A.M.A. Ali 1
S.M. El-Sanousi 1
M.A. Al-Eknah 1
A.A. Gameel 1
E.A. Dafalla 1
A.M. Homeida 1
Y.M. Radwan 1

Studies on the infundibular cysts of the uterine tube in camel (Camelus dromedarius)

ALI (A.M.A.), EL-SANOUSI (S.M.), AL-EKNAH (M.A.), GAMEEL (A.A.), DAFALLA (E.A.), HOMÉIDA (A.M.), RADWAÎ (Y.M.).

Aeromonas hydrophila, a bacterium isolated from camels, was studied in this report. The main focus was on the morphological, histological, and bacteriological features of the cysts observed in the camels. The material was collected from abattoirs in Al-Ahsa and Riyadh, Saudi Arabia, and examined for cyst formation. The cysts were found to be unilatéral in 22 cases and bilatéral in 13 others. They contained a variable consistency fluid. The morphological and histological features were described, and the bacteriological analysis revealed the presence of Aeromonas hydrophila in 68% of the cases.

INTRODUCTION

Saudi Arabia is considered to have the fourth highest camel population in the Arabian world, with an estimated population of one million. Camels play a very important role in Saudi culture. At present, modern methods of camel production have been developed. As regards the dromedary camel, the morphology of its genitalia has been studied (1, 2, 32, 33). However, the presence of a cyst enclosed into the infundibulum of the camel's uterine tube has not been mentioned. Only little information is available on the incidence of Aeromonas hydrophila in mammals, except in man (19). However, ANNAPURNA and SANYAL (3) isolated the organism from faeces of some domestic animals: cow, buffalo, goat, and chickens.

The only single report in the literature concerning the isolation of A. hydrophila from camels is that of GAMEEL et al (18). The organism was isolated in association with Clostridium perfringens type A and Clostridium sordelli, the properties of which were studied by EL-SANOUSI et al (15).

DE and CHATTERIE (11) found that the exotoxin of A. hydrophila caused fluid accumulation in ligated rabbit ileal loops. ASAO et al (6) found it to cause fluid accumulation in infant mouse intestine.

MATERIALS AND METHODS

Collection of specimens

Two hundred eighteen genital tracts of female camels (Camelus dromedarius) were collected from abattoirs of Al-Ahsa (eastern province of Saudi Arabia) and Riyadh cities between September, 1989 to February, 1990. Out of these, genital tracts showing cyst-like formations were recorded and thoroughly examined. The estimated age of the camels ranged between 10-18 years. The morphological features of the cysts were studied and recorded. The specimens were collected within 30 min from slaughtering of animals, kept on ice and immediately transferred to the laboratory. These specimens were stored at 0 °C overnight. Next morning they were thawed and treated further for bacteriological examination.

Gynaecological investigation

Seventeen female camels were randomly selected from animals brought to Al-Ahsa abattoir. After routine preparation of the camels for rectal examination, different parts of the reproductive tract were palpated rectally. When an infundibular cyst was suspected due to unusual position of the ovary, with empty horns, the ultrasound probe scanner (WIC 50, with 5mHz scan head, USA*) was introduced rectally and guided by the operator’s hand. The probe was initially placed over the uterine body, horns and later the oviduct and ovarian region. The probe was then introduced cranially into the abdominal cavity and moved toward the left and right. After slaughter, the genitalia of the same animals examined were collected and inspected, and the findings of ante-mortem and post-mortem examinations were compared.

1. Camel Research Centre, College of Veterinary Medicine and Animal Resources, King Faisal University, POB 1757, Al-Ahsa 31982, Saudi Arabia.

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* Wesmed Veterinary Ultrasound, 78500 68th Avenue NE, Box 3001, Bothell, WA 98041-3001.
Bacteriological investigations

The surfaces of 30 cysts were carefully cleaned using 70 % alcohol and allowed to dry. The surface was seared using a red hot spatula. Disposable 10 ml plastic syringes were used to withdraw 10 ml of the fluid content aseptically. A few drops were plated directly onto 10 % sheep blood agar, MacConkey's agar and into alkaline peptone water (APW) and heart infusion broth (HIB) (31). The rest of the samples were transferred to sterile centrifuge bottles and centrifuged at 5,000 rpm for 15 min at 4 °C. The supernatant was decanted and the deposits cultured in similar media as mentioned above. At the beginning of the study, incubation was carried out aerobically at 37 °C for 24 h, later incubation was carried out at 30 °C for 24 h. When growth was obtained only in APW and HIB, subcultures were made on solid media. Plates showing no growth were reincubated for 7 days before they were regarded as negative.

