DOI: http://dx.doi.org/10.18782/2320-7051.5957

**ISSN: 2320 – 7051** *Int. J. Pure App. Biosci.* **5 (5):** 1406-1414 (2017)



# 

**Review** Article

### Historical Perspectives and Epidemiology of Bovine Tuberculosis

Neelam Rani<sup>1\*</sup>, Piyush Tomar<sup>1</sup>, P. K. Kapoor<sup>1</sup>, N. K. Mahajan<sup>1</sup>, Naresh Jindal<sup>1</sup>and Rajesh Chhabra<sup>2</sup>

<sup>1</sup>Department of Veterinary Public Health and Epidemiology <sup>2</sup>College Central Laboratory, College of Veterinary Sciences, Lala Lajpat Rai University of Veterinary and Animal Sciences, Hisar-125 004 \*Corresponding Author E-mail: neelamvet2011@gmail.com Received: 25.09.2017 | Revised: 19.10.2017 | Accepted: 23.10.2017

#### ABSTRACT

Bovine tuberculosis is an important bacterial zoonotic disease of animals caused by Mycobacterium bovis which affect animal health, production and overall economy of the country. Among animals, the disease transmission occurs mainly through inhalation. In calves, the disease may occur via ingestion of contaminated milk from infected dam and rarely via congenital or cutaneous route. The disease transmits to human through ingestion of contaminated milk or milk products and also through inhalation. The disease has a slow progression which creates difficulty in early diagnosis and treatment. Conventional diagnostic methods should be used in combination with modern molecular and immunological techniques for early and accurate diagnosis. Treatment is usually not recommended in animals. Vaccination is being carried out in some countries as a preventive measure. For control of the disease, slaughter of the reactor animals in developed countries and segregation of the suspected or reactor animals in developing countries is preferred. This article focuses on the historical perspectives and epidemiology of bovine tuberculosis.

*Key words:* Anergic, Mycobacterium bovis, Mycobacterium tuberculosis complex, Wildlife, zoonotic.

#### **INTRODUCTION**

Bovine tuberculosis (bovine TB) is an OIE listed chronic, bacterial zoonotic disease caused by *Mycobacterium bovis* (*M. bovis*), a member of the *Mycobacterium tuberculosis* complex (MTC). It is one of the major infectious disease among domestic animals and certain wildlife population<sup>25,26</sup>. World Health Organization (WHO) classified bovine tuberculosis among seven neglected zoonotic diseases having potential to infect man<sup>15</sup>. It is an economically significant disease because of trade restriction and serious public health consequences mainly in developing countries<sup>35,45</sup>. It is a known fact that humans and animals have had close interactions and this interaction is increasing due to the advancement in livestock production systems to meet out the demand for animal products. This contributes to the transmission of infectious diseases from animals to human<sup>21,40</sup>.

Cite this article: Rani, N., Tomar, P., Kapoor, P.K., Mahajan, N.K., Jindal, N. and Chhabra, R., Historical Perspectives and Epidemiology of Bovine Tuberculosis, *Int. J. Pure App. Biosci.* **5**(5): 1406-1414 (2017). doi: http://dx.doi.org/10.18782/2320-7051.5957

Cattle and buffalo are considered as the natural host of *M. bovis*<sup>39</sup>. Zebu cattle are much more resistant to tuberculosis than crossbred and exotic breeds. Mycobacterium tuberculosis has also been reported in a wide range of domestic and wildlife animal species, more frequently those living in close contact with human for a long time<sup>2,28</sup>. It is estimated that *M. bovis* is responsible for 5% of all TB infections in humans<sup>10,22</sup>. In developed countries, there is dramatic decline in the incidence of human tuberculosis due to *M. bovis* due to mandatory pasteurization of milk in combination with tuberculin skin test and slaughter of reactor cattle<sup>29</sup>. During the year 2015, estimated incidence of human TB due to M. bovis was 149,000 cases and 47,400 cases in worldwide and in South-East Asia respectively whereas mortality rate was 13,400 cases and 2,280 cases worldwide and in South-East Asia respectively<sup>56</sup>.

