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YERSINIA ENTEROCOLITICA AMONG SHEEP CARCASSES (With One Table)

By

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ميكروب اليرسينيا انتيروكوليتيكا في ذبائح النشآن

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تم عزل ميكروب اليرسينيا انتيروكوليتيكا من 7.5% من ذبائح النشآن التي تم فحصها . وقد وجد أن أعلى نسبة لعزل الميكروب كانت من سطح الذبائح 7.5% يليها السطح الداخلي وكان 5% ونسبة 3.75% في محتويات القولون .

SUMMARY

Y. enterocolitica was recovered from 7.5% of the examined slaughtered sheep carcasses. The highest recovery rate was from the outer surface (7.5%) followed by the inner surface (5%) and least the rectal contents (3.75%).

INTRODUCTION

Y. enterocolitica organism was reported as a new human pathogen named *Bacterium enterocoliticum* 50 years ago in the USA by SCHLEIFSTEIN and COLEMAN (1939). The organism was named later as *Y. enterocolitica* according to BERGY (1984).

Since the last decade several authors had incriminated *Yersinia enterocolitica* as a meat borne pathogen causing gastrointestinal infection in man (HANNA *et al.*, 1976; MALLARET *et al.*, 1979 and LEE *et al.*, 1981).

Y. enterocolitica infections in man were studied by ZEN-YOJI & MARUYAMA, 1972; SCHIEVEN & RANDALL, 1974; RABSON *et al.*, 1975 and MARTIN *et al.*, 1982. Authors frequently incriminate food of animal origin and water as a source of the oral infection in man (RABSON & KOORNHOF, 1972 and MARTIN *et al.*, 1982) and house hold animals for contact infection (GUTMAN *et al.*, 1973).

Strains of *Y. enterocolitica* were isolated from aborted lamb suffering from acute enteritis (BREWER and CORBEL, 1983) and also from feces of slaughtered sheep (LUDES, 1983 and LUDES & WEISS, 1984). Two of the isolated strains are serologically similar to those which are pathogenic for man.

The present study was planned to monitor sheep slaughtered at Cairo abattoir for *Y. enterocolitica*.

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MATERIALS and METHODS

For the purpose of monitoring slaughtered sheep (80 animals) for *Y. enterocolitica*, the outer and inner surfaces of the carcasses were swabbed in double, in addition, the rectal contents were also sampled by taking 10 g of rectal contents in sterile test tubes. All swabs and samples were transferred to the laboratory in an ice box within 2 hours.

The materials obtained were subjected to :

- 1- Enrichment using Phosphate buffered saline (P.B.S) according to BREWER & CORBEL (1983) and Trypticase soya broth (T.S.B) according to DUDLEY & SHOTTS (1979).
- 2- Plating on solid media (DUDLEY & SHOTTS, 1979 and BREWER & CORBEL, 1983) using S.S agar, MacConkey agar and Cellobiosargenine lysine agar plates.
- 3- Identification of isolates:
 - Morphologically using Gram's stain after SWANINATHAN *et al.* (1982).
 - Biochemically according to BERCAVIER and MOLLARET (1984).

RESULTS and DISCUSSION

Obtained results were similar to those reported by LUCES & WEISS (1984). The fact that *Y. enterocolitica* was recovered from carcass surfaces in a higher frequency than from the fecal samples reflect the possibility of cross contamination from the already infected rectal contents to the carcass surfaces during evisceration and primary processing. This is because the unhygienic state prevailing at the old slaughter house in Cairo and the neglect of prevention of contamination during carcass preparation.

MENGE *et al.* (1986) in a laboratory investigation on mice supports the view of GUTMAN *et al.* (1973) that the oral route is the possible way for human infection. While SHAYEGANI (1986) reported that *Y. enterocolitica* maintained at a low temperature are more likely to be pathogenic when ingested.

Therefore, it could be concluded that cross contamination, must be prevented during abattoir practice. Moreover, decontamination should be performed for reduction of contamination of meat with *Y. enterocolitica*.

Table (1): Frequency of *Y. enterocolitica* in the examined samples.

Samples	No.	+ ve samples	
		No.	%
Rectal contents	80	3	3.75
Carcass:			
Outer surface	80	6	7.5
Inner surface	80	4	5.0
Total animal	80	6	7.5

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