Advancement in the diagnosis of mitochondrial diseases

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Abstract

Mitochondrial diseases are multi-systemic, heterogeneous groups of diseases that are associated with various neuromuscular problems, cardiovascular disorders, metabolic syndrome, cancer, and obesity. Mitochondrial diseases are due to mutations in mitochondrial DNA (mtDNA) or nuclear DNA (nDNA) that encode mitochondrial components. Mitochondrial diseases are complex diseases involving multiple organ systems, and the symptoms include deafness, blindness, dementia, movement disorders, cardiovascular diseases, and renal dysfunction. Neurological and neuromuscular syndromes are the most common symptoms of mitochondrial diseases. However, clinical phenotype heterogeneity is a huge problem for the diagnosis of mitochondrial diseases, as patients with the same mutations exhibit different clinical symptoms. Also, the heteroplasmy/homoplasmy conditions complicate the diagnosis process. Here, in this review, we discuss these challenges and problems in mitochondrial disease diagnosis, focusing on the mutational profile of both primary and secondary mitochondrial diseases. We also review the utilization of next-generation technology and multi-omics strategy to improve the diagnosis. The discussion addresses the current evidence of those applications and the challenges that need some improvement for better diagnosis yield.

Keywords: Genomics, next-generation sequencing, mtDNA, mitochondrial diseases

INTRODUCTION

Mitochondrial diseases are caused by mutations in mitochondrial DNA (mtDNA) or nuclear DNA (nDNA) that encodes mitochondrial components. Mitochondrial diseases are complex diseases involving multiple organ systems, and the symptoms include deafness, blindness, dementia, movement disorders, cardiovascular diseases, and renal dysfunction. Neurological and neuromuscular syndromes are the most common symptoms of mitochondrial diseases. In addition, cardiovascular diseases, endocrine disorders,
and cancers may also be associated with mtDNA mutations\textsuperscript{[3,4]}. To date, there are about 350 causal genes for mitochondrial diseases\textsuperscript{[5]}. Since mitochondria play a significant role in energy production via the oxidative phosphorylation (OXPHOS) system and Krebs cycle, disruptions in their genome and protein functions may affect various important cellular processes such as fatty acid metabolism, pyrimidine biosynthesis, calcium homeostasis, cell signaling, beta-oxidation and heme biosynthesis\textsuperscript{[6]}.

Several factors are involved in determining disease onset, clinical symptoms, phenotypic variability, and mitochondrial penetrance\textsuperscript{[7]}. One such factor is replicative or mitotic segregation that leads to several conditions known as homoplasmy and heteroplasmy, threshold effect, clonal expansion, and the mtDNA bottleneck\textsuperscript{[8]}. The ratio of wild type and mutant mtDNA is important in determining disease onset and clinical symptoms. The vast majority of mtDNA mutations are present in the heteroplasmy condition, and only some mtDNA mutations are homoplasmic and deleterious. Mitochondria are randomly segregated; thus, there is a possibility that the daughter cells can shift from wild type to mutant and vice versa. A random genetic drift results in clonal expansion, leading to acceleration of mtDNA mutation rate\textsuperscript{[8]}. Rapid segregation of mammalian heteroplasmy with homoplasmy mtDNA between generations suggests that the mtDNA bottleneck occurs during development. Subsequently, this condition leads to differences in heteroplasmy levels in different mature oocytes of a woman\textsuperscript{[1]}.

mtDNA diseases have five unique characteristics, based on the mtDNA disease pedigree\textsuperscript{[1,7]}. First, mtDNA mutations are inherited maternally. Second, mutations arise as a result of several factors, including lack of histone protection, deficiency in DNA repair mechanisms, and increased levels of reactive oxygen species (ROS) through OXPHOS activities in the mitochondria. Pathogenic mtDNA mutations comprise rearrangement mutations and point mutations in genes affecting mitochondrial protein translation and causing specific OXPHOS defects. Moreover, replicative segregation of mitochondria gives rise to homoplasmic and heteroplasmic conditions. These conditions cause differential phenotypes in various processes such as the transcription, translation, enzyme complex formation, respiratory complexes, biochemical levels, and cellular phenotypes\textsuperscript{[1,4,7,8]}. Third, the impact of mtDNA mutations is subject to the amount of mitochondrial ATP production. In this case, tissues with the highest requirements for ATP can be affected the most, such as the central nervous system (CNS). Fourth, mtDNA repairs occur synonymously and replace mutations rapidly, about 5-10 times faster than nuclear OXPHOS genes due to mammalian mtDNA genes evolving faster compared to a single copy of nDNA. Finally, aging could result in decreased OXPHOS activities as well as mitochondrial dysfunction, which could be due to an accumulation of somatic cell mtDNA mutations\textsuperscript{[10]}.

