Study on Occurrence of Contagious Equine Metritis in the Genital Tract of Equine.

Mona M. Sobhy 1*, A. Fathi 3, Kh.A. Abougazia 1, M.R. Oshba 3 and M.H.R. Koth 1
1*Department of Reproductive Diseases, Animal Reproduction Research Institute, Agriculture Research Centre (ARC), Cairo, Egypt.
2Immunity Units, Animal Reproduction Research Institute, Agriculture Research Centre (ARC), Cairo, Egypt.
3Ultrasonography and Laparoscope Unit, Animal Reproduction Research Institute, Agriculture Research Centre (ARC), Cairo, Egypt.

CONTAGIOUS Equine Metritis (CEM) is a transmissible venereal disease of equine, causes infertility in mares and spontaneous abortion in pregnant ones. Theiological agent of CEM is Gram negative bacteria; Taylorellaequigenitalis. In this study, a number of 45 clitoral swabs were collected from suspected mares and twenty swabs were fromurethral fossa of stallions. Swabs were kept cool during transportation on Amies with charcoal medium. At the lab, all swabs during 48 hours since they have been collected then cultured on tryptose chocolate blood agar (TBA) plates, and incubated at 37°C in microaerophilic atmosphere of 5%-10% CO2, in hydrogen of 7 days. The suspected colonies have been examined for biochemical characters of catalase, oxidase and phosphatase. All mares were scannedby ultrasound scanner (Sono Scape sonar), vaginal and/or rectal, checked for the presence of any uterine fluid which may indicate the presence of infection. On the other hand, culture investigations revealed 2 (10%) positive cases among stallions and 5 (11.1%) mare’s culture positive for T. equigenitalis. The bacterium is catalase, oxidase and phosphatase positive. The colonies were confirmed by immunofluorescent test and its sensitivity was 100%. Ultrasound examination of three mare’s uteri showing pyometra where their lumen measured 40.6-57.3 mm fully filled with echogenic particles. While, the uteri of the other two mares showingendometritis which appearedas an echogenicuterine lumen measured 18.6 mm and 37.6 mm with echogenic particles scattered on it. Biosecurity practices can help in preventing spread of CEM by bacterial culture tests on breeding stallions prior to breeding season. Also, urethral swab for CEM testing should be a stallion’s annual breeding exam. According to this study an ultrasound scan at the breeding season to check for the soundness of the mare uteri is strongly recommended.

Keywords: Contagious Equine Metritis, Mares, Stallion, Ultrasound.

Introduction

Contagious equine metritis (CEM) was first described in the United Kingdom (UK) in 1977, and then become world-wide [1]. Contagious equine metritis is an inflammatory disease of the proximal and distal reproductive tract of the mare caused by Taylorellaequigenitalis, causing mares temporary infertile [2]. Taylorellaequigenitalis is a Gram-negative, microaerophilic coccus bacterium, transmitted during mating. Infected mares had a profuse, mucopurulent vaginal discharge up to 40% of affected mares due to inflammation.
of uterus and cervix, resulting in infertility in few days after breeding and return to estrus[3]. Acute infection cause mucoid inflammation of endometrium withobvious discharge from vulva seen 1-6 days after infection post-breeding [4].

CEM is rare abortions, but infected mares can produce subclinical carrier foals. Most mares are clitoral carriers and the poor hygienic measures during breeding may spread the organism. The sites of T. equigenitalis colonization are in the clitoral fossa [5].

Stallion is asymptomatic and carries the organism on their external genitalia for a period of time [6]. Both mares and stallions become chronic carriers acting as a reservoir of T. equigenitalis and source of infection [7]. Infected stallions are passive carriers of the colonised bacteria on their external genitalia where it invades urethra and sex glands, causing pus and contaminated semen [8].

The transmission is high with breeding, where the contaminated instruments and equipment’s act as an indirect means of infection. The bacteria spread in semen collected for artificial insemination [9]. Organism has the fastidious nature and slow growth difficult to culture and requires multiple culture samples for one week [10].

The organism can be eliminated by treatment with systemic antibiotics at least 7 days or 21 days combined with antiseptic washing of the affected sites[11]. The infected equine can be treated by chlorhexidine and applying ointments 0.2% nitrofurazone [12]. Diagnostic tests must be repeated 21 days after the end of treatment. Trimethoprim-sulfamethoxazole and application 1% silver sulfadiazine cream can be used. Fertility appears to return normal after treatment of horses [13]. The present study was aimed at assessing the occurrence of Contagious Equine Metritis in equine by identification of infected carrier animals and on their treatment or elimination from breeding programs to avoid the introduction of the disease.

Materials and Methods

Sampling

From mares 45 clitoral swabs were collected, where 20 swabs were taken from the urethral fossa of stallions. All sanitary and hygienic measures were followed by using sterile saline and cotton to clean the areas of sample taken, and use new pair of disposable gloves for each animal. Swabs were kept on ice or at 4°C, and delivered to a qualified laboratory within 24 hrs. transportation and plated out at lab during 48 hours after collection of samples. All swabs were transferred in a transport medium Amies with charcoal then labeled and kept under refrigeration for variety of laboratory tests has been developed as described by Swerczek [14].

Cultural and biochemical characteristics

All swabs were cultured on plates of tryptose chocolate blood agar (TBA); by adding 5% horse blood with a rich peptone agar, trimethoprim 1µg/ml, clindamycin 5µg/ml and amphotericin B 10µg/ml then cultured at 37°C and kept in microaerophilic atmosphere of 5% to 10% CO₂ in hydrogen during a minimum period of 7 days [15]. The suspected colonies were subjected to biochemical investigations with catalase, oxidase and phosphatase[16].