#### Historical perspectives:

Tuberculosis has been co-evolved with human civilization. The earliest references related to TB can be found in the language Samskritam (Sanskrit). In ancient Chinese and Arabic literature, there is documentation of a TB-like disease<sup>23,36</sup>. Tuberculosis was referred to as Yakshma, in the ancient Indian scriptures, The Vedas. The word "tuberculosis" appears to have been derived from the Latin word *tubercula* i.e. a small lump<sup>13</sup>. The discovery of tubercle bacillus was announced by Robert Koch on 24<sup>th</sup> March 1882 during the monthly evening meeting of the Berlin Physiological Society. Since 1982, 24th March is being celebrated "World as TB Dav" commemorating the centenary of this event<sup>58</sup>. The work of Villemin in 1865<sup>11</sup> and Koch in  $1882^7$  shown the cross adaptability of the tubercle bacilli from one species to other species and revealed that tuberculosis could be transmitted from animals to humans<sup>11</sup> This was supported by demonstration of *M. bovis* in a child with tuberculous meningitis by Ravenel in 1902. In developed countries, due to better public health practices, massive vaccination with Calmette-Guerin bacillus (BCG) vaccine and the advent of streptomycin, para-amino salicylic acid (PAS) and isoniazid, there was decline in morbidity and mortality rates due to TB during the 20<sup>th</sup> century but it continued to affect the developing countries like India. In the mid-1980s due to emergence of acquired immunodeficiency syndrome (AIDS), incidence of TB began to increase in both developed and developing world. In April 1993, WHO declared TB to be a "global emergency" after recognizing the global burden of TB<sup>58</sup>. During late 1990s, drugresistant TB (DR-TB) with multidrug-resistant TB (MDR-TB) and during first decade of the 21<sup>st</sup> century, extensively drug-resistant TB (XDR-TB) was emerged<sup>8</sup>. There are reports of occurrence of extremely drug-resistant TB (XXDR-TB), super XDR-TB, totally drugresistant TB (TDR-TB)<sup>52</sup> from some parts of the world including India<sup>48</sup>.

#### The etiological agent:

The Mycobacterium tuberculosis complex (MTC) includes five Mycobacterium species-M. tuberculosis, M. canettii, M. africanum, M. microti, M. bovis and two subspecies- M. caprae and M. pinnipedii. These species are characterized by 99.9% similarity at nucleotide level and their identical 16S rRNA sequences<sup>19,42</sup>. but differs in terms of phenotype, host tropism and pathogenicity. The name "Mycobacterium", meaning fungus like bacterium and is derived from mould-like appearance of Mycobacterium tuberculosis when grown on liquid media. They are characterized phenotypically as Gram positive bacilli or cocco-bacilli, strictly acid-fast, vary from  $0.5-10\mu m \times 0.2-0.6\mu m$ , non-motile, noncapsular, non-spore former and facultative intracellular aerobes. They are not heatresistant, being readily killed by normal milk pasteurization conditions but resistant to drying and can persist in environment for long periods. Their cell wall is rich in lipids *i.e.* mycolic acid which is responsible for acid fastness, hydrophobicity and provides resistance to many disinfectants, laboratory stains, physical injuries, antibiotics and lysosomal enzymes<sup>3</sup>. The genome of MTC consists of double stranded circular

deoxyribonucleic acid (DNA), consisting of approximately 4.4 million base pairs. The genome has a high (G + C) content approximately 65%<sup>4</sup>. Almost all members of M. tuberculosis complex have 0-25 copies of IS6110 gene but only one copy of IS6110 found in M. bovis. Among the various repetitive sequences only IS6110 and IS1081 are insertion sequences and others are short sequences with no known function. The direct repeat (DR) region in M. tuberculosis complex strains is composed of multiple direct variant repeat sequence (DVRS) each of which is composed of a 36 bp direct repeat and a non repetitive spacer sequence of similar size. It has been shown that there is extensive polymorphism in the direct repeat region by the variable presence of direct variant repeat sequence and this polymorphism is used for differentiation of mycobacterium species and to define epidemiological relationships<sup>24</sup>.