The complexity and multi-systemic involvement in mitochondrial diseases render early diagnosis difficult. With ongoing advances in next-generation sequencing (NGS), the early diagnosis of mitochondrial diseases becomes feasible, and accurate diagnosis can be made even before the symptoms occur. In this review, we discuss the mitochondrial diseases, the challenges in their diagnosis, and future recommendations to assist in the diagnosis. We also provide the details of several companies that offer NGS services to diagnose mitochondrial diseases.

**OVERVIEW OF mtDNA AND MITOCHONDRIAL DISEASES**

**mtDNA**

Mitochondria have their own genome, which is known as mtDNA\textsuperscript{[11]}. The first complete sequence of the human mitochondrial genome was published in 1981\textsuperscript{[12]}. Following that, in 1999, the mtDNA sequence was revised, and its final full genomic sequence was published\textsuperscript{[13]}. In humans, mtDNA spans about 16,500 bp and consists of the heavy and light strands\textsuperscript{[12,13]}. The heavy strand is rich in guanine bases and encodes 12 subunits of the oxidative phosphorylation (OXPHOS) system, two ribosomal RNAs (12S and 16S), and 14 tRNAs. The light strand encodes one subunit of OXPHOS and eight tRNAs. Altogether, mtDNA contains...
37 genes encoding two rRNAs, 22 tRNAs, and 13 proteins subunits\[^{[12,13]}\]. To date, there are about 1000-2000 mitochondrial proteins; however, only 600 of these proteins have determined functions. mtDNA is a multicopy genome, ranging from just ~100 copies in sperm to > 100,000 copies in mature oocytes\[^{[14]}\]. The mtDNA genomes can exist either in homoplasmy (all genomes have an identical mtDNA genotype) or heteroplasmy (a combination of genomes with different mtDNA genotypes). These conditions can lead to various diseases or a normal phenotype, known as the phenotypic threshold effect. The inheritance of mtDNA is strictly maternal since mitochondrial endonuclease degrades the mtDNA within paternal mitochondria after fertilization\[^{[1,7,15]}\].

There are five complexes in the OXPHOS system, namely complexes I (NADH ubiquinone oxidoreductase), complex II (succinate ubiquinone oxidoreductase), complex III (ubiquinone-cytochrome c reductase), complex IV (cytochrome c oxidase) and complex V (ATP synthase) which is important for energy production\[^{[6,12]}\]. Embedded in this system are two electron carriers, ubiquinone coenzyme Q and cytochrome C, which serve as electron transporters important for energy production. Leakage of electrons from complexes I and III in the electron transport chain (ETC) can lead to the formation of ROS and oxidative stress conditions\[^{[9]}\]. Oxidative stress occurs where there is an imbalance between the production of ROS, such as superoxide anion, H\(_2\)O\(_2\), and the antioxidant defense mechanism to remove these radical molecules\[^{[6]}\]. Uncontrolled oxidative stress may induce various dysfunctions within the mitochondria and cells, which could then lead to disease development [Figure 1].
Different types of mitochondrial diseases

