Research Article

Serodiagnosis and Risk Factors Associated with Infectious Agents of Reproductive Diseases in Bovines of Chiquinquirá, District of Boyacá (Colombia)

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Received 22 April 2022; Revised 8 June 2022; Accepted 1 July 2022; Published 16 July 2022

Academic Editor: Remo Lobetti

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The productivity of cattle farms is affected by infectious and noninfectious factors that generate economic losses and cause reproductive failure represented by low conception rates, embryonic mortality, abortions, and fetal mummification. The infectious agents that most impact the reproductive health of the bovine species from conception to birth are bovine herpesvirus type 1 (BoHV-1) causing infectious bovine rhinotracheitis (IBR), bovine viral diarrhea virus (BVDV), bovine parainfluenza virus type 3 (PI3), Neospora caninum, and Leptospira spp. The objective of this study was to diagnose the presence of BoHV-1, bovine viral diarrhea (BVD), PI3, Neospora caninum, and Leptospira spp. by serology and identify the risk factors associated with infectious agents of reproductive interest in bovines of Boyacá (Colombia). A descriptive cross-sectional study was developed, with simple random sampling, where a sample size of 601 female cattle of Holstein, Jersey, and Normande breeds of different age groups was determined. Blood samples were taken and processed using the indirect ELISA technique (SYNBIOTICS®, SERELISA® BVD p80 Ab Mono Blocking, Ingezim R.12.NC.K, PRIMACHECK VPI-3®) and the MAT test for the diagnosis of bovine leptospirosis. The data were processed with the statistical program Epi Info™. The highest apparent seroprevalence was established for infectious bovine rhinotracheitis (61.1%), followed by BVD (37.6%), PI3 (40.9%), neosporosis (51.1%), and leptospirosis (14.8%). Variables such as age > 4 years and Holstein breed for IBR and >4 years for BVD were established risk factors. Considering our results, we suggest implementing prevention and control plans that include vaccination as a prophylactic measure and biosecurity tools that reduce the probability of contagion and transmission of pathogens.

1. Introduction

The sustainability of a livestock farm begins with an adequate state of animal health, which depends mainly on the effective management of the multiple pathogens found in the farm [1]. In particular, the control of etiological agents of infectious diseases that generate significant negative impacts on the reproductive efficiency of cattle, both in the beef and dairy industry, as well as concomitant problems for human health and the environment, is due to the use of antimicrobials as a treatment for reproductive pathologies [2,3].

Infectious agents such as bacteria, viruses, protozoa, chlamydiae, and fungi are known to directly impact the reproductive health of cattle [4]. Bovine herpesvirus type 1 (BoHV-1), bovine viral diarrhea virus (BVDV), bovine parainfluenza virus type 3 (PI3), Neospora caninum, and Leptospira spp. are considered the main causative agent of reproductive disturbances in cattle around the world. Resulting in the presentation of clinical manifestations generated by the diseases infectious bovine rhinotracheitis (IBR), bovine viral diarrhea (BVD), bovine parainfluenza, bovine type 3 (PI3), neosporosis, and bovine leptospirosis, respectively [5–8]. The reproductive consequences of
infection in an animal vary from outbreaks of abortion that can affect a large proportion of the pregnant herd to more subtle symptoms such as calving intervals, decrease in conception rate, and early embryonic mortality, among others that go unnoticed or undiagnosed [2]. The economic losses derived from reproductive events caused by infectious agents are high; therefore, proper reproductive efficiency management is essential for any livestock farm’s economic success [9].

Considering the above impacts, prevention and control practices, preventive vaccination tools, and effective and efficient treatment protocols form the basis for preparing the immune system of each animal for possible exposure to pathogens on livestock farms. However, the implementation of other strategies focused on improving collective immunity through external factors such as nutrition, environment, and genetics also play a significant role in the management of animal production [1].

Based on the best available scientific information, effective recognition, prevention, and treatment of reproductive disorders are essential in the practice of veterinary professionals and farm owners. Thus, prevention and control protocols must be based on diagnostic tests that fit into the overall biosecurity production programs and must have measures to prevent pathogens’ introduction and spread [2,3]. Therefore, the objective of this study was to evaluate serological evidence for the presence of infections caused by BoHV-1, BVD, PI3, Neospora caninum, and Leptospira spp. and to identify risk factors potentially associated with such infectious agents in cattle in Boyacá (Colombia).