#### **Epidemiology:**

Bovine TB has emerged as an economically significant disease affecting both animals and human. It is distributed globally except Antarctica, Caribbean islands, parts of South America and Australia, Iceland, Denmark, Sweden, Norway, Finland, Austria, Switzerland, Luxembourg, Latvia, Slovakia, Lithuania, Estonia, the Czech Republic, Canada, Singapore, Jamaica, Barbados and Israel. Eradication programs are in progress in other European countries, Japan, New Zealand, the United States, Mexico, and some countries of Central and South America where it has been eradicated by following strict test and slaughter policies<sup>47</sup>. It is a major public health problem mainly in developing countries including India. Factors which influence the occurrence of disease are sex, breed and social management of livelihood conditions<sup>1</sup>. Hereditary and maternal factors also influence incidence of TB in cattle progeny. Infected cattle are the main source of infection for other animals and human beings also. Organisms are excreted in exhaled air, sputum, faces, milk, urine, vaginal and uterine and discharges discharges from open peripheral lymph nodes<sup>55</sup>. It is more prevalent in dairy workers because of their close association with animals.

#### Transmission:

The transmission between animals occurs mainly through aerosols. The transmission rate is increased by close contact among animals and intensive breeding<sup>14</sup>. Other rare routes like cutaneous, congenital and genital have also been reported. Suckling calves can get the infection through consumption of infected Sometimes, tuberculosis milk. can be foodborne  $also^{22}$ . The infected bull may also disease through transmit artificial insemination with the use of infected semen<sup>55</sup>. Others factors like long survival period of the organism in environment also contribute to an increased risk of infection<sup>2,12</sup>. It is quite likely that animals can be infected with humanoriginated M. tuberculosis, becoming a reverse zoonosis, which in turn may act as a source of infection for humans.

#### Pathogenesis:

of pathogenesis The mechanism of tuberculosis caused by *M. tuberculosis* and *M.* bovis is almost same as they have 99.9% similarity at nucleotide level and identical 16S rRNA sequences<sup>54</sup>. After entry into the lungs, the bacteria are phagocytosed by the alveolar macrophages. Various chemokines and cytokines releases as a result of interaction of mycobacteria with macrophage receptors that serve as signals. By inhibiting phagolysosomal fusion, M. tuberculosis, M. bovis and M. avium can survive inside the macrophages<sup>38,53</sup>. It will result in migration of macrophages and dendritic cells to the site of infection. The dendritic cells engulf bacteria and migrate to the lymph nodes. In response to mediators produced by infected cells, primed T cells migrate back to infection site in lungs and leads to granuloma formation. The granuloma is formed by necrotic cells in the center of the tubercle surrounded by epitheloid cells and multinucleated giant cells, encapsulated by connective tissue<sup>25,26</sup>. The necrotic core of cells can often become calcified as the tubercle matures<sup>5,16</sup>. The infection can spread via hematogenous route to lymph nodes and to other organs<sup>6</sup>.

#### Clinical signs and symptoms:

The disease usually take months to develop and can affect any part of the body but generally affect lungs and nearby lymph nodes. In early stages, the infected animals remain asymptomatic. Infection can reactivate during stress or in old age. During late stages, symptoms include progressive common emaciation, low grade fluctuating fever and capricious appetite. Pulmonary involvement characterized by moist cough that is worse in the morning and during cold weather<sup>50</sup>. In advanced cases, blood vessels, air passages or alimentary tract may be obstructed by enlarged lymph nodes. Sometimes affected lymph nodes may rupture and drain. Involvement of the digestive tract is manifested by intermittent diarrhea and constipation in some cases. In terminal stage, animals may become severely emaciated and distress<sup>49</sup>. severe respiratory develop Tuberculosis mastitis is of major public health importance and difficult to differentiate it from other forms of mastitis<sup>31</sup>.

#### **Diagnosis:**

The increase in incidence of infection caused by *M. bovis* and *M. tuberculosis* in animals and humans has become the subject of investigation. The detection of early infection is dependent on cell mediated immunity as reflected in the response to diagnostic tests such as tuberculin test and cytokine assays. The introduction of molecular techniques has greatly reduced identification time and improved the level of detection in clinical specimens.

#### Smear microscopy and Histopathology:

Smear can be made from discharges of affected lymph nodes or nasal swabs and stained with Ziehl-Neelson'S (ZN) stain or auramine-rhodamine stain. The red coloured acid fast bacilli appears when examined under microscope after staining. But staining method have less sensitivity and specificity. During necropsy of suspected animal, tissue samples are collected and examined for histopathological lesions. The whitish hard caseous nodules present on different visceral organs and gritty sound will be there when slicing of these granulomas. Ziehl- Neelsen staining is done but to increase the sensitivity, auramine-rhodamine stain can also be used. The central to periphery of the nodule consists of necrosed mass, calcium deposits, acid fast bacilli, lymphocytes, epitheloid cells and fibrous connective tissue. The stained slides are observed under light microscope for the presence of red stained acid-fast bacilli<sup>57</sup>.

