

## عنوان مقاله: Genetic diversity of clinical isolates of Mycobacterium tuberculosis in Northeast of Iran

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## خلاصه مقاله:

AbstractBackground : Mycobacterium tuberculosis is still one of the most dangerous human pathogens. Identification of the relationships between different clinical strains has remained a high priority for epidemiology research. Methods : In this study, we used MLSA (Multilocus sequence analysis) to generate a highly robust phylogeny of M. tuberculosis. MLSA, based on single nucleotide polymorphism (SNP) was performed on five genes fragments from the Rpsl ( $\tau \cdot \tau$  bp), MprA ( $\Delta\Delta\Lambda$  bp), LipR ( $\tau \tau \tau$  bp), KatG ( $\tau \wedge \Delta$  bp) and Fgd $\iota$  ( $\tau \not \sim \rho$  bp), in order to identify polymorphic nucleotide sites, and the discriminatory power of each locus for all genes was measured with Hunter-Gaston Index (HGI). Results : In this study, a sequence type (ST) number was assigned to each unique allelic profile, and  $\Lambda$  sequence types were identified from  $\tau \cdot$  strains, these imply that there is a high diversity of strains in this area. Conclusion : Our results showed that the presence of high genetic diversity among clinical isolates of M. tuberculosis in Northeast of Iran. There is no evidence for recent transmission. Keywords : Mycobacterium tuberculosis, Multi-locus sequence analysis; Molecular epidemiology; Tuberculosis; KatG; Rpsl $\iota$ . IntroductionMycobacterium tuberculosis (M. tuberculosis), the causative agents of tuberculosis (TB), is one of the most successful human pathogens, infecting nearly one-third of the people all around the world, causing over  $\Lambda$  million new cases and  $\iota \cdot \tau$  million deaths each year [ $\iota - \tau$ ]. Identification of the .[relationships between different clinical strains of M. tuberculosis has great significance to the public health [ $\tau$ 

## كلمات كليدى:

Mycobacterium tuberculosis, Multi-locus sequence analysis, Molecular epidemiology, Tuberculosis, KatG, Rpsl

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