Mutations of mtDNA or nuclear DNA (nDNA) that encode mitochondria components lead to an ineffective mitochondrial respiratory chain, and hence mitochondrial dysfunction [1,7,15]. The mitochondrial respiratory chain is an essential pathway for aerobic metabolism. Tissues and organs that are highly dependent on aerobic metabolism are prone to mitochondrial diseases [17]. Mitochondrial diseases can affect single or multiple organ systems [1,7,15], thus the association with various disease conditions including neurodegenerative conditions, cardiovascular diseases, neurometabolic problems, cancer and metabolic diseases such as obesity [3,4,10]. Mitochondrial diseases are considered rare, with a prevalence of 1:1,000,000. However, recent epidemiological studies suggested that at least 1 in 5000 individuals are affected by mitochondrial diseases [18,19]. Mitochondrial diseases may occur at any age [20], where mutations in mtDNA account for about 75% of adult diseases [18] but only about 25% of childhood-onset diseases [21,22]. Initially, it was postulated that nDNA abnormalities are present in childhood, while mtDNA abnormalities are only present in late childhood. Recently, many mtDNA diseases also present in childhood, and nDNA mutations are also present in adult life [23]. Mitochondrial diseases are classified into several categories, including primary and secondary mitochondrial diseases (SMD), which are discussed in detail in the next section [Figure 2].

**Primary mitochondrial disease**

As the mitochondrial components are inter-dependent of mtDNA and nDNA, most of the primary mitochondria diseases are due to these components. About 90% of mitochondrial proteins are from nuclear genes, which include the majority of the OXPHOS system. Also, mtDNA replication, transcription, and translation are dependent on these nuclear-encoded proteins. Primary mitochondrial diseases (PMD) are genetically inherited and diagnosed by identifying mutations in the mtDNA or nDNA [21], which could result in mitochondrial dysfunction [Tables 1 and 2]. Such examples are mitochondrial encephalomyopathy, lactic acidosis and stroke-like episodes (MELAS), neuropathy, ataxia and retinitis pigmentosa (NARP), Leigh syndrome, myoclonic epilepsy with ragged-red fibers (MERRF), Leber hereditary optic neuropathy (LHON), Kern-Sayre syndrome (KSS), mitochondrial neurogastrointestinal encephalopathy (MNGIE), and Alpers.