#### Culture isolation:

The culture isolation and biochemical characterization of *M. bovis* from apparently healthy animals shows active transmission in the infected herds<sup>20</sup>. The egg based LJ media and stone brinks media are most commonly used in veterinary bacteriology. An agar-based medium such as middle brook 7H10 and 7H11 or blood based agar medium may also be used. Mycobacterium tuberculosis, M. avium and many of the atypical mycobacteria require glycerol for growth and M. bovis grows in absence of glycerol and its growth can be supported by pyurvate. The luxuriant growth of M. tuberculosis on glycerol containing media, giving the characteristic rough, tough and buff colonies is known as eugenic, the growth of *M. avium* on media containing glycerol is also eugenic but M. bovis has sparse, thin growth on glycerol containing media which is called as  $dysgenic^{43}$ .

#### Tuberculin skin test:

Affected animals develop immune response, which can be detected by tuberculin skin test. Tuberculin is a purified protein derivative made by growing TB bacteria, heat killed followed by filteration or chemical treatment. The purified protein derivative (PPD) should have potency of 2000IU. In cattle with diminished allergic sensitivity, a higher dose of PPD is needed and the volume of each injection dose must not exceed 0.2ml<sup>32</sup>. After 72 hours of tuberculin intradermal injection, characteristic swelling appears at the point of injection. The diameter of the swollen area more than 4mm is a positive test result, indicating exposure to one type of mycobacteria. Different types of tuberculin skin test could be used to increase the

specificity such as comparative tuberculin skin test, stormont test, short thermal test  $etc^{17}$ .

#### Interferon Gamma (IFN-gamma) assay:

This in-vitro assay detects specific cell mediated immune response by the circulating lymphocytes. The use of defined Mycobacterium antigens such as ESAT-6 and promise CFP-10 shows for improved specificity. The use of these antigens may also offer the ability to differentiate BCGvaccinated from unvaccinated animals. The assay is based on the release of IFN-gamma from sensitized lymphocytes during 16-24 hours incubation period with specific antigen. The detection of bovine IFN-gamma is carried out with a sandwich ELISA that uses two monoclonal antibodies to bovine gammainterferon<sup>18</sup>.

#### Lymphocyte proliferation assay:

It is a more specific in-vitro assay which compares the reactivity of peripheral blood lymphocytes to PPD from *M. bovis* (PPD-B) and PPD from *M. avium* (PPD-A) and can be performed on whole blood or purified lymophocytes from peripheral blood samples. Results are usually analyzed as the difference in value obtained in response to PPD-B and PPD-A. It is not used for routine diagnosis because the test is time-consuming and the logistics are complex and the use of radioactive nucleotides<sup>46</sup>.

### Enzyme-linked immunosorbent assay (ELISA):

It appears to be the most suitable of antibodydetection tests and can be a complement, rather than an alternative, based on humoral immunity. It is a valuable complementary tool in-order to identify possible anergic cows that may be acting as reservoirs of the agent. Advantage lies in its simplicity, but limited sensitivity because of the late and irregular development of humoral immune response in cattle during the course of the disease. Improvement may be possible by using a combination of different antigens including proteins such as MPB 70 and MPB 83, which are specific but lack sensitivity<sup>32</sup>.

Polymerase chain reaction (PCR):

It is a very sensitive technique and can detect the presence of an organism when present at very low levels. It is used to detect the presence of genetic material (DNA) that is unique and specific to a particular organism and amplifies a portion of DNA that is specific for that organism and followed by gene sequencing<sup>34</sup>.

#### **Spoligotyping:**

It is also called spacer oligonucleotide typing, a method for simultaneously detection and typing of *M. tuberculosis* complex bacteria. This method is based on PCR amplification of highly polymorphic direct repeat (DR) locus in the *M. tuberculosis* genome. The DR region in *M. bovis* BCG contains direct repeat sequences of 36bp, which is interspersed by the non-repetitive DNA spacers of 35-41bp in length. Other MTC strains contain one or more IS6110 elements in DR-region<sup>37</sup>.

