately 1).  **ALP** = 1/6 | |  |  |  | | --- | --- | --- | | WLP = 0.125  pFOF = 0.062 | WLP = 0.111  pFOF = 0.043 | WLP = 0  pFOF = 0.002 | | WLP = 0.143  pFOF = 0.099 | WLP = 0.111  pFOF = 0.043 | WLP = 0.111  pFOF = 0.043 |  |  |  |  | | --- | --- | --- | | WLP = 0  pFOF = 0.002 | WLP = 0.556  pFOF = 1.000 | WLP = 0  pFOF = 0.002 | | WLP = 0  pFOF = 0.002 | WLP = 0  pFOF = 0.002 | WLP = 0  pFOF = 0.002 | |
| 5  6 | **Repeat steps 1 to 4 (4,999 more times)** and obtain the median TLP andALP thereafter. The 5% prevalence is then matched with the median TLP and median ALP for the chosen clustering level.  Then repeat steps 1 to 5 for other prevalence scenarios |  |  |

*Table S2: The changes in the probability of PRRSV RNA RT-rtPCR detection in FOFs with increases in the proportion of PRRSV viremic piglets within a litter*

|  |  |  |
| --- | --- | --- |
| **SN** | **Proportion of PRRSV-positive pigs (WLP)** | **Probability of PRRSV RNA RT-rtPCR detection in FOF.** |
| 1 | 0.00% | 0.22% |
| 2 | 1.00% | 0.29% |
| 3 | 5.00% | 0.86% |
| 4 | 10.00% | 3.26% |
| 5 | 15.00% | 11.59% |
| 6 | 20.00% | 33.76% |
| 7 | 25.00% | 66.45% |
| 8 | 30.00% | 88.50% |
| 9 | 35.00% | 96.77% |
| 10 | 40.00% | 99.15% |
| 11 | 45.00% | 99.78% |
| 12 | 50.00% | 99.94% |
| 13 | 55.00% | 99.99% |
| 14 | 60.00% | 100.00% |
| 15 | 65.00% | 100.00% |
| 16 | 70.00% | 100.00% |
| 17 | 75.00% | 100.00% |
| 18 | 80.00% | 100.00% |
| 19 | 85.00% | 100.00% |
| 20 | 90.00% | 100.00% |
| 21 | 95.00% | 100.00% |
| 22 | 100.00% | 100.00% |

*Table S3: Relationship between PP, ALP, and TLP at different clustering levels and room sizes*

|  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- |
| **PP**  **(%)** | **Number of crates**  **(%)** | **Clustering**  **(%)** | **TLP**  **(%)** | **ALP**  **(%)** | **Number of litters with at least one PRRSV-positive piglet** | | **Number of litters expected to test PRRSV-positive by FOF sampling** |
| 1.00 | 10 | 5.00 | 10.00 | 0.43 | 1 | 0 | |
| 5.00 | 10 | 5.00 | 40.00 | 3.55 | 4 | 0 | |
| 10.00 | 10 | 5.00 | 70.00 | 14.24 | 7 | 1 | |
| 15.00 | 10 | 5.00 | 80.00 | 27.20 | 8 | 3 | |