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**Figure 2.** Common diseases associated with mtDNA mutations and the genes involved [24]. LHON: leber hereditary optic neuropathy; CPEO: chronic progressive external ophthalmoplegia; MERRF: myoclonic epilepsy with ragged-red fibers; MND: motor neuron disease; MELAS: mitochondrial encephalomyopathy, lactic acidosis, and stroke-like episodes; PD: Parkinson’s disease.
| Syndrome                           | Locus       | Disease                        | Allele       | NT change | AA change | Ho | He | Ref.               |
|-----------------------------------|-------------|--------------------------------|--------------|-----------|-----------|----|----|--------------------|
| Primary mitochondrial disease     |             |                                |              |           |           |    |    |                    |
| Leigh syndrome/dystonia           | MTATP6     | LS/NARP                        | T8993C       | T-C       | L156P     | -  | +  | [48]               |
| Leigh syndrome                    | MTATP6     | NARP                           | T8993G       | T-G       | L156R     | -  | +  | [49,50]            |
| Leigh syndrome                    | MTATP6     | LS                             | T9176G       | T-G       | L217R     | +  |    | [51]               |
| Leigh syndrome                    | MTATP6     | LS/FBSN                        | T9176C       | T-C       | L217P     | +  |    | [52,53]            |
| Leigh syndrome/ataxia/NARP-like   | MTATP6     | LS/ataxia/NARP-like disease    | T9185C       | T-C       | L220P     | +  |    | [54]               |
| Leigh syndrome                    | MTATP6     | LS                             | T9191C       | T-C       | L222P     | +  |    | [54]               |
| Leigh syndrome                    | MTND3      | LS                             | T10158C      | T-C       | S34P      | +  |    | [22,56,57]         |
| Leigh syndrome                    | MTND3      | LS/LS-like disease/ESOC        | T10191C      | T-C       | S45P      | +  |    | [58]               |
| Leber hereditary optic neuropathy | ND1        | LHON                           | G3460A       | G-A       | A52T      | -  |    | [69,70]            |
| Leber hereditary optic neuropathy | ND4        | LHON                           | G11778A      | G-A       | R340H     | -  |    | [71]               |
| Leber hereditary optic neuropathy | ND6        | LHON                           | T14484C      | T-C       | M64V      | -  |    | [69,72,73]         |
| Encephalomyopathy                 | MTND1      | MELAS                          | T3308C       | T-C       | M1T       | -  |    | [74]               |
| Encephalomyopathy, MELAS          | MTND1      | MELAS/LHON                     | G3376A       | G-A       | E24K      | -  |    | [75]               |
| Encephalomyopathy, MELAS          | MTND1      | MELAS/LHON/LS                  | G3946A       | G-A       | E214K     | +  |    | [60]               |
| Encephalomyopathy, MELAS          | MTND1      | MELAS/LHON/LS                  | T3949C       | T-C       | Y215H     | -  |    | [60]               |
| Encephalomyopathy, MELAS          | MTND4      | MELAS                          | A1084G       | A-G       | T109A     | +  |    | [76,77]            |
| Encephalomyopathy, MELAS          | MTND5      | MELAS                          | A12770G      | A-G       | E145G     | -  |    | [78]               |
| Encephalomyopathy, MELAS          | MTND5      | MELAS/LHON/LS                  | A13045C      | A-C       | M237L     | -  |    | [78]               |
| Encephalomyopathy, MELAS          | MTND5      | MELAS/LHON/LS                  | A13084T      | A-T       | S250C     | -  |    | [79]               |
| Encephalomyopathy, MELAS          | MTND5      | MELAS/LS                       | G13513A      | G-A       | D393N     | -  |    | [80]               |
| Encephalomyopathy, MELAS          | MTND5      | MELAS/LS                       | A13514G      | G-A       | D393G     | -  |    | [81]               |