## Restriction fragment length polymorphism (RFLP):

It is considered as gold standard for the molecular typing of *M. tuberculosis* due to its high discriminative power and reproducibility. It can also be used for outbreaks identification and can facilitate contact tracing of tuberculosis. However, this technique requires large amount of DNA and is therefore restricted to the mycobacterial cultures which take around 20 to 40 days to obtain sufficient DNA needed<sup>27</sup>.

#### Treatment:

usually In animals, treatment is not recommended. Although anti-tuberculosis drug pyrazinamide is ineffective against M. bovis, the use of isoniazid and rifampicin could be used effectively. In developed countries, the reactor animals are sent to slaughter houses and in developing countries, the rector animals sent to gaushalas. But the reactor animals still remains a source of infection for other animals and human beings<sup>30</sup>.

#### **Prevention and control:**

*Bacillus Calmette* and *Guerin* (BCG) vaccine developed in 1921 is the commonly available live attenuated vaccine for tuberculosis till date. But this vaccine has shown variable

efficacy in cattle. In support with an international venture, a number of vaccines capable of replacing BCG with primary immunogens and also as boosters for BCG are being studied<sup>41</sup>. To reduce the risk for tuberculosis transmission, most of the highincome countries implement tuberculosis control programs but such control programs are not routinely implemented in India. However, in spite of intensive test and slaughter policy in many countries, a low level tuberculosis problem still remains that is difficult to eradicate because of wildlife reservoirs<sup>9</sup>. Variations in the tuberculin testing frequency, tuberculin types used and the interpretation of the test results affect the apparent incidence and prevalence of the disease recorded. The identification and early disposal of infected animals form the basis of national bovine tuberculosis eradication programme worldwide.

#### CONCLUSION

It is necessary to diagnose the disease at an early stage to minimize the risk of transmission to animals and human beings both. New diagnostic techniques are being used for early and efficient disease diagnosis but the molecular mechanisms behind the pathogen escape from immune system are complex. Vaccines other than BCG vaccine are also under clinical trial. Working association between physicians, private sector, and religious bodies local nonprofit organizations should be strengthened for better dissemination of awareness about the disease.

#### REFERENCES

- Acevedo, P., Romero, B., Vicente, J., Caracappa, S., Galluzzo, P., Marineo, S., Vicari, D., Torina, A., Casal, C., de la Fuente, J. and Gortazar, C., Tuberculosis Epidemiology in Islands: Insularity, Hosts and Trade, *PloS One.* 8 (7): 71074 (2013).
- Ayele, W. Y., Neill, S. D., Zinsstag, J., Weiss, M. G. and Pavlik, I., Bovine tuberculosis: an old disease but a new threat to Africa, *Int j Tuberc Lung Dis.* 8 (8): 924-937 (2004).

- Birhanu, T., Mezgebu, E., Ejeta, E., Gizachew, A. and Nekemte, E., Review on Diagnostic Techniques of Bovine Tuberculosis in Ethiopia, *Rep Opinion*. 7 (1): 7-14 (2015).
- 4. Bishai, W., The *Mycobacterium tuberculosis* genomic sequence: anatomy of a master adapter, *Trends Microbiol.* 6: 464-465 (1998).
- Bodnar, K. A., Serbina, N. V. and Flynn, J. L., Fate of *Mycobacterium tuberculosis* within murine dendritic cells, *Infection and Immunity*. 69 (2): 800-809 (2001).
- Buddle, B. M., Wedlock, D. N. and Denis, M., Progress in the development of tuberculosis vaccines for cattle and wildlife, *Vet Microbiol.* **112** (2-4): 313-323 (2006).
- Calmette, A., In Tuberculosis Bacillus Infection and Tuberculosis in Man and Animals, translated by Soper, W. B. and Smith, G.H., Williams and Wilkins Co. Baltimore (1923).
- Central TB Division. Revised National Tuberculosis Control Programme. Guidelines on Programmatic Management of Drug Resistant TB (PMDT) in India. New Delhi: Central TB Division, Directorate General of Health Services, Ministry of Health and Family Welfare, (2012).
- Corner and L.A., The Role of Wild Animal Populations in the Epidemiology of Tuberculosis in Domestic Animals: How to Assess the Risk, *Vet Microbiol.* 112 (2-4): 303-312 (2006).
- Cosovi, O.I., Grange, A. M., Daborn, C. J., Raviglione, M. C., Fujikura, T., Cousins, D., Robinson, R. A., Huchzermeyer, H. F., de Kantor, I., Meslin, F. X., Zoonotic tuberculosis due to *Mycobacterium bovis* in developing countries, *Emerg Infect Dis.* 4 (1): 59-70 (1998).
- Davies, P. D. O., Tuberculosis in humans and animals: are we a threat to each other? *J. R. Soc. Med.* 99 (10): 539-540 (2006).
- 12. Drewe, J. A., Pfeiffer, D. U. and Kaneene, J. B., Epidemiology of *Mycobacterium*

