

Seroprevalence and Molecular Epidemiology of Brucellosis in Pakistan

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Objective

To detect the presence of brucella in serum samples of occupationally exposed human and animals by conventional screening methods.

To perform epidemiology of brucella molecular based tests including genus and species specific PCR.

To check the brucella prevalence in occupationally exposed human.

Introduction

Livestock sector contributes more than 58% to agriculture-based economy of Pakistan. Diseases of socio-economic importance are posing an enormous pressure to the growth of this sector. Zoonotic diseases are generally neglected in wake of epizootics having epidemic potential. One Health is a multi-sectoral approach to control zoonotic diseases at animal level to mitigate risk of transfer to the humans and environment. Despite various control programs, zoonosis is known to cause public health emergencies at various regional and national levels. OIE declared brucellosis as a model bacterial disease to control zoonosis in developing countries. Genus *Brucella* is expanding with its discovery in various amphibian species and marine mammals and demands control efforts at various levels. Reporting of zoonosis is less than actual prevalence in third world countries like Pakistan where disease is considered endemic but no official data is available. In this study, brucellosis was used as a model disease to emphasize the significance of One Health.

Methods

In total, 183 occupationally exposed human and 324 animal blood samples were collected from five different geographical areas of Punjab and one region from KP. For detection of brucella, rose bengal plate test (RBPT) and cELISA were carried out on serum samples. For molecular epidemiology genus specific PCR *BCSP31* and species specific PCR *IS711* were conducted. Fifty-seven milk samples as environmental samples were also collected. For the testing of milk for the detection of brucella, Milk Ring Test (MRT) was applied.

Results

Serologically in animals 26(8%) samples were found positive by RBPT & 31(9%) by cELISA. Disease was detected in 42(13%) & 59(18%) samples by applying molecular methods using genus specific PCR *BCSP31* & species specific PCR *IS711*. Disease was recorded in humans as 16(8%), 24(13%), 33(18%), 56(30%) by RBPT, cELISA, PCR *BCSP31* & PCR *IS711*, respectively. Out of 57 milk samples collected from different areas were tested by Milk Ring Test (MRT) & 12(21%) samples were found positive.

Conclusions

It is a significant finding that raw milk is a constant source of disease exposure to farmers, milking men and general users. Disease prevalence was more in people associated with milking activities possibly due to use of raw milk. This study validates the prevalence of brucellosis in Pakistan with significant presence of disease in occupationally exposed individuals emphasizing the close collaboration between veterinary and human health sectors.

This study will broaden our knowledge of disease prevalence and epidemiology in Pakistan. The data produced from this study will help in future control and eradication of this important zoonosis using one health approach.

Keywords

Brucellosis; Occupationally exposed; Pakistan; animals

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