| Encephalomyopathy, MELAS          | MTND6      | MELAS                          | G14453A      | G-A       | A74V      | -  |    | [82]               |
| Encephalomyopathy, MELAS          | MTYB       | MELAS/PD                       | G1787delI4   | TCAA-del  | I14frameshift | -  | [83] |
| Encephalomyopathy, MELAS          | MTTL1      | MELAS                          | A3243G       | A-G       | rRNA        | -  | [84] |
| Encephalomyopathy, epilepsy       | MTCO1      | Therapy-resistant epilepsy     | C6489A       | C-A       | L196L     | -  |    | [85]               |
| Encephalomyopathy, multisystem disorder | MTCO1  | Multisystem disorder           | G6930A       | G-A       | G343Ter   | -  |    | [86]               |
| Encephalomyopathy, multisystem disorder | MTCO1  | Myopathy and cortical lesions  | 6015del5     | Del 5 bp  | Frameshift, 42 peptide | -  | [87] |
| Encephalomyopathy                 | MTCO2      | Encephalomyopathy              | T7587C       | T-C       | M1T       | -  |    | [88]               |
| Encephalomyopathy, lactic acidosis | MTCO2     | Multisystem disorder           | G7896A       | G-A       | W104Ter   | -  |    | [89]               |
| Encephalomyopathy                 | MTCO3      | Lactic acidosis                | 8042del2     | AT-del    | M153Ter   | -  |    | [90]               |
| Encephalomyopathy, MELAS          | MTCO3      | Lactic acidosis/MELAS/PEM/NAION| T9957C       | T-C       | F251L     | -  |    | [92]               |
| Encephalomyopathy, lactic acidosis | MTAP7b     | Lactic acidosis/seizures       | 9205del2     | TA-del    | Ter227M   | +  |    | [93]               |
| Encephalomyopathy, multisystem disorder | MTYB   | Multisystem disorder           | A15579G      | A-G       | Y278C     | -  |    | [94]               |
| Mitochondrial myopathy            | MTYB       | Septo-optic dysplasia          | T14849C      | T-C       | S35P      | -  |    | [95]               |
| MM, exercise intolerance          | MTYB       | EXIT                          | G14846A      | G-A       | G34S      | -  |    | [96]               |
| MM, exercise intolerance          | MTYB       | MM                            | G15059A      | G-A       | G901Ter   | -  |    | [96]               |
| MM, exercise intolerance          | MTYB       | EXIT                          | G15084A      | G-A       | W113Ter   | -  |    | [96]               |
| MM, exercise intolerance          | MTYB       | EXIT                          | G15150A      | G-A       | W135Ter   | -  |    | [97]               |
| MM, exercise intolerance          | MTYB       | EXIT                          | G15168A      | G-A       | W141Ter   | -  |    | [96]               |
| MM, exercise intolerance          | MTYB       | EXIT/encephalomyopathy        | G15242A      | G-A       | G166Ter   | -  |    | [98]               |
| Region | Gene/Locus | OMIM | Function | Chromosome | Inheritance | Clinical phenotype |
|--------|------------|------|----------|------------|-------------|-------------------|
| Complex I | ACAD9 | 611103 | Assembly and activity | 3q26 | AR | Hypertrophic cardiomyopathy encephalopathy |
| | FOXRED1 | 613622 | Assembly | 11q24.2 | AR | LS |
| | NUBPL | 613621 | Assembly | 14q12 | AR | Encephalomyopathy |
| | NDUF1 | 300078 | HP fraction | Xq24 | X-linked | LS, progressive neurodegenerative disorder |
| | NDUF2 | 602137 | HP fraction | 5q31.2 | AR | LS |
| | NDUF3 | 603834 | HP fraction | 12p13.32 | AR | LS |
| | NDUF4 | 603835 | HP fraction | 2q37.3 | AR | LS |
| | NDUF5 | 612638 | IP fraction | 19p13.3 | AR | Fatal infantile lactic acidosis, encephalocardiomyopathy |
| | NDUF6 | 609653 | HP fraction | 12q22 | AR | LS |
| | NDUF7 | 609435 | HP fraction | 19p13.31 | AR | Encephalopathy, optic atrophy |
| | NDUF8(CIA30) | 606934 | Assembly | 15q13.3 | AR | Cardiomyopathy |
| | NDUF9 | 609653 | Assembly | 15q13.3 | AR | Early-onset progressive encephalopathy |
| | NDUF10 | 612911 | Assembly | 3p21.31 | AR | Neonatal encephalopathy |
| | NDUF11A (HRPAP2) | 611776 | Assembly | 6q16.1 | AR | Infantile encephalopathy |