*bovis*. In: Thoen, C. O., Steele, J. H., Kaneene, J. B., Zoonotic Tuberculosis: *Mycobacterium* bovis and Other Pathogenic Mycobacteria: John Wiley & Sons (2014).

Rani *et al* 

- Dubos, R. and Dubos, J., The white plague. Tuberculosis, man and society. Boston: Little, Brown and Company, (1952).
- 14. Elias, K. H. D., Asseged, B., Wondwossen, T. and Gebeyehu, M., Status of bovine tuberculosis in Addis Ababa dairy farms, *Scientific and Technical Review of OIE*. **27** (3): 915-923 (2008).
- 15. Ereqat, S., Nasereddin, A., Levine, H., Azmi, K. and Al-Jawabreh, A., Greenblatt, C. L., Abdeen, Z., Bar-Gal, G. K., First-Time Detection of *Mycobacterium bovis* in Livestock Tissues and Milk in the West Bank, Palestinian Territories, *PLoS Negl Trop Dis.* 7 (9): e2417 (2013).
- Hertz, C. J., Kiertscher, S. M., Godowski, P. J., Bouis, D. A., Norgard, M. V., Roth, M. D., Modlin and R. L., Microbial lipopeptides stimulate dendritic cell maturation via Toll-like receptor 2, *J Immunol.* 166 (4): 2444–2450 (2001).
- 17. Iwnetu, R., Hombergh Van Den, J., Woldeamanuel. Y., Asfaw, M. and Gebrekirstos, С., Is tuberculous lymphadenitis over-diagnosed in Ethiopia? Comparative Performance of Diagnostic Tests for Mycobacterial Lymphadenitis in a High-Burden Country, Scand J Infect Dis. 41 (6-7): 462-468 (2009).
- Kubica, T., Agzamova, R., Wright, A., Rakishev, G., Rushgerdes, S. and Niemann, S., *Mycobacterium bovis* isolates with *Mycobacterium tuberculosis* specific characteristics, *Global Veterinarian*. 12 (5): 763-765 (2006).
- 19. Le Roex, N., van Helden, P.D., Koets, A.P. and Hoal, E.G., Bovine TB in Livestock and Wildlife: What's in the

631-637 (2013). 20. Leite, C. Q., Anno, I. S., Leite, S. R.,

Roxo, E., Morlock, G. P. and Cooksey, R.
C., Isolation and identification of *mycobacterium* from livestock specimens and milk obtained in Brazil, *Mem Inst Oswaldo Cruz.* 98 (3): 319-323 (2003).

Genes?, Physiological Genomics. 45 (15):

- Mbugi, E. V., Katale, B. Z., Kendall, S., Good, L., Kibiki, G. S., Keyyu, J. D., Godfrey-Faussett, P., Helden, P. and Matee, M. I., Tuberculosis cross-species transmission in Tanzania: Towards a One-Health concept, *Onderstepoort J Vet Res.* 79 (2): 501 (2012).
- Michel, A.L., Mueller, B. and Van helden, P. D., *Mycobacterium bovis* at the animalhuman interface: A problem or not, *Vet Microbiol.*, 140 (1-3): 371-381 (2010).
- Mohan, A. and Sharma, S. K., History, In: Sharma SK, Mohan A., *Tuberculosis*. 2nd ed. New Delhi: Jaypee Brothers Medical Publishers, 7-15 (2009).
- Narayanan, S., Das, S., Garg, R., Hari, L., Rao, V.B., Frieden, T.R., Molecular epidemiology of tuberculosis in a rural area of high prevalence in south India: implications for disease control and prevention, *J Clin Microbiol.* 40: 4785-4788 (2002).
- 25. O. I. E., Bovine Tuberculosis: Terrestrial Manual, Chapter 2.4.7. 1-16 (2009).
- 26. O.I.E., Terrestrial Manual (2009).
- O'Brian, R., Danilowicz, B. S., Bailey, L., Flynn, O., Costello, E. D., O'Grady and Rodgers, M., Characterization of the *Mycobacterium bovis* restriction fragment length polymorphism DNA probe pUCD and performance comparison with standard methods, *J Clin Microbiol.* 38 (9): 3362-3369 (2000).
- Ocepek, M., Pate, M., Zolnir-Dovc, M. and Poljak, M., Transmission of *Mycobacterium tuberculosis* from Human to Cattle, *J of Clin Microbiol.* 43 (7): 3555-3557 (2005).
- Palmer, M. V., Thacker, T. C., Waters, W. R., Gort, C. A. and Corner, L. A., *Mycobacterium bovis:* A Model Pathogen

