Original Research Article

Antibiotic Susceptibility Pattern and Multiple Antibiotic Resistance Index of Salmonella enterica Isolates from Horses in Bikaner

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A B S T R A C T

In present study, 160 horses were screened for enteric Salmonella and their antimicrobial resistance pattern in Bikaner. Presence of InvA gene in faecal samples was used as marker for Salmonella shedding. Antimicrobial resistance analysis against 10 different antibiotics was done on 11 isolates of Salmonella enterica. The results revealed that the Salmonella isolates were sensitive to ceftriaxone + tazobactum (100 per cent), cefoperazone (81.81 per cent), and cefuroxime (81.81 per cent). All (100 per cent) the strains were found resistant to penicillin-G and 72.73 per cent isolates were resistant to tetracycline. Six isolates exhibited multiple drug resistance against 3 to 9 antibiotics with Multiple Antibiotic index ranging from 0.10 to 0.90. MAR, index of 0.70 was found in maximum number of isolates (3) which were resistant to seven different antimicrobials.

Keywords
Horses, Salmonella enterica, Antibiogram and Multiple Antibiotic Resistance Index

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Introduction

Salmonellosis is caused by enteric or systemic infection with bacteria of Salmonella spp. Horses can be affected with salmonellosis without showing any apparent clinical signs and symptoms, and transient shedding of organisms in faeces. Such latent carriers can turn into a clinical case under stress condition. Case fatality rate of salmonellosis in equines is up to 60 per cent (Constable et al., 2017).

Horses are at high risk of nosocomial salmonellosis infection in veterinary teaching hospitals because of exposure to a common source of Salmonella organisms (e.g., contaminated feed) or by lateral transmission from infected animals (Kim et al., 2001). In some instances, these facilities have been forced to close temporarily (≥ 3 months) because of serious outbreaks of clinical disease caused by Salmonella infection in horses (Ernst et al., 2004). In the past few
decades, emergence of antibiotic resistance among different species of bacteria is on the rise (Davies and Davies, 2010). The problem is compounded when it occurs in pathogenic bacteria like *Salmonella enterica* having public health significance. Bikaner city have a good equine population and in past no study have been carried out to know the status of antibiotic resistance against salmonellosis in horses in this area. Therefore, the present study was conducted to know the antibiogram profile of *Salmonella enterica* isolated from horses of Bikaner to evaluate and monitor the extent of antibiotic resistance in the isolates of this region. Multiple Antibiotic Resistance (MAR) index was also calculated which has been shown to be a cost effective and valid method of bacteria source tracking.

**Materials and Methods**

**Collection of samples**

Rectal swabs were taken in triplicate over a period of 24 hours by using transport swab containing cary-blair medium from 160 horses belonged to different stud farms and veterinary hospitals of Bikaner area in Rajasthan state.

**Sample processing**

*Salmonella* were detected using selective enrichment faecal cultures. For this, faecal swab samples were used to inoculate faecal material into 10 ml of mannitol selenite broth, which were incubated overnight at 37 °C. The swabs were then streaked onto xylose lysine deoxycholate (XLD) agar plates and incubated at 37°C for 18-24 hours for isolation and identification of *Salmonella* pure colonies. On XLD agar plates colony morphology of *Salmonella* was appeared as red colonies with black centres (Plate. 1), and on microscopic examination gram-negative, rod-shaped bacterial cells after gram’s staining. For confirmatory diagnosis isolated pure colonies of *Salmonella* were further incubated for 24 hours at 37 °C in nutrient broth for DNA extraction and characterization by highly conserved 457 bp nucleotide sequence within the invasion gene (*invA*) of *Salmonella* spp. by polymerase chain reaction (PCR) assay (Plate. 2).

**In-vitro chemotherapeutic sensitivity**

In the present investigation, all *Salmonella* spp. isolates were subjected to resistotyping with 10 different antibiotics. *In-vitro* antibiotic sensitivity pattern of the isolates were determined by disc diffusion method of Bauer *et al.*, (1966). The *Salmonella* spp. isolates were tested against commonly used antibiotics *viz.* cefoperazone (CPZ), cefuroxime (CXM), ceftriaxone/tazobactum (CIT), penicillin (P), amoxycillin/clavulanic acid (AMC), ampicillin/subactum (A/S), streptomycin (S), erythromycin (E), tetracycline (TE) and gentamicin (GEN).