| 20.00 | 10 | 5.00 | 90.00 | 40.68 | 9 | 4 | |
| 25.00 | 10 | 5.00 | 90.00 | 52.83 | 9 | 5 | |
| 30.00 | 10 | 5.00 | 100.00 | 63.14 | 10 | 6 | |
| 35.00 | 10 | 5.00 | 100.00 | 71.21 | 10 | 7 | |
| 40.00 | 10 | 5.00 | 100.00 | 78.01 | 10 | 8 | |
| 45.00 | 10 | 5.00 | 100.00 | 83.62 | 10 | 8 | |
| 50.00 | 10 | 5.00 | 100.00 | 87.88 | 10 | 9 | |
| 1.00 | 10 | 33.00 | 10.00 | 0.45 | 1 | 0 | |
| 5.00 | 10 | 33.00 | 30.00 | 8.48 | 3 | 1 | |
| 10.00 | 10 | 33.00 | 50.00 | 19.49 | 5 | 2 | |
| 15.00 | 10 | 33.00 | 50.00 | 27.50 | 5 | 3 | |
| 20.00 | 10 | 33.00 | 60.00 | 32.27 | 6 | 3 | |
| 25.00 | 10 | 33.00 | 60.00 | 38.56 | 6 | 4 | |
| 30.00 | 10 | 33.00 | 70.00 | 42.51 | 7 | 4 | |
| 35.00 | 10 | 33.00 | 70.00 | 48.40 | 7 | 5 | |
| 40.00 | 10 | 33.00 | 70.00 | 52.21 | 7 | 5 | |
| 45.00 | 10 | 33.00 | 80.00 | 58.21 | 8 | 6 | |
| 50.00 | 10 | 33.00 | 80.00 | 61.85 | 8 | 6 | |
| 1.00 | 10 | 61.00 | 10.00 | 0.45 | 1 | 0 | |
| 5.00 | 10 | 61.00 | 20.00 | 10.31 | 2 | 1 | |
| 10.00 | 10 | 61.00 | 30.00 | 17.35 | 3 | 2 | |
| 15.00 | 10 | 61.00 | 30.00 | 20.87 | 3 | 2 | |
| 20.00 | 10 | 61.00 | 40.00 | 27.25 | 4 | 3 | |
| 25.00 | 10 | 61.00 | 40.00 | 30.81 | 4 | 3 | |
| 30.00 | 10 | 61.00 | 50.00 | 37.26 | 5 | 4 | |
| 35.00 | 10 | 61.00 | 50.00 | 40.65 | 5 | 4 | |
| 40.00 | 10 | 61.00 | 60.00 | 46.87 | 6 | 5 | |
| 45.00 | 10 | 61.00 | 60.00 | 50.57 | 6 | 5 | |
| 50.00 | 10 | 61.00 | 70.00 | 56.75 | 7 | 6 | |
| 1.00 | 10 | 100.00 | 10.00 | 0.46 | 1 | 0 | |
| 5.00 | 10 | 100.00 | 10.00 | 10.19 | 1 | 1 | |
| 10.00 | 10 | 100.00 | 10.00 | 10.23 | 1 | 1 | |
| 15.00 | 10 | 100.00 | 20.00 | 20.15 | 2 | 2 | |
| 20.00 | 10 | 100.00 | 20.00 | 20.22 | 2 | 2 | |
| 25.00 | 10 | 100.00 | 30.00 | 30.13 | 3 | 3 | |
| 30.00 | 10 | 100.00 | 30.00 | 30.19 | 3 | 3 | |
| 35.00 | 10 | 100.00 | 40.00 | 40.11 | 4 | 4 | |
| 40.00 | 10 | 100.00 | 40.00 | 40.17 | 4 | 4 | |
| 45.00 | 10 | 100.00 | 50.00 | 50.09 | 5 | 5 | |
| 50.00 | 10 | 100.00 | 50.00 | 50.14 | 5 | 5 | |
| 1.00 | 56 | 5.00 | 10.71 | 0.64 | 6 | 0 | |
| 5.00 | 56 | 5.00 | 35.71 | 7.35 | 20 | 4 | |
| 10.00 | 56 | 5.00 | 50.00 | 18.28 | 28 | 10 | |
| 15.00 | 56 | 5.00 | 58.93 | 27.65 | 33 | 15 | |