AD: Alzheimer’s disease; ADPD: Alzheimer’s plus Parkinson’s diseases; CPEO: chronic progressive ophthalmoplegia; DEAF: deafness, sensorineural hearing loss; ESOC: epilepsy, strokes, optic atrophy and cognitive decline; EXIT: exercise intolerance; FBSN: familial bilateral striatal necrosis; HCM: hypertrophic cardiomyopathy; LHON: Leber hereditary optic neuropathy; MM: mitochondrial myopathy; MERRF: myoclonic epilepsy with ragged-red fibers disease; MELAS: mitochondrial encephalopathy, lactic acidosis and stroke-like episodes; NAION: non-arteritic anterior ischemic optic neuropathy; NARP: neurogenic muscle weakness, ataxia and retinitis pigmentosa; NIDDM: non-insulin dependent diabetes mellitus; PEM: progressive external ophthalmoplegia; PD: Parkinson’s disease; SNHL: sensorineural hearing loss; SIDA: sideroblastic anemia; Ho: homoplasmic; He: heteroplasmic; Rep: reported/provisional mutations that have been reported one or a few times in association with disease but are not confirmed as pathogenic; P.M.: (point mutation/polymorphism) status indicates that some published reports have determined the mutation to be a non-pathogenic population variant.
NDUFAF5 (C20orf17) 612360 Assembly 20p12.1 AR LS
NDUFAF6 612392 Assembly 8q22.1 AR LS
NDUF83 603839 HP fraction 2q31.3 AR Fatal infantile lactic acidosis
NDUF89 601445 HP fraction 8q24.13 AR Hypotonia, lactic acidosis
NDUF10 603843 HP fraction 16p13.3 AR Lactic acidosis, cardiomyopathy
NDUF11 300403 HP fraction Xp11.3 X-linked Intrauterine growth restriction, lactic acidosis
NDUF51 157665 IP fraction 2q33-q34 AR LS
NDUF52 602985 IP fraction 1q23 AR Encephalopathy, cardiomyopathy
NDUF53 603846 IP fraction 11p11.11 AR LS
NDUF54 602694 IP fraction 5q11.1 AR LS
NDUF56 603848 IP fraction 8pter-p15.33 AR Fatal infantile lactic acidosis
NDUF57 601825 HP fraction 19p13.3 AR LS
NDUF58 602141 HP fraction 11q13 AR LS
NDUFV1 161015 FP fraction 11q13 AR LS
NDUFV2 600532 FP fraction 18p11 AR Cardiomyopathy, hypotonia, encephalopathy