Indirect immunofluorescence antibody tests (IFAT)

The Pourquier IIIF Taylorella equigenitalis Test is an indirect immunofluore scence (IIF) test, a simple and cost-effective method based on the direct detection of Taylorella equigenitalis for confirming colonies by IDEXX kits.

Ultrasonography examination

Vaginal and/or rectal examinations of 45 mares were carried out by ultrasound scanner (SonoScapeSonar) checking for presence of any uterine fluids which may indicate infection.

Results

By culture examination found 5 (11.1%) positive cases of T. equigenitalis in mare and 2 (10%) positive cases in stallion. The colonies are up to 2-3mm in diameter, soft, full edge yellow / greyish colour produces, tiny and round after 48 h. The colonies may be enlarged and turned whitish at 37°C with further incubation. The bacterium is nonmotile, catalase, oxidase and phosphatase positive for biochemical characteristics. The indirect immunofluore scence tjest was applied to discriminate T. equigenitalis from T. asinigenitalis. This test constitutes a rapid, sensitive and specific tool for confirming presumptive colonies of T. equigenitalis (Fig.1).

The scanned of 45 mares with ultrasound scanner detected 3 cases of uterine pyometra which appears as lumen measured 40.6 - 57.3mm fully filled with echogenic particles (Image 1) and 2 mares affected with endometritis appearing anechoic uterine lumen measured 18.6 mm- 37.6 mm with some echogenic particles (Image 2).
Fig. 1. Positive sample by indirect immunofluorescent test for *T. equigenitalis* under fluorescent microscope Zeiss at 10 X.

Image 1: Ultrasound image of mare uterus showing pyometra which appears as lumen measured 40.6 - 57.3 mm fully filled with echogenic particles.

Image 2: Ultrasound image of mare uterus showing endometritis which appears as anechoic uterine lumen measured 18.6 mm - 37.6 mm with some echogenic particles.
Discussion

Contagious equine metritis (CEM) is a contagious venereal disease of horses caused by *Taylorella equigenitalis* [17]. Culture positive was investigated in 2 (10%) cases stallions and 5 (11.1%) mares belonging to different localities in Egypt. These results agreed with previously demonstrated in United States with Timoney [18].

The colonies were confirmed for *Taylorella equigenitalis* by IDEXX indirect immunofluorescent testkits (Fig.1) and discriminated bacterium from *T. asinigenitalis*. IFT sensitivity was 100% in samplesand its specificity 97.2% screened by bacteriological tests constitutes a satisfactory way to determine CEM status [19].

The scanning of 45 mares with ultrasound scanner detected 3 (6.7%) uterine pyometra which appears as lumen measured 40.6 - 57.3 mm fully filled with echogenic particles (Image, 1) and 2 (4.4%) affected with endometritis appears as measured 18.6 mm- 37.6 mm (Image, 2) [20].

The equine breeding industry depends on the health of the uterus, and without a healthy uterus a mare cannot successfully conceive or carry a foal to term [21]. In this study detected low prevalence of CME can be treated or eliminated from breeding. Bacterial infections of uterus are leading cause of subfertility and infertility in mares and represent a major economic loss to the equine industry. Infected mares affected bycervicitis, metritis, with vaginal and cervical discharges. Mares had a short estrus cycle and failed to conceive or give full-term foal, which may carry the organism [22]. Mares may remain infected for several months and chronically infected mares show no signs [23].

*T. equigenitalis* spread to stallions via indirect mechanisms at shared breeding and to mares via artificial insemination [17]. CEM should be a diagnostic consideration even where this organism is to be absent. Infected horses must be quarantined and treated with disinfectants and antibiotics for several weeks [24].

Biosecurity practices can prevent the spread of CEM by conducting bacterial culture tests on breeding stallions prior to breeding season[25]. Urethral swab for CEM testing must be annual breeding examfor stallion[26]. The use of the technique of ultrasonography and laboratory findings was very helpful and of great benefit, facilitated the selection of the suspected cases among the disease carriers and identify the endometritis in mare, that greatly saved time, effort and cost.

Conclusion

The control starts before reproductive season to know the status of the animal and improving hygienic measures to prevent the spread of infection among animals.

Control of CEM depends upon early identification of infected carrier horses, followed by treatment or elimination from breeding program. Horses over two years of age are quarantined and screened for *T. equigenitalis*.

The utmost preventing measures are:
- Stallions, teasers and mares should be examined for CEM before breeding by swabbing and testing at the laboratory.
- Strict hygienic measures should be applied during handling mares and stallions.
- Stop mating using the infected stallion.
- Isolate and treat the infected horse.
- Biosecurity measures during the daily handling.
- Clean and disinfect semen equipment between stallions.
- Keep records for horse movements and semen shipments.

Acknowledgment: I will be appreciating to stuff of immunity unit in Animal Reproduction Research Institute for their helpful and support during this research.

Funding statement: Research workers paid the full costs of the research.

Conflict of interest: We didn’t found any problem during the work.

Ethical consideration: All studies described in this research were carried out at the department of Reproductive Diseases Laboratory, located at Animal Reproduction Research Institute, Agriculture Research Center, Giza, Egypt. Code No. 13/732/3/4/7.
References