2. Materials and Methods

2.1. Study Area. The study was conducted in the town of Chiquinquirá, located in the west of the department of Boyacá (latitude 5°36′48″ north, longitude 0°15′21″ meridian of Bogotá, 2000 to 3200 m.a.s.l, and mean temperature of 15°C). The township is the economic and commercial center of the western region of the department, where the agricultural sector relies on milk production and its derivatives [10].

2.2. Sample Size. The livestock census conducted by the Colombian Agricultural Institute - ICA reported that for the year 2020, Chiquinquirá had 31082 heads of cattle [11]. As a result, a sample size of 601 unvaccinated animals was determined with a confidence interval of 95% (CI) and an accepted error of 4% implementing the following equation:

\[ n = \left( \frac{Z_{a/2} \sqrt{p(1-p)}}{E} \right)^2 \]

where \( n \) = sample size, \( E \) = accepted error, \( p \) = expected value of the proportion, and \( \alpha \) = tail probability. Parallel to the sampling, an epidemiological survey was designed to collect information such as the age group and the breed of the animal and those reproductive events related to the diseases evaluated.

2.3. Sample Collection and Processing. Blood samples were collected by puncturing the coccygeal vein using the multiterple needle gauge no. 21 x 1 (Vacutainer®). Seven ml of biological material was collected and deposited in the BD Vacutainer® tubes without anticoagulant. The samples were refrigerated (4°C) and transported to the Veterinary Parasitology Laboratory of the Universidad Pedagógica y Tecnológica de Colombia. In the laboratory, the blood samples were centrifuged at 2500 r.p.m. for 10 minutes to separate the cells from the serum. Then, with a Pasteur pipette, the serum or the supernatant was transferred to the 1.5 ml Eppendorf type storage tube for storage at −20°C until testing [12].

An indirect enzyme-linked immunosorbent assay (ELISA) for the detection of specific antibodies against IBR, BVD, PI3, and neosporosis was carried out, using the commercial kits SYNBIOTICS® (sensitivity 96%; specificity 98%), SERELISA® BVD p80 Ab Mono Blocking, (sensitivity 98%; specificity 100%), PRIMACHEK VPI-3® (sensitivity 97%; specificity 99%), and Ingezim R.12NC.K1 (Ingenasa S.A) (sensitivity 70%; specificity 100%), by following the manufacturer’s protocol. In addition, leptospirosis was diagnosed by the microscopic agglutination test (MAT); animals were considered to be positive when titers were ≥ 1: 100 (sensitivity 60%; specificity 100%).

2.4. Statistical Analysis. An observational, descriptive, cross-sectional study was carried out with a simple random sampling of animals, where the sampling unit was female cattle. The apparent prevalence (AP) and the real prevalence (RP) were determined with the WinEpi statistical program. After consolidating and filtering the database, the data were analyzed using statistical program Epi Info® version 7.2.4.0. To estimate the prevalence ratio (PR), the proportion of animals and herds affected by the disease exposed to a factor was compared to the same proportion of the population not exposed to that factor. This PR was used to measure the association between reproductive agents and hypothesized causal factors, as well as the significance of these associations using Fisher’s exact test [13]. The dependent variable included the serological results obtained; the independent variables were all considered in the epidemiological study. Once these factors were established, logistic regression was performed to establish possible correlations.

2.5. Ethical Considerations. The study was conducted under the laws 576 of 2000 and 84 of 1989 of the Republic of Colombia. Informed consent was obtained from the owners of the specimens prior to sample collection.

3. Results

The apparent prevalence (AP) and the real prevalence (RP) of the infectious diseases were determined where seropositivity was 61.1% and 62.9% for IBR, respectively, (positive predictive value (PP+) 98.8% and negative predictive value (NP-) 93.8%), 37.6% and 38.4% for BVD, 40.9% and 41.6%
for PI3, 51.1% and 73% for neosporosis, and 14.8% and 24.7% for bovine leptospirosis, respectively, (Table 1).

Regarding the age groups evaluated, individuals >4 years old presented the highest seroprevalence for IBR and BVD (68.9% and 50.6%, respectively). In contrast, for PI3, neosporosis, and leptospirosis, cattle 1–4 years old presented the highest seropositivity (Table 1). Concerning the breeds evaluated, Holstein presented the highest seroprevalence to IBR and PI3 and Jersey presented the highest seropositivity to N. caninum, while BVD and Leptospira spp. were more prevalent in cattle of the Normande breed (Table 2).