| 20.00 | 56 | 5.00 | 66.07 | 34.97 | 37 | 20 | |
| 25.00 | 56 | 5.00 | 71.43 | 41.26 | 40 | 23 | |
| 30.00 | 56 | 5.00 | 75.00 | 46.64 | 42 | 26 | |
| 35.00 | 56 | 5.00 | 78.57 | 51.69 | 44 | 29 | |
| 40.00 | 56 | 5.00 | 82.14 | 56.51 | 46 | 32 | |
| 45.00 | 56 | 5.00 | 85.71 | 61.27 | 48 | 34 | |
| 50.00 | 56 | 5.00 | 87.50 | 66.01 | 49 | 37 | |
| 1.00 | 56 | 33.00 | 7.14 | 1.75 | 4 | 1 | |
| 5.00 | 56 | 33.00 | 14.29 | 7.92 | 8 | 4 | |
| 10.00 | 56 | 33.00 | 19.64 | 13.02 | 11 | 7 | |
| 15.00 | 56 | 33.00 | 23.21 | 18.04 | 13 | 10 | |
| 20.00 | 56 | 33.00 | 28.57 | 23.13 | 16 | 13 | |
| 25.00 | 56 | 33.00 | 33.93 | 27.93 | 19 | 16 | |
| 30.00 | 56 | 33.00 | 39.29 | 32.86 | 22 | 18 | |
| 35.00 | 56 | 33.00 | 42.86 | 37.92 | 24 | 21 | |
| 40.00 | 56 | 33.00 | 48.21 | 42.97 | 27 | 24 | |
| 45.00 | 56 | 33.00 | 53.57 | 47.89 | 30 | 27 | |
| 50.00 | 56 | 33.00 | 58.93 | 52.86 | 33 | 30 | |
| 1.00 | 56 | 61.00 | 5.36 | 2.06 | 3 | 1 | |
| 5.00 | 56 | 61.00 | 8.93 | 6.45 | 5 | 4 | |
| 10.00 | 56 | 61.00 | 14.29 | 11.27 | 8 | 6 | |
| 15.00 | 56 | 61.00 | 19.64 | 16.34 | 11 | 9 | |
| 20.00 | 56 | 61.00 | 23.21 | 21.61 | 13 | 12 | |
| 25.00 | 56 | 61.00 | 28.57 | 26.62 | 16 | 15 | |
| 30.00 | 56 | 61.00 | 33.93 | 31.36 | 19 | 18 | |
| 35.00 | 56 | 61.00 | 39.29 | 36.18 | 22 | 20 | |
| 40.00 | 56 | 61.00 | 44.64 | 41.28 | 25 | 23 | |
| 45.00 | 56 | 61.00 | 48.21 | 46.54 | 27 | 26 | |
| 50.00 | 56 | 61.00 | 53.57 | 51.62 | 30 | 29 | |
| 1.00 | 56 | 100.00 | 1.79 | 2.00 | 1 | 1 | |
| 5.00 | 56 | 100.00 | 5.36 | 5.57 | 3 | 3 | |
| 10.00 | 56 | 100.00 | 10.71 | 10.91 | 6 | 6 | |
| 15.00 | 56 | 100.00 | 16.07 | 16.23 | 9 | 9 | |
| 20.00 | 56 | 100.00 | 21.43 | 20.42 | 12 | 11 | |
| 25.00 | 56 | 100.00 | 25.00 | 25.18 | 14 | 14 | |
| 30.00 | 56 | 100.00 | 30.36 | 30.52 | 17 | 17 | |
| 35.00 | 56 | 100.00 | 35.71 | 35.85 | 20 | 20 | |
| 40.00 | 56 | 100.00 | 41.07 | 41.18 | 23 | 23 | |
| 45.00 | 56 | 100.00 | 46.43 | 45.19 | 26 | 25 | |
| 50.00 | 56 | 100.00 | 50.00 | 50.13 | 28 | 28 | |
| 1.00 | 102 | 5.00 | 9.80 | 0.91 | 10 | 1 | |
| 5.00 | 102 | 5.00 | 29.41 | 8.75 | 30 | 9 | |