Complex II SDH-A 600857 FP subunit 5p15 AR LS
SDH-B 185470 IP subunit 1p36.1-p35 AD Phaeochromocytoma and paraganglioma
SDH-C 602413 Membrane subunit 1q21 AD Autosomal dominant paraganglioma type 3
SDH-D 602690 Membrane subunit 11q23 AD Autosomal dominant paraganglioma type 1, phaeochromocytoma
SDHA4F1 612848 Assembly 19q12-q13.2 AR Leukoencephalopathy
SDHA4F2 613019 Assembly 11q12.2 AD Autosomal dominant paraganglioma type 2

Complex III BCS1L 603647 Assembly 2q33 AR Encephalopathy, hepatic failure and tubulopathy, LS, GRACLE syndrome, Bjornstad syndrome

Complex IV UQCRC2 614461 Assembly 6p21.3 AR Lactic acidosis and renal tubular dysfunction
UQCRC3 616097 Assembly 1q2.3 AR Lactic acidosis, hypoglycemia, hypotonia
UQCR10 191330 Electron transfer 1p36.1-p35 AD Hypoglycemia, lactic acidosis
UQCRQ 612080 Electron transfer 5q31.1 AR Severe neurological phenotype

Complex V ATP5E 606153 ATPase activity 20q13.3 AR Neonatal lactic acidosis, mental retardation, peripheral neuropathy
ATP5A1 164360 ATPase activity 18q21.1 AR Neonatal encephalopathy
ATP8A2 605870 ATPase activity 13q12.13 AR Cerebellar ataxia, mental retardation
ATP12A2 608981 Assembly 17q11.2 AR Neonatal encephalopathy, lactic acidosis
TMEM70 604273 Assembly 8q21.11 AR Neonatal encephalopathy, cardiomyopathy
| Gene   | Description                                                                 | Location | Inheritance | phenotypes                                                                 |
|--------|------------------------------------------------------------------------------|----------|-------------|-----------------------------------------------------------------------------|
| ANT1   | Adenine nucleotide translocator isoform 1                                  | 4q35     | AD-AR       | AD-PEO, multiple mtDNA deletions                                           |
| C10orf75 | Twinkle helicase                                                            | 10q24    | AD          | AD-PEO, SANDO syndrome                                                     |
| DGUOK  | Deoxyguanosine kinase                                                        | 2p13     | AR          | Hepatocerebral mtDNA depletion syndrome                                    |
| FBXL4  | Mitochondrial dNTP pool maintenance                                          | 6q16.1-q16.2 | AR      | Encephalomyopathy and myopathy, mtDNA depletion                            |
| MFN1   | Mitofusin, mitochondrial fusion                                              | 1p36-p35 | AD          | Charcot-Marie-Tooth disease-2A2 (CMT2A2), multiple deletions              |
| MGME1  | mtDNA maintenance                                                            | 20p11.23 | AR          | CPEO and myopathy, mtDNA deletion                                         |
| MPV17  | Regulation of mtDNA copy number                                              | 2p23-p21 | AR          | Hepatocerebral MDDS                                                        |
| OPA1   | Dynamin-related protein                                                      | 3q28-q29 | AR          | AD-optic atrophy, multiple deletions                                      |
| POLG   | Polymerase gamma mtDNA replication                                           | 17q23-q24 | AD          | Alpers syndrome, AD-PEO, and AR-PEO, male infertility, SANDO syndrome, SCAE* AD-PEO |
| POLG2  | Catalytic subunit of DNA polymerase gamma                                    | 8q23.1   | AR          | Encephalomyopathy, renal tubulopathy, MNIGE, AD-PEO                       |
| RRM2B  | Ribonucleotide reductase M2 B dNTP pool                                     | 13q12.2-q13 | AR      | Encephalomyopathy with methylmalonic aciduria                            |
| SUCLA2 | Succinate-CoA ligase, ADP-forming, beta subunit                              | 2q11.2-q11 | AR          | Encephalomyopathy with methylmalonic aciduria                            |
| SUCLG1 | Succinate-CoA ligase, alpha subunit                                          | 1q41     | X-linked    | Deafness-dystonia or Mohr-Tranebjaerg syndrome                            |
| TFAM   | Mitochondrial transcription factor A                                         | 1q25     | AD-AR       | Leukoencephalopathy and lactic acidosis                                    |
| TK2    | Thymidine kinase MtDNA synthetase                                            | 16q22    | AR          | Leukoencephalopathy                                                        |
| TYMP   | Thymidine phosphorylase                                                      | 3p26.3   | AR          | Leukoencephalopathy                                                        |
| AARS2  | Alanyl-tRNA synthetase                                                       | Xq22     | X-linked    | Cardiomyopathy, leuoencephalopathy                                       |
| CARS2  | Cysteinyl-tRNA synthetase                                                    | 13q34    | AR          | Myoclonic epilepsy                                                        |
| C12orf65 | Mitochondrial translation                                                    | 12q24.31 | AR          | Encephalomyopathy, optic atrophy, axonal neuropathy, paraparesis          |
| DARS2  | Aspartyl-tRNA synthetase                                                     | 1q25.1   | AR          | Leukoencephalopathy and lactic acidosis                                    |
| EARS2  | Glutamyl tRNA synthetase                                                     | 16p12.2  | AR          | Leukoencephalopathy                                                        |
| EFG1   | Elongation factor G1 mitochondrial translation defect                        | 3q25     | AR          | Severe hepato-encephalopathy and lactic acidosis                          |
| FARS2  | Phenylalanyl-tRNA synthetase                                                 | 6p25.1   | AR          | Alpers syndrome, spastic paraplegia                                       |
| GARS   | Glycyl-tRNA synthetase                                                       | 7p14.3   | AD          | Charcot-Marie-Tooth disease                                               |
| GFM1   | Mitochondrial translation elongation                                          | 3q25.32  | AR          | Encephalopathy/hepatic failure                                             |
| GFM2   | Mitochondrial translation elongation                                          | 5q13.3   | AR          | Neurodevelopmental disorder, dysmorphic features                          |
| GTPBP3 | GTP-binding protein                                                          | 19p13.11 | AR          | Cardiomyopathy, encephalopathy                                            |
| HARS2  | Histidyl-tRNA synthetase                                                     | 5q31.3   | AR          | Perrault syndrome                                                         |