Statistical significance was established between the breeds and age groups evaluated through the presence of antibodies against IBR. Likewise, the presence of BVD was associated with the age groups of the cattle. Furthermore, a significant statistical association was determined between seropositivity to the disease and the reproductive events taken into account in the epidemiological survey. Cattle >4 years old, Holstein breed, AI, fetal death, cows with a history of inseminations without reproductive success, and non-certified semen were considered possible risk factors for IBR. In addition, individuals older than four years were established as possible risk factors for BVD occurrence. Natural mating and weak calves at birth presented the same condition but for neosporosis seropositivity (Table 3).

Logistic regression analysis identified that the age group >4 years and the Holstein breed were risk factors for IBR; furthermore, bovines with a history of fetal death in Table 1: Prevalence of IBR, BVD, PI3, neosporosis, and leptospirosis in cattle in Chiquinquirá, Boyacá.

| Disease               | No of animals tested | Positive | AP (%) | RP (%) | PP+ (%) | NP - (%) |
|-----------------------|----------------------|----------|--------|--------|---------|----------|
| IBR (BoHV-1)          | 601                  | 367      | 61.1   | 62.9   | 98.8    | 93.8     |
| BVD                   | 601                  | 225      | 37.6   | 38.4   | 100     | 98.8     |
| PI3                   | 601                  | 245      | 40.9   | 41.6   | 98.6    | 97.9     |
| Neosporosis           | 601                  | 307      | 51.1   | 73     | 100     | 55.2     |
| Leptospirosis         | 601                  | 89       | 14.8   | 24.7   | 100     | 88.4     |

Table 2: Prevalence of IBR, BVD, PI3, neosporosis, and leptospirosis by the age group and the breed in cattle in Chiquinquirá, Boyacá.

| Disease               | Category   | No of animals tested | Positive | Seropositive (%) |
|-----------------------|------------|----------------------|----------|------------------|
| IBR (BoHV-1)          | <1 year    | 106                  | 54       | 50.9             |
|                       | 1–4 years  | 183                  | 98       | 53.6             |
|                       | >4 years   | 312                  | 215      | 68.9             |
| BVD                   | <1 year    | 106                  | 19       | 17.9             |
|                       | 1–4 years  | 183                  | 49       | 26.8             |
|                       | >4 years   | 312                  | 158      | 50.6             |
| PI3                   | <1 year    | 106                  | 43       | 40.6             |
|                       | 1–4 years  | 183                  | 80       | 43.7             |
|                       | >4 years   | 312                  | 123      | 39.4             |
| Neosporosis           | <1 year    | 106                  | 51       | 48.1             |
|                       | 1–4 years  | 183                  | 101      | 55.2             |
|                       | >4 years   | 312                  | 155      | 49.7             |
| Leptospirosis         | <1 year    | 106                  | 11       | 10.4             |
|                       | 1–4 years  | 183                  | 32       | 17.5             |
|                       | >4 years   | 312                  | 46       | 14.7             |
| Breed                 |            |                      |          |                  |
| IBR (BoHV-1)          | Holstein   | 427                  | 282      | 66.0             |
|                       | Jersey     | 60                   | 29       | 48.3             |
|                       | Normande   | 114                  | 54       | 47.4             |
| BVD                   | Holstein   | 427                  | 160      | 37.5             |
|                       | Jersey     | 60                   | 22       | 36.7             |
|                       | Normande   | 114                  | 44       | 38.6             |
| PI3                   | Holstein   | 427                  | 179      | 41.9             |
|                       | Jersey     | 60                   | 20       | 33.3             |
|                       | Normande   | 114                  | 47       | 41.2             |
| Neosporosis           | Holstein   | 427                  | 211      | 49.4             |
|                       | Jersey     | 60                   | 35       | 58.3             |
|                       | Normande   | 114                  | 61       | 53.5             |
| Leptospirosis         | Holstein   | 427                  | 71       | 16.6             |
|                       | Jersey     | 60                   | 7        | 11.7             |
|                       | Normande   | 114                  | 21       | 18.4             |
pregnancy, cattle with a history of inseminations without reproductive success, and dairy cows pregnant with uncertified semen could have 2.4383, 2.1222, and 2.9633 possibilities of being positive for the disease, respectively. For BVD, individuals older than 4 years were considered risk factors for the disease. On the other hand, cattle that are pregnant through natural mating and weak calves at birth may have 1.8936 and 2.6266 more possibilities of being seropositive for *Neospora* (Table 4).