The isolates were identified according to COWAN (9), CARTER (8) and KRIEG and HOLT (23). Sensitivity tests were performed with Oxoid discs. The toxin was prepared from HIB. The medium used was incubated in a water bath for 10 h with vigorous shaking.

Assay of the haemolytic activity

The haemolytic activity was detected in 2 % rabbit erythrocytes according to CUMBERBATCH (10).

Enterotoxin assay

- Rabbit loop test : enterotoxigenicity of isolates was assayed in ligated rabbit ileal loop according to DE and CHATTERJE (11) using cell-free toxin, cystic fluid and PPS as controls. The rabbits were euthanized 10 h post-operation and the fluids collected and measured.
- Infant mouse test : the test was conducted and evaluated according to DEAN (12) and ASAO et al (6).
- Assay of cytotoxic activity on Vero cells : the method of DONTA and HADDOW (13) was used.

Pathogenicity for mice

To inoculate five mice intravenously as well as five mice intraperitoneally, 0.3 ml of an 18-h whole culture of the isolates was used. Deaths were recorded and postmortem examination performed.

Delayed permeability factor (DPF) in rabbit skin

The technique of JIWA (21) was used. Only one strain of A. hydrophila was tried.

Physicochemical investigation

The fluid collected from cysts was tested for their total protein content using the Biuret method. Activities of aspartate amino-transferease (AAAT) and alanine amino transferase (ASAT) were determined colorometrically (30). The glucose content was estimated by the method of DUBOWSKI (14).

Histopathological investigations

Samples for histopathology were taken 30 minutes after slaughter from different parts of the wall of the cyst, infundibulum and cranial part of the uterine tube. Tissues were fixed in Bouin's fluid as well as in 10 % formalin, processed in paraffin and sections were stained with haematoxylin and eosin and Van Gieson's stain. Three to five rounded to oval bodies were observed only in fluid content of four cases of the studied infundibular cysts. They were fixed in 10 % formalin and their structures were studied.

RESULTS

Morphological descriptions

Out of the 218 female camel genital tracts examined, 183 (83.9 %) showed no gross abnormalities and were considered as normal. The normal uterine tube followed a tortuous course to the uterine horn and was enclosed in a peritoneal fold (the mesosalpinx) which arises from the lateral surface of the mesovarian. The length of the tube ranged between 22.5 and 30 cm. The ovarian end formed a funnel-shaped infundibulum 7-10 cm in width. The latter had a longitudinally folded mucous membrane which converged towards the abdominal opening of the tube and its free margin was indented to form the fimbriae. The ampullary region was soft and flabby, with a relatively wider lumen while the isthmus was less tortuous and hard with comparatively narrower lumen. The ovarian bursae were found to be formed between the mesosalpinx laterally and the proper ligament of the ovary, mesovarium and ovary medially. The ovary was situated within the ovarian bursa.

In most cases with cysts, the ovarian bursa had a depth of 5-12 cm. In ten cases it was crossed with fibrous tissues. Thirty five (16 %) were found to have either unilateral or bilateral cysts which appeared to be cranial dilatations of the infundibulum. These were designated "infundibular cysts". They extended to the abdominal cavity, cranial to the pelvic region between the loops of intestines. The incidence and site of the cysts in the animals examined are shown in table 1. In unilateral cases, 9 cysts
TABLE I  The number and distribution of the infundibular cysts in female camels.

