INTERNATIONAL JOURNAL OF MYCOBACTERIOLOGY XXX (2016) XXX-XXX



Of Mycobacteriology

Available at www.sciencedirect.com

## **ScienceDirect**

journal homepage: www.elsevier.com/locate/IJMYCO



# Deletion of region of difference 181 in Mycobacterium tuberculosis Beijing strains

Ehsan Sharifipour<sup>a</sup>, Parissa Farnia<sup>a</sup>, Mohadeseh Mozafari<sup>a,\*</sup>, Shiva Irani<sup>b</sup>, Ali Akbar Velayati<sup>a</sup>

#### ARTICLE INFO

## Article history:

Received 14 September 2016 Accepted 20 September 2016 Available online xxxx

## Keywords:

Iran

Mycobacterium tuberculosis Region of differences

## $A\ B\ S\ T\ R\ A\ C\ T$

Objectives/background: The region of differences (RDs) polymorphisms is a potential molecular epidemiology method to distinguish origins of Mycobacterium tuberculosis. To date, 68 RDs have been identified in M. tuberculosis. This study was designed to determine the frequency of RD deletions in M. tuberculosis strains that were isolated from patients with pulmonary tuberculosis who were referred to the National Research Institute of Tuberculosis and Lung Disease for diagnosis and treatment. Therefore, highly polymorphic regions (RD1, RD150, and RD181) among M. tuberculosis strains isolates were investigated.

Methods: A total of 250 M. tuberculosis isolates were identified by conventional and molecular methods. Subsequently, spoligotyping and RD typing (RD1, RD150 and RD181) were performed to genotype these strains.

Results: The most frequent spoligotype belonged to Haarlem (n = 85, 34.0%) followed by CAS (n = 54, 21.6%), T1 (n = 27, 10.8%), and Beijing (n = 28, 11.2%) lineages. Deletion in RD181 was identified only among the Beijing lineage (Fig. 1).

Conclusion: As we found a deletion in RD181 in the Beijing strains only, we propose to use this marker as an identification tool for genotyping of the Beijing strain.

Peer review under responsibility of Asian African Society for Mycobacteriology. http://dx.doi.org/10.1016/j.ijmyco.2016.09.071

<sup>&</sup>lt;sup>a</sup> Mycobacteriology Research Centre, National Research Institute of Tuberculosis and Lung Disease (NRITLD), Shahid Beheshti University of Medical Sciences, Tehran, Iran

<sup>&</sup>lt;sup>b</sup> Department of Biology, Science and Research Branch, Islamic Azad University, Tehran, Iran

<sup>\*</sup> Corresponding author.

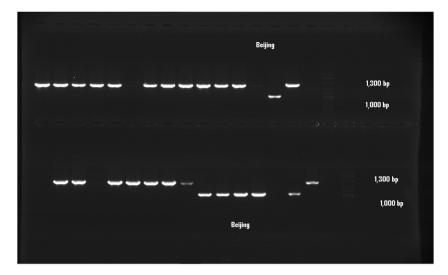


Fig. 1 – Electrophoresis fractionation of PCR products in 1% agarose gel to detect the existing deletion in RD181. The deletion can be diagnosed in the genomic region of RD181 in Beijing strains. The molecular weight of PCR products was 1000 bp. A molecular weight marker (100-bp ladder) was used. PCR = polymerase chain reaction; RD = region of difference.