### 4. Discussion

In this study, the disease with the highest seropositivity in the individuals evaluated was BoHV-1. Seroprevalences

| Table 3: Possible risk factors associated with IBR, BVD, PI3, neosporosis, and leptospirosis infections and results are shown as the prevalence ratio (PR) and the 95% confidence interval (95% CI). Significance is denoted by a value of \( p < 0.05 \). |
| --- |
| **Disease** | **Variable** | **Category** | **PR** | **CI (95%)** | **p value** |
| IBR (BoHV-1) | Age group | <1 year | 0.7495 | 0.5980–0.9394 | 0.012894936 |
| IBR (BoHV-1) | 1–4 years | 0.7674 | 0.6271–0.9394 | 0.008220621 |
| IBR (BoHV-1) | >4 years | 1.5248 | 1.2421–1.8718 | 0.000289377 |
| IBR (BoHV-1) | Breed | Holstein | 1.5617 | 1.2850–1.8978 | 0.000153039 |
| IBR (BoHV-1) | Jersey | 1.7263 | 0.5556–0.9493 | 0.02413266 |
| IBR (BoHV-1) | Normande | 0.6789 | 0.5497–0.8383 | 0.00069915 |

| Table 4: Analysis of different variables as possible risk factors for seropositivity to IBR, BVD, PI3, neosporosis, and leptospirosis infections in cattle in Chiquinquirá, Boyacá. |
| --- |
| **Disease** | **Variable** | **OR** | **Lower confidence interval (95%CI)** | **Upper confidence interval (95% CI)** | **p value** |
| IBR | >4 years | 1.8185 | 1.2845 | 2.5746 | 0.0007 |
| IBR | Holstein | 1.4936 | 0.9572 | 2.3304 | 0.0772 |
| IBR | IA | 0.7294 | 0.4425 | 1.2025 | 0.2161 |
| IBR | Fetal death | 2.4383 | 1.5021 | 3.9581 | 0.0003 |
| IBR | Cows with a history of inseminations without reproductive success | 2.1222 | 1.3943 | 3.2301 | 0.0004 |
| IBR | Uncertified semen | 2.9633 | 1.6967 | 5.1755 | 0.0001 |
| BVD | >4 years | 3.3344 | 2.3476 | 4.736 | 0.0002 |
| Neosporosis | Natural mating | 1.8936 | 1.3109 | 2.7353 | 0.0007 |
| Neosporosis | Weak calves at birth | 2.6266 | 1.661 | 4.1537 | 0.0005 |
ranging from 35.65% to 73.13% are reported in Colombia [14–18]. On the other hand, diagnosis by serology for Neospora indicated that 51.1% of the cattle evaluated were seropositive, and these results are similar to national-level seroprevalences ranging from 2.8% to 57.5% [15,19–26]. Likewise, it has been reported that 40.9% of cattle had antibodies against P13, a value that is in agreement with other studies performed in Colombia, which are between 11.20% and 88% [15,27–30]. It was demonstrated in this study that there is a wide distribution of seropositivity to BoHV-1, BVDV, P13, leptospirosis, and neosporosis in Chiquinquirá, accompanied by a high transmission of these pathogens.

Regarding BVD, the values determined in the present study are 37.6%, which are between the ranges reported for the disease, which range from 29.7% to 76.4% [8,15,16,21,23,31]. Considering the seroprevalence determined for leptospirosis, 14.8% is a lower value than those established nationally, between 23.1% and 74.5% for the disease [16,20,23,32–34]. The findings are considered due to the risk of animal infection by Leptospira spp. associated with environmental factors such as rainfall and adverse climate, management factors, and farming systems such as cobeeding with other productive species or the presence of domestic animals (canines) and wild animals (rodents) in the herd [35].

Similarly, it is relevant to note that cattle are commonly grouped in herds, where prevalence estimates depend on the sampling strategy; thus, the presence of antibodies may vary from one investigation to another [36]. Hence, a limitation of this type of study is the ELISA that is intended for use as an aid in identifying animals with an adaptive immune response, indicating recent or prior infection. This has been reported by Kipyego et al. [37] in bovine herpesvirus type 1; however, the cattle sampled had no history of vaccination.