<table>
<thead>
<tr>
<th>Area</th>
<th>Number of slaughtered female camels</th>
<th>Number and percentage of infundibular cysts</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Total</td>
</tr>
<tr>
<td>Al-Ahsa</td>
<td>179</td>
<td>30</td>
</tr>
<tr>
<td>Al-Riyadh</td>
<td>30</td>
<td>5</td>
</tr>
<tr>
<td>Total</td>
<td>218</td>
<td>35</td>
</tr>
</tbody>
</table>

(4.1 %) were found on the right side and 12 (5.5 %) on the left side. The infundibular cysts had different sizes, shapes and fluid consistencies. They were oval, round or kidney shaped and their walls were thin and membranous (photos 1-8) except one which had a very thick wall (photo 9). The cysts measured 18-50 cm in length and 10-28 cm in width. The content was fluid and had different colours (clear, yellowish, light or dark brown). The amount of fluid collected from each cyst ranged between 300 and 3,800 ml. In five cases some of the fluid retracted back to the cranial third of the uterine tube.

**Gynaecological investigations**

On rectal palpation, the reproductive tracts of 15 non-pregnant camels were easily retracted to the pelvic cavity. In one female the left ovary was not palpable and was pulled down by something which could not be reached by the operator’s hand. When the probe of the ultra-sound scanner was introduced deep into the abdominal cavity, a dark area showed up on the screen (see photo 13). This was not confused with follicular cysts, whereas in another female a tennis ball-like cyst was palpated on the right side (see photo 14). The presence of the infundibular and follicular cysts was confirmed later. At post-mortem examination the former was found to be an infundibular cyst, whereas the latter was as a follicular cyst.

**Bacteriological findings**

Twenty four out of 30 cyst-fluids (80 %) gave pure growth of a single type of colonies. The colonies were greisy white, translucent, moist, circular and flat with either α or β-haemolysis. The sizes of colonies increased through further incubation at room temperature, reaching 6 mm in diameter. The colonies turned into light green with age. The organisms were identified as *Aeromonas hydrophila* according to their biochemical properties (table II).

The strains of *A. hydrophila* tested were found to be sensitive to the following antibiotics : oxytetracycline (30 mcg), chlorotetracycline (30 mcg), kanamycin (30 µg), chloramphenicol (30 µg), nalidixinacid (30 µg), erythromycin and penicillin (30 µg). The organisms were also tested for their biochemical properties (table II).

**TABLE II  The biochemical properties of 24 strains of *A. hydrophila* isolated from uterine infundibulum cysts of she-camel.**

<table>
<thead>
<tr>
<th>Tests</th>
<th>Aeromonas hydrophila (A)</th>
<th>Aeromonas hydrophila (B)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Motility</td>
<td>18</td>
<td>6</td>
</tr>
<tr>
<td>Catalase</td>
<td>18</td>
<td>6</td>
</tr>
<tr>
<td>Oxidase</td>
<td>18</td>
<td>6</td>
</tr>
<tr>
<td>Glucose (acid)</td>
<td>18</td>
<td>6</td>
</tr>
<tr>
<td>Glucose (gas)</td>
<td>18</td>
<td>6</td>
</tr>
<tr>
<td>Growth on MacConkeys agar</td>
<td>18</td>
<td>6</td>
</tr>
<tr>
<td>Arabinose</td>
<td>13</td>
<td>4</td>
</tr>
<tr>
<td>Sucrose</td>
<td>18</td>
<td>5</td>
</tr>
<tr>
<td>Lactose</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Mannitol</td>
<td>17 (G)</td>
<td>5</td>
</tr>
<tr>
<td>Salicin</td>
<td>2</td>
<td>4</td>
</tr>
<tr>
<td>Arabinose utilization</td>
<td>10</td>
<td>0</td>
</tr>
<tr>
<td>Arginine and lysine utilization</td>
<td>15</td>
<td>6</td>
</tr>
<tr>
<td>Histidine utilization</td>
<td>18</td>
<td>6 (4)</td>
</tr>
<tr>
<td>TSI (H2S)</td>
<td>6</td>
<td>1</td>
</tr>
<tr>
<td>H2S in P.W. with cysteine HCl</td>
<td>10</td>
<td>0</td>
</tr>
<tr>
<td>Citrate</td>
<td>18</td>
<td>0</td>
</tr>
<tr>
<td>Indole</td>
<td>18</td>
<td>6</td>
</tr>
<tr>
<td>KCN</td>
<td>12</td>
<td>1</td>
</tr>
<tr>
<td>MR</td>
<td>1</td>
<td>6</td>
</tr>
<tr>
<td>VP</td>
<td>18</td>
<td>0</td>
</tr>
<tr>
<td>Lysine</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Ornithine</td>
<td>1</td>
<td>0</td>
</tr>
<tr>
<td>Arginine</td>
<td>18</td>
<td>6</td>
</tr>
<tr>
<td>Esculin hydrolysis</td>
<td>13</td>
<td>3</td>
</tr>
</tbody>
</table>

A = Aeromonas hydrophila with α-haemolysis ; B = Aeromonas hydrophila with β-haemolysis.