| 10.00 | 102 | 5.00 | 39.22 | 18.14 | 40 | 19 | |
| 15.00 | 102 | 5.00 | 46.08 | 24.96 | 47 | 25 | |
| 20.00 | 102 | 5.00 | 50.98 | 30.44 | 52 | 31 | |
| 25.00 | 102 | 5.00 | 55.88 | 35.47 | 57 | 36 | |
| 30.00 | 102 | 5.00 | 60.78 | 40.36 | 62 | 41 | |
| 35.00 | 102 | 5.00 | 64.71 | 45.31 | 66 | 46 | |
| 40.00 | 102 | 5.00 | 68.63 | 50.15 | 70 | 51 | |
| 45.00 | 102 | 5.00 | 73.53 | 54.99 | 75 | 56 | |
| 50.00 | 102 | 5.00 | 77.45 | 59.81 | 79 | 61 | |
| 1.00 | 102 | 33.00 | 4.90 | 2.07 | 5 | 2 | |
| 5.00 | 102 | 33.00 | 9.80 | 6.85 | 10 | 7 | |
| 10.00 | 102 | 33.00 | 14.71 | 11.85 | 15 | 12 | |
| 15.00 | 102 | 33.00 | 19.61 | 16.82 | 20 | 17 | |
| 20.00 | 102 | 33.00 | 24.51 | 21.76 | 25 | 22 | |
| 25.00 | 102 | 33.00 | 30.39 | 26.75 | 31 | 27 | |
| 30.00 | 102 | 33.00 | 35.29 | 31.67 | 36 | 32 | |
| 35.00 | 102 | 33.00 | 40.20 | 36.66 | 41 | 37 | |
| 40.00 | 102 | 33.00 | 45.10 | 41.68 | 46 | 43 | |
| 45.00 | 102 | 33.00 | 50.00 | 46.69 | 51 | 48 | |
| 50.00 | 102 | 33.00 | 54.90 | 51.68 | 56 | 53 | |
| 1.00 | 102 | 61.00 | 2.94 | 1.99 | 3 | 2 | |
| 5.00 | 102 | 61.00 | 6.86 | 6.04 | 7 | 6 | |
| 10.00 | 102 | 61.00 | 11.76 | 10.99 | 12 | 11 | |
| 15.00 | 102 | 61.00 | 17.65 | 15.90 | 18 | 16 | |
| 20.00 | 102 | 61.00 | 22.55 | 20.82 | 23 | 21 | |
| 25.00 | 102 | 61.00 | 27.45 | 25.77 | 28 | 26 | |
| 30.00 | 102 | 61.00 | 32.35 | 30.74 | 33 | 31 | |
| 35.00 | 102 | 61.00 | 37.25 | 35.70 | 38 | 36 | |
| 40.00 | 102 | 61.00 | 42.16 | 40.81 | 43 | 42 | |
| 45.00 | 102 | 61.00 | 47.06 | 45.89 | 48 | 47 | |
| 50.00 | 102 | 61.00 | 51.96 | 50.88 | 53 | 52 | |
| 1.00 | 102 | 100.00 | 0.98 | 1.21 | 1 | 1 | |
| 5.00 | 102 | 100.00 | 5.88 | 5.14 | 6 | 5 | |
| 10.00 | 102 | 100.00 | 10.78 | 10.31 | 11 | 11 | |
| 15.00 | 102 | 100.00 | 15.69 | 15.68 | 16 | 16 | |
| 20.00 | 102 | 100.00 | 20.59 | 20.74 | 21 | 21 | |
| 25.00 | 102 | 100.00 | 25.49 | 25.65 | 26 | 26 | |
| 30.00 | 102 | 100.00 | 30.39 | 30.54 | 31 | 31 | |
| 35.00 | 102 | 100.00 | 35.29 | 35.44 | 36 | 36 | |
| 40.00 | 102 | 100.00 | 40.20 | 40.33 | 41 | 41 | |
| 45.00 | 102 | 100.00 | 45.10 | 45.23 | 46 | 46 | |
| 50.00 | 102 | 100.00 | 50.00 | 50.12 | 51 | 51 | |