Cattle older than four years presented the highest seropositivity to BVDV, which is in agreement with previous studies. González-Bautista et al. [8] found the highest seroprevalence for this same age group in Sotaquirá. In Chiquinquirá, a significant statistical association was found between IBR seropositivity, breeds, and age groups evaluated, which differs from those reported by Doria-Ramos et al. [30], but is in agreement with the study conducted by Kipyego et al. [37], who reported a positive correlation with aging of cattle, where the odds of older dairy cattle being BoHV-1 antibody positivity which were 1,200 times higher with each additional year of age. Likewise, Doria-Ramos et al. and González-Bautista et al. [8,30] reported a statistically significant correlation between aging and certain breeds with seropositivity to BVDV, which is similar to the findings reported in this study.

Regarding P13, neosporosis, and leptospirosis, cattle aged 1–4 years presented the highest seropositivity, which is not in agreement with the studies by Betancur et al. [27], Betancur et al. [32], Cardona et al., and Bedoya et al. [22,25], who found the highest seroprevalences for P13, leptospirosis, and neosporosis, respectively, in older cattle, and this may occur because infected cattle remain infected for life; this effect is likely to represent the risk of cumulative exposure to the agent in the environment as the animals’ age [37,38].

Holstein and Jersey cattle are the breeds that had the highest amount of antibodies against P13 and neosporosis, which does not agree with Fernandez et al. [29], who indicated that the most prevalent breed for P13 was Jersey and with the higher seropositivity of the Holstein breed found by Cruz-Estupiñan et al. [26]; however, this can occur due to the different management practices implemented on the farms and because these are dairy herds. No significant statistical association was obtained between P13, leptospirosis, and neosporosis with the breed and age group of the cattle evaluated. This agrees with the findings of Fernandez et al. [29], Betancur et al. [32], Cruz-Estupiñan et al. [26], and Ansari-lari [7] but differs with Bedoya et al. [25], who found an association with age and established it as a risk factor for infection with Neospora. Doria-Ramos et al. [30] reported a relationship between the breed and age with the presentation of P13; however, in Chiquinquirá, no association was found between seropositivity for antibodies against P13 and the described variables.

In the present study, artificial insemination (AI) showed a significant association with seropositivity to BoHV-1 and leptospirosis. The transmission of pathogens occurs when there are no prior preventive measures in reproductive technologies; hence, mitigation of this risk should be considered during their development. Likewise, implementing biosecurity protocols and excellent quality control is imperative to prevent contamination and transmission of diseases while using these tools [39,40]. The quality of semen used to inseminate cattle is associated with the different diseases evaluated here. The use of noncertified semen, that is, semen freedom of these agents might become a risk factor for IBR, as previously reported [41–44]. There is significant evidence that these infections have other impacts on dairy cows, which are reflected in reduced conception rates, which is why the cows with a history of inseminations without reproductive success were considered as a risk factor for IBR appearance in the females evaluated.

Dystocic parturition, natural mating, fetal death, and weak calves at birth presented a significant statistical association with the seropositivity to neosporosis antibodies (p ≤ 0.05), and bovines with a history of fetal death in pregnancy, cattle with a history of inseminations without reproductive success, and dairy cows pregnant with uncertified semen have possibilities of being positive for the IBR. This is associated with what was reported by de Barros et al. [45] who mentioned that the correlations between the reproductive variables of seronegative animals were normal, while these relationships were weakened in seropositive animals.

5. Conclusion

High seroprevalences of antibodies against BoHV-1, BVDV, bovine parainfluenza virus type 3, N. caninum y and Leptospira spp. were found in bovine females in Chiquinquirá, Colombia, indicating the dissemination of pathogens in different herds and a high transmission level. Cattle older than 4 years presented the highest seroprevalence for BoHV-
and BVDV, while cattle aged 1–4 years had higher seropositivity to PI3, neosporosis, and leptospirosis. Regarding breeds, Holstein showed the highest prevalence for BoHV-1 and PI3, Jersey for neosporosis, and Normande for BVD and leptospirosis. Bovine >4 years and Holstein breed for IBR and >4 years for BVD were considered as risk factors.

**Data Availability**

The data used to support the research findings are included within the article.

**Conflicts of Interest**

The authors declare that there are no conflicts of interest regarding the publication of this paper.

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