Test bacteria broth culture swabbed on Muller Hinton agar plates with sterile cotton swabs. When broth culture was dried, the antibiotic discs were placed with the help of automatic disc dispenser in front of flame. These petri dishes were incubated for 15-20 hours and observed for zone of inhibition. Diameters of zone of inhibition (mm) were observed and results were interpreted as sensitive, intermediate and resistant based on the CLSI guidelines.

**MAR (Multiple Antibiotic Resistance) Index Study**

The MAR Index of an isolate is defined as a/b, where ‘a’ represents the number of antibiotics to which the isolate was resistant and ‘b’ represents the number of antibiotics to which the isolate was subjected (Jayaraman *et al.*, 2012).
Identification of MDR (Multi Drug Resistance)

Multi Drug Resistance is defined as resistance to more than two classes of antibiotics among all the tested antibiotics. The Multi Drug Resistance (MDR) characters of the isolates were identified by observing the resistance pattern of the isolates to the antibiotics.

Results and Discussion

The detailed results of antibiotic sensitivity test of Salmonella isolates are given in table 1. The results of antibiogram showed that all (100 per cent) of the isolates were resistant to penicillin-G followed by tetracycline (72.73 per cent), amoxycillin + clavulanic (54.55 per cent), gentamicin (45.45 per cent), ampicillin + sulbactum (36.36 per cent), erythromycin (36.36 per cent), streptomycin (36.36 per cent), cefoperazone (9.09 per cent) and cefuroxime (9.09 per cent) (Fig. 1).

Ceftriaxone + tazobactum were found most effective against Salmonella spp. during in-vitro study with sensitivity in all isolates (100 per cent). Cefoperazone and cefuroxime were found reasonably effective as both the antibiotics showed 81.81 per cent sensitive against Salmonella isolates.

Table 1. Antibiotic sensitivity patterns of Salmonella isolates of horse origin

<table>
<thead>
<tr>
<th>S.No.</th>
<th>Antibiotics</th>
<th>Sensitive</th>
<th>Intermediate</th>
<th>Resistant</th>
</tr>
</thead>
<tbody>
<tr>
<td>1.</td>
<td>Ceftriaxone + tazobactum</td>
<td>11(100.00)</td>
<td>00 (00.00)</td>
<td>00 (00.00)</td>
</tr>
<tr>
<td>2.</td>
<td>Cefoperazone</td>
<td>09(81.81)</td>
<td>01 (09.09)</td>
<td>01 (09.09)</td>
</tr>
<tr>
<td>3.</td>
<td>Ampicillin + sulbactum</td>
<td>06(54.54)</td>
<td>01 (09.09)</td>
<td>04 (36.36)</td>
</tr>
<tr>
<td>4.</td>
<td>Cefuroxime</td>
<td>09(81.81)</td>
<td>01 (09.09)</td>
<td>01 (09.09)</td>
</tr>
<tr>
<td>5.</td>
<td>Streptomycin</td>
<td>05(45.45)</td>
<td>02(18.18)</td>
<td>04(36.36)</td>
</tr>
<tr>
<td>6.</td>
<td>Amoxycillin + clavulanic</td>
<td>02(18.18)</td>
<td>03 (27.27)</td>
<td>06(54.54)</td>
</tr>
<tr>
<td>7.</td>
<td>Penicillin-G</td>
<td>00 (00.00)</td>
<td>00 (00.00)</td>
<td>11(100.00)</td>
</tr>
<tr>
<td>8.</td>
<td>Tetracycline</td>
<td>00(00.00)</td>
<td>03(27.27)</td>
<td>08(72.73)</td>
</tr>
<tr>
<td>9.</td>
<td>Erythromycin</td>
<td>04(36.36)</td>
<td>03(27.27)</td>
<td>04(36.36)</td>
</tr>
<tr>
<td>10.</td>
<td>Gentamicin</td>
<td>03(27.27)</td>
<td>03(18.18)</td>
<td>05(45.45)</td>
</tr>
</tbody>
</table>