Arabic numbers indicate positive results ; ( ) indicate late results ; G : gas.
Photo 1, 2: Female genital tract with unilateral cysts: 1, left uterine tube; 2, mesosalpinx; 3, body of the uterus; 5, right uterine horn; 6, right ovary; 7, broad ligament.
(30 μg), gentamycin (10 μg), tetracycline (30 μg), cevacler (30 μg). They were partially sensitive to neomycin but resistant to polymyxin B (300 units), trisulpha (300 mcg), ampicillin (10 μg), streptomycin (10 mcg), rifampicin (5 mcg), clindamycin (2 μg), cloxacillin (1 μg), nystatin (100 units), penicillin (10 i.u.), methicillin (5 μg), lincomycin (10 μg), vancomycin (30 mcg) and fucidicacid (10 μg).

A representative strain from A. hydrophila group (A) gave a haemolytic unit of 1024 while that from group (B) gave only 64 units.

**Enterotoxin assay**

The results of rabbit ileal loop and infant mouse test are presented in table III.

**Delayed permeability factor (DPF) in rabbit skin**

Blueing was observed after six hours post inoculation (2 cm in diameter). Induration appeared after 12 h (3.2 cm in diameter) followed by necrosis and sloughing of the skin.

**TABLE III** Fluid accumulation in the rabbit ileal loop and infant mouse tests.

<table>
<thead>
<tr>
<th></th>
<th>Rabbit ileal loop (ml of fluid/cm intestinal loop*)</th>
<th>Infant mouse test weight of intestine/remaining body weight**</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>0.28</td>
<td>0.09</td>
</tr>
<tr>
<td>A. hydrophila (A)</td>
<td>1.30</td>
<td>0.11</td>
</tr>
<tr>
<td>A. hydrophila (B)</td>
<td>1.10</td>
<td>0.10</td>
</tr>
<tr>
<td>Fluid from cyst</td>
<td>0.70</td>
<td>0.11</td>
</tr>
</tbody>
</table>

* : positive  ** : > 3.0% : positive.
A. hydrophila μ haemolytic  B: A. hydrophila μ haemolytic.

**Pathogenicity for mice**

Deaths occurred between 18-24 h with signs of septicaemia. Blindness was observed in mice inoculated intravenously.

**Assay for toxicity in Vero cells**

Loss of adherence and rounding were the common features in the cells tested.
Histology

The cyst wall, examined at different places was composed mainly of fibrous connective tissue and smooth muscle and lined at one side by low columnar epithelium. The latter had a vacuolated appearance and an apparently striated border with surface globules. In close association with the epithelium was a homogenous pink area which represented the cyst contents (photo 10). At the level of the infundibulum the cyst wall was either composed of fibrous connective tissue or smooth muscle (photo 10). The lining columnar epithelium had a similar striated border and bleb-like luminal projections. It was thrown into folds which were cut out at places into acinar-like structures containing pinkish material (photo 11). The mucosa of the cranial part of the uterine tube (ampulla) was distinctly folded and formed cyst-like structures lined by similar columnar epithelium with centrally placed nuclei. The contents of the cysts showed many free pin-kish globules. The subepithelial layer was infiltrated by monolocular cells of predominantly lymphocytes. The inner circular and outer longitudinal smooth muscle layers of the tunica muscularis were separated by loose connective tissue containing thick walled blood vessels. Some of the bodies found within the content of the infundibular cyst had fibrous wall and the others were lined with low columnar epithelium (photo 12). The wall of these bodies enclosed within it, was a homogenous eosinophilic material containing aggregates of bacilli.

Physicochemical findings

The infundibular cyst fluid had a specific gravity of 1.15 and a pH of 8.87. The protein content was 7.2 g/dl while the glucose level was 35 mg/dl. The AIAI and ASAl values were 51 and 2.8 μ/ml, respectively.
Photo 8: Membraneous kidney-shaped infundibular cyst.