Int. J. Pure App. Biosci. 5 (5): 1406-1414 (2017)

ISSN: 2320 - 7051

- at the Interface of Livestock, Wildlife, and Humans, Vet Med Int. 1: 17 (2012).
- 30. Peters, D., Farm Animals, Signs and Symptoms of Tuberculosis in Cattle. U.K. 1-8 (2010).
- 31. Radostits, O. M., Gay, C. C., Blood, D.C. and Hinchelift, K.W., Disease caused by bacteria-Mycobacterium. In: Veterinary Medicine: A Text Book of Disease of Cattle, Sheep, Pig, Goat and Horses, 9th ed., Harcourt Publ. Ltd., London, pp. 909-918 (2000).
- 32. Radostits, O., Blood, D. and Gray, C., Veterinary Medicine, a Text Book of the Diseases of cattle, sheep, Pigs, Goats and Houses, 8<sup>th</sup> ed., London: Ballier Tindals, 830-838 (2007a).
- 33. Ravenel, M. P., Intercommunicability of Human and Bovine Tuberculosis, J. Comp. Pathol. Thera. 55 (9): 140-147 (1902).
- 34. Regassa, A., Tassew, A., Amenu, K., Megersa, B., Abuna, F. and Mekibib, B., A cross-sectional study on bovine tuberculosis in Hawassa town and its surrounding Southern Ethiopia, Tropical Animal Health Production. 42: 915-920 (2010).
- 35. Rodriguez-Campos, S., Smith, N.H., Boniotti, M.B. and Aranaz, A., Overview and Phylogeny of *Mycobacterium* **Tuberculosis** *Complex* Organisms: Implications for Diagnostics and Legislation of Bovine Tuberculosis, Research in Veterinary Science, Suppl. 97: 5-19 (2014).
- 36. Rubin, S. A., Tuberculosis: Captain of all these men of death, Radiol Clin North Am. **33 (4):** 619-639 (1995).
- 37. Ruettger, A., Nieter, J., Skrypnyk, J., Engelmann, I., Ziegler, I., Moser, I., Monecke, S., Ehricht, R. and Sachse, K., Rapid Spoligotyping of Mycobacterium tuberculosis Complex Bacteria by Use of a Microarray System with Automatic Data Processing and Assignment, J Clin Microbiol. 50 (7): 2492-2495 (2012).
- 38. Russell and D. G. *Mycobacterium* tuberculosis: here today. and here Copyright © Sept.-Oct., 2017; IJPAB