Table 2. Multiple Antibiotic Resistance index of Salmonella enterica isolates

<table>
<thead>
<tr>
<th>S.No.</th>
<th>isolates</th>
<th>No.of antibiotics to which isolate is resistant(a)</th>
<th>MAR index=a/b</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Salmonella enterica</td>
<td>1</td>
<td>1/10= 0.10</td>
</tr>
<tr>
<td>2</td>
<td>Salmonella enterica</td>
<td>1</td>
<td>1/10= 0.10</td>
</tr>
<tr>
<td>3</td>
<td>Salmonella enterica</td>
<td>1</td>
<td>1/10= 0.10</td>
</tr>
<tr>
<td>4</td>
<td>Salmonella enterica</td>
<td>2</td>
<td>2/10= 0.20</td>
</tr>
<tr>
<td>5</td>
<td>Salmonella enterica</td>
<td>2</td>
<td>2/10= 0.20</td>
</tr>
<tr>
<td>6</td>
<td>Salmonella enterica</td>
<td>3</td>
<td>3/10= 0.30</td>
</tr>
<tr>
<td>7</td>
<td>Salmonella enterica</td>
<td>4</td>
<td>4/10= 0.40</td>
</tr>
<tr>
<td>8</td>
<td>Salmonella enterica</td>
<td>7</td>
<td>7/10= 0.70</td>
</tr>
<tr>
<td>9</td>
<td>Salmonella enterica</td>
<td>7</td>
<td>7/10= 0.70</td>
</tr>
<tr>
<td>10</td>
<td>Salmonella enterica</td>
<td>7</td>
<td>7/10= 0.70</td>
</tr>
<tr>
<td>11</td>
<td>Salmonella enterica</td>
<td>9</td>
<td>9/10= 0.90</td>
</tr>
</tbody>
</table>
**Fig.1** Percentage antibiotic sensitivity patterns of *Salmonella* isolates of horse origin

![Bar chart showing antibiotic sensitivity patterns of Salmonella isolates of horse origin.](chart)

**Plate.1** *Salmonella* species: red colonies, with black centers on XLD agar

![Plate showing Salmonella species on XLD agar.](plate1)

**Plate.2** Amplification of a 457bp fragment of *Salmonella* genus specific InvA gene in apparently healthy horses

![Amplification gel showing InvA gene amplification.](plate2)

(P* = +ve Control; C* = -ve Control; L = Ladder)
These results are comparable to the findings of earlier workers who determined different resistance patterns among *Salmonella* strains of equine origin and other animals in India (Agarwal *et al.*, 2004; Chandra *et al.*, 2006; Gaind, 2007; Singh *et al.*, 2007; Singh *et al.*, 2009; Khan *et al.*, 2015; Kalambhe *et al.*, 2016; Kulkarni *et al.*, 2019), might be attributable to change in antimicrobial drug therapy in equids. Growing resistance towards antimicrobial drugs has been prevalent worldwide among members of *Enterobacteriaceae* from animal origin especially in *Salmonella* sp. The high resistance to penicillin-G and tetracycline antibiotics might be due to over use of drugs or transmission of resistant strains of *Salmonella* from other sources i.e. human, other animals and environment etc. in the absence of hygienic practices.

Multiple drug resistance was recorded in 6 (54.54 per cent) out of the 11 isolates which showed resistance against 3 to 9 antibiotics (table 2). The multiple drug resistance Index calculated ranged from 0.10 to 0.90 with 0.70 being the predominant MAR index in 3 isolates (table 2).

Multiple antibiotic resistance (MAR) has been attributed to the presence of plasmids which contain one or more resistance genes, each encoding a single antibiotic resistance phenotype (Daini *et al.*, 2005). Findings of the present study were in accordance with Singh *et al.*, (2012), Khan *et al.*, (2015) and Kalambhe *et al.*, (2016) who also reported multiple drug resistance against 3 to 9 antimicrobials in several *Salmonella* isolates of animal and human origin in India. Emergence of multiple drug resistant *Salmonella* strains has often been held responsible for frequently occurring outbreaks and hyper-endemicity of salmonellosis in India (Singh *et al.*, 2006). India has become home for multiple drug resistant strains of different *Salmonella* serotypes. The phenomena namely the localization of resistance genes on conjugative plasmids, co-integrate formation which gave rise to new plasmids and the localization of resistance genes on transposes has led to the efficient spread of multiple drug resistance in *Salmonella* (Helmuth, 2000).

In conclusion the present study revealed that ceftriaxone + tazobactum is the best effective antibiotic to treat *Salmonella* infection in horses. The susceptibility pattern of *Salmonella* isolates of horses from different sources of Bikaner region showed that they were highly resistant to penicillin-G and tetracycline.

**References**


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