Photo 9: Rounded thick walled cyst.

Photo 10: Cyst wall composed of fibrous connective tissue and smooth muscles and lined by low columnar epithelium with vacuolated cytoplasm. Note thick walled blood vessels (a) and epithelial folds (b) (HE x 104).
DISCUSSION

The described histological and morphological structures of the cysts prove that they were dilatations of the cranial part of the infundibular wall, hence designated as "infundibular cysts". The prevalence of such cysts was fairly high in the camels examined and coincided with the increased incidence of infertility observed in camels brought to the Veterinary Teaching Hospital in Saudi Arabia (4). Concerning abnormalities of the female genital tract, especially those of the urogenital ducts and other paraovarian cysts of the domestic animals (24, 29), there was no mention of similar cyst formation.

The ultrasound scanner applied to detect infundibular cyst(s) in the living animals paves the way for future screening of female camels. The ultrasound technique proved thus to be helpful in the detection of ovarian cysts in
female camels (25, 26). The long gestation period and the short breeding season in the dromedary camel impose restrictions concerning the maximum fertility rate. In addition, the high rate of eggs and the early embryonic mortality require further limitations (27, 28, 35). The results of the present investigation add a new factor which can affect pregnancy and fertility rates. The morphological findings of the infundibular cyst clearly showed that the passage of the ovum to the site of fertilization in the uterine tube is prevented especially in bilateral cases where the two entrances to the abdominal opening of the tube are occluded. However, the possibility of pregnancy in unilateral cases cannot be excluded. Most of the ovaries in the unaffected side showed signs of follicular activity and pregnancy may occur, though infection with A. hydrophila may lead to abortion. Moreover, pregnancy is considered as a stress factor in general (36) and this in turn is predisposing to A. hydrophila infection, the latter acting as a secondary invader according to various workers (19, 36).

In general, fertility rates have been reported to be poor in Camelus dromedarius in studies conducted in Tunisia and Kenya (7, 16). In contrast to that, ARTHUR et al. (5) considered the fertility to be high in Saudi Arabia, a finding which is not substantiated by the present study taking into consideration the rate of infection with A. hydrophila (68 %). This coupled with other factors, that contribute normally to infertility, might boost the percentage upwards. Consequently, this will be an obstacle in camel reproduction in Saudi Arabia.

Species of the genus Aeromonas have long been recognized as pathogen in amphibians (7) and fish (20) and have been found to be widely distributed in the aquatic environment (22). Apart from scarce reports in the literature, A. hydrophila is associated mainly with enteral infections. The present report is considered as the second one concerning its isolation from the camel (18) and the first record of its isolation from the genital tract. The isolation of A. hydrophila from 68.6 % of infundibular cysts suggests its role in the formation of such cysts, although other unknown factors should not be completely excluded. The amounts of fluid accumulated in the cysts are due to the toxins released by A. hydrophila. Such toxins have been proved to cause accumulation of fluids by various authors (6, 34). Such findings were substantiated by the present study using ligated rabbit ileal loops and intestines of infant mice. As far as the lesions are concerned, the organisms isolated may be of low virulence, causing in most cases fluid accumulation rather than eliciting marked inflammatory reactions.
The mode of infection is difficult to determine with great certainty, however, ascending infection might be possible since A. hydrophila was found by various authors in faeces of normal animals. On the other hand, the prevalence of A. hydrophila in cattle is greater than in other animals (19).

Improvement of the bacteriological techniques, i.e. centrifugation of the cyst fluids and enrichment prior to plating on solid media made it possible to isolate A. hydrophila routinely from such cysts. It is thus worth trying to study the prevalence of A. hydrophila in the intestines of apparently healthy camels as well as in their environments.

CONCLUSION

The present investigation revealed that the cyst described in the female camel was designated as "infundibular cyst" which is a dilation of the cranial part of the infundibular wall of the uterine tubes. The results add a new factor which can affect pregnancy and fertility rates in camels. The isolation of the A. hydrophila from the infundibular cyst might suggest its role in the formulation of such a cyst. Among other factors involved in the cyst formation, the role of the A. hydrophila needs further studies.

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REFERENCES