- tomorrow, Nature Reviews Molecular Cell Biology. 2(8): 569-577 (2001).
- 39. Samuel, K. A., Oti, K. G., Ephraim, M. M., Isaac, K. A., Galyuon, Darlington, O., Bonsu, F. A., Bedzra, K. D., Gyasi, R. K., Slaughter surveillance for tuberculosis among cattle in three metropolitan abattoirs in Ghana, J of Vet and Ani Health. 6 (7): 198-207 (2014).
- 40. Shitaye, J. E., Tsegaye, W. and Pavlik, I., Bovine tuberculosis infection in animal and human populations in Ethiopia, Rev. Vet. Med. 52 (8): 317-332 (2007).
- 41. Singh, R., Rajni, Meena, A. and Meena, L. S., Multidrug resistant and Extensively drug resistant TB: A Nuisance to Medical Science, J Bacteriol Parasitol. 2: 105.doi:10.4172/2155-9597.1000105 (2011).
- 42. Smith, N. H., Gordon, S. V., de la Rua-Domenech, R., Clifton-Hadley, R. S. and Hewinson, R. G., Bottlenecks and broomsticks: the molecular evolution of Mycobacterium bovis, Nature Reviews Microbiol. 4 (9): 670-681 (2006).
- 43. Soolingen, van D., Molecular Epidemiology of Tuberculosis and other Mycobacterial infections: main methodologies and achievements. Journal of Veterinary Diagnostic Investigation, 249: 1-26 (2008).
- 44. Sreevatsan, S., Pan, X. I., Stockbauer, K. Connell, N. D., Kreiswirth, B. Е., N., Whittam, T. S., Musser, J. M., Restricted structural gene polymorphism in the Mycobacterium tuberculosis complex indicates evolutionarily recent global dissemination, Proc Natl Acad Sci USA. 94 (18): 9869-9874 (1997).
- 45. Tenguria, K. R., Khan, F. N., Quereshi, S. Pandey, A., Review Article and Epidemiological Study of Zoonotic Tuberculosis Complex (ZTBC), World J Sci Tech. 1 (3): 31-56 (2011).
- 46. Tewodros, F. and Girja, L., A review on diagnostic techniques of bovine tuberculosis, African Journal of Basic and Applied Sciences. 4 (6): 192-199 (2012).

Rani *et al* 

(2009).

- 5-1414 (2017) ISSN: 2320 7051 strains in Iran, *Chest.* **136 (2):** 420-425
- The centre for food security and public health (CFSPH), Bovine tuberculosis. Iowa state university, College of veterinary medicine (2009).

Rani *et al* 

- Udwadia, Z. F., Amale, R. A., Ajbani, K. K., Rodrigues, C., Totally drug-resistant tuberculosis in India, *Clin Infect Dis.* 54 (4): 579-581 (2012).
- 49. Une, Y. and Mori, T., Tuberculosis as a zoonosis from a veterinary perspective, *Comp Immunol, Microbiol, Infect Dis.* 30 (5-6): 415-425 (2007).
- Van Rhijn, I., Godfroid, J., Michel, A. and Rutten, V., Bovine Tuberculosis as a Model for Human Tuberculosis: Advantages over Small Animal Models, *Microbes Infect.* 10 (7): 711-715 (2008).
- Varello, K., Pezzolato, M., Mascarino, D. Ingravalle, F., Caranelli, M. and Bozzeta, E., Comparison of histological techniques for the diagnosis of bovine tuberculosis in the framework of eradication programs, *African Journal of Basic and Applied Sciences*, 1 (1-2): 26-30 (2006).
- Velayati, A. A., Masjedi, M. R., Farnia, P., Tabarsi, P., Ghanavi, J., Ziazarifi, A. H., Hoffner, S. E., Emergence of new forms of totally drug-resistant tuberculosis bacilli: super extensively drug-resistant tuberculosis or totally drug-resistant

- Vergne, I., Chua, J., Singh, S. B. and Deretic, V., Cell biology of *Mycobacterium tuberculosis* phagosome, *Annu Rev Cell Dev Biol.* 20: 367-394 (2004).
- 54. Verhagen, L. M., Hof, S. V. D., Duetekom, H. V., Harmans, P.W., Kremer, K., Borgdorff, M. W. and Soolingen, D. V., Mycobacterial factors relevant for transmission of tuberculosis, *J Infect Dis.* 203 (9): 1249-1255 (2011).
- 55. Verma, A. K., Tiwari, R., Chakraborty, S., Neha, Saminathan, M., Dhama, K. and Singh, S.V., Insights into Bovine Tuberculosis (bTB), Various Approaches for Its Diagnosis, Control and Its Public Health Concerns: An Updat, *Asian J of Ani and Vet Adv.* 9 (6): 323-344 (2014).
- 56. W. H.O., global report (2016).
- W.H.O., Global Tuberculosis Control. Geneva, WHO report 35-6 wide-needle aspiration in the diagnosis of Tuberculous Lymphadenitis in Africa, *AIDS*. 5: 213-298 (2012).
- World Health Organization, TB: A global emergency. WHO, Report on the TB epidemic. WHO/TB/94.177. Geneva: World Health Organization (1994).