Targeted next generation sequencing on Algerian patients with limb girdle muscular dystrophy type 2

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Limb girdle muscular dystrophies type 2 (LGMD2) represent a large group of heterogeneous inherited muscle disorders. At present, the spectrum of these myopathies extends to more than 20 diseases. They are characterized by a high molecular heterogeneity, clinical overlaps, but a paucity of specific biomarkers. Indeed, even by critical clinical evaluation and muscle biopsy analyses, diagnosis is still difficult. To potentially remediate this difficulty, we applied targeted next generation sequencing technology to analyze 306 neuromuscular disorders associated genes. For this purpose, we studied 18 patients from five families presenting LGMD2 phenotype with ambiguous features for some patients. Putative pathogenic mutations were confirmed by Sanger sequencing. The data analysis of next generation sequencing done for the selected families allowed us to identify the putative causative molecular alterations in every family. Indeed, six different homozygous mutations were selected: 3 in the DYSF gene (c.5509G>A_p.Asp1837Asn, c.2643+1G>A et c.1834C>T_ p.Q612X), 1 in the LAMA2 gene (c.8244+1G>A), 1 in the GMPPB (c.458c>T_p.Thr153Ile) gene and 1 in the CPT2 (c.338C>T_p.Ser113Leu) gene. All these variants correlated well with the clinical features. Our result showed the accuracy and efficiency of next-generation sequencing in gene diagnosis of genetically heterogeneous diseases. It also demonstrated the usefulness of this approach in studying genes that would have been difficult to suspect following a clinical examination.

Biography

Benhassine T is currently working at University of Science and Technology Houari Boumediene, Algeria.

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