| IARS2  | Isoleucyl tRNA-Synthetase                                                    | 1q41     | AR          | Cataract, deafness, neuropathy/Leigh syndrome                             |
| KARS   | Lysyl-tRNA synthetase                                                        | 16q23.1  | AR          | CMT disease/deafness                                                       |
| LARS   | Leucine-tRNA synthetase                                                      | 5q32     | AR          | Hepatopathy                                                               |
| LARS2  | Leucyl-tRNA synthetase                                                       | 3p21.31  | AR          | Perrault syndrome                                                         |
| Gene    | Reference ID | Function                                      | Chromosome | Disorder                                                                 |
|---------|--------------|-----------------------------------------------|------------|---------------------------------------------------------------------------|
| MRPL3   | 607118       | Mitochondrial translation                      | 3q22.1     | AR Cardiomyopathy, mental retardation                                      |
| MRPS7   | 611974       | Mitochondrial translation                      | 17q25.1    | AR Deafness, hepatic and renal failure                                     |
| MRPL12  | 602375       | Mitochondrial translation                      | 17q25.3    | AR Growth retardation, encephalopathy                                     |
| MRPS16  | 609204       | Mitochondrial translation                      | 10q22.1    | AR Neonatal lactic acidosis, corpus callosum agenesis                     |
| MRPS22  | 605810       | Mitochondrial translation                      | 3q23       | AR Cardiomyopathy, tubulopathy                                           |
| MRPL44  | 611849       | Mitochondrial translation                      | 2q36.1     | AR Cardiomyopathy                                                         |
| MTFMT   | 611766       | Mitochondrial translation                      | 15q22.31   | AR LS                                                                     |
| MT01    | 614667       | tRNA modification                              | 6q13       | AR Cardiomyopathy                                                         |
| NARS2   | 612803       | Asparaginyl-tRNA synthetase                    | 11q14.1    | AR Alpers syndrome/nonsyndromic deafness and Leigh syndrome               |
| PARS2   | 612036       | Prolyl-tRNA synthetase                         | 1p32.3     | AR Alpers syndrome                                                        |
| RARS2   | 611523       | Arginyl-tRNA synthetase                        | 6q16.1     | AR Pontocerebellar hypoplasia                                              |
| RMND1   | 614917       | Mitochondrial translation                      | 6q25.1     | AR Encephalopathy                                                         |
| SARS2   | 612804       | seryl-tRNA synthetase                          | 19q13.2    | AR Hyperuricemia, pulmonary hypertension, renal failure                   |
| TARS2   | 612805       | Threonyl-tRNA synthetase                       | 1q21.2     | AR Encephalomyopathy                                                      |
| TSFM    | 604723       | Mitochondrial translation elongation           | 12q13-q14  | AR Encephalomyopathy, hypertrophic cardiomyopathy                         |
| TUFM    | 602389       | Mitochondrial translation elongation           | 16p11.2    | AR Leukodystrophy with micropolygyria                                     |
| TRMT5   | 611023       | Mitochondrial RNA methyltransferase            | 14q23.1    | AR Cardiomyopathy/exercise intolerance                                     |
| TRMT10C | 615423       | tRNA methyltransferase                         | 3q12.3     | AR Hypotonia, feeding difficulties, deafness                              |
| TRMU    | 610230       | Mitochondrial translation                      | 22q13.31   | AR Liver failure, deafness                                                |
| VARS2   | 612802       | Valyl-tRNA synthetase                          | 12p11.21   | AR Myopathy, lactic acidosis, and sideroblastic anemia-2                   |
| ABCB7   | 301310       | Iron transport                                 | Xq13.1-q13.3| X-linked sideroblastic anemia with ataxia                                 |
| BOLA3   | 613183       | Iron-sulfur cluster biosynthesis               | 2p13.1     | AR Encephalomyopathy, cardiomyopathy                                      |
| FDXL1   | 614585       | Iron-sulfur cluster biosynthesis               | 19p13.2    | AR Myopathy, lactic acidosis                                               |
| FRDA (FXN)| 606829     | Frataxin修剪.* repeat,                       | 9q13       | AR Friedreich ataxia, neuropathy, cardiomyopathy, diabetes                |
| GLRX5   | 205950       | Iron-sulfur cluster biosynthesis               | 3p22.1     | AR Sideroblastic anemia                                                    |
| IBA57   | 615316       | Iron-sulfur cluster biosynthesis               | 1q42.13    | AR Myopathy, encephalopathy                                               |
| ISCA2   | 615317       | Iron-sulfur cluster biosynthesis               | 14q24.3    | AR Leukodystrophy                                                         |
| ISCU    | 255125       | Iron-sulfur cluster biosynthesis               | 12q23.3    | AR Myopathy, lactic acidosis, exercise intolerance                         |
| LYRM4   | 613311       | Iron-sulfur cluster biosynthesis               | 6q25.1     | AR Lactic acidosis, Failure to thrive                                     |
| LYRM7   | 615831       | Iron-sulfur cluster biosynthesis               | 5q23.3-q31.1| AR Encephalomyopathy, lactic acidosis                                     |
| NUF1    | 608100       | Iron-sulfur cluster biosynthesis               | 2p13.3     | AR Lactic acidosis, multiple respiratory chain deficiency                   |
| APTX    | 606350       | CoQ10 deficiency                               | 9p13.3     | AR Cerebellar ataxia, Oculomotor apraxia                                  |
| CABC1   | 606980       | CoQ10 deficiency                               | 1q42.2     | AR Cerebellar ataxia, lactic acidosis                                      |
| COQ2    | 609825       | CoQ10 deficiency                               | 4q21-q22   | AR Encephalomyopathy, nephropathy                                         |
| COQ4    | 612898       | CoQ10 deficiency                               | 9q34.13    | AR Encephalomyopathy, mental retardation                                  |
| COQ5    | 616359       | CoQ10 deficiency                               | 12q24.31   | AR Encephalomyopathy, cerebellar ataxia                                   |
| COQ6    | 614647       | CoQ10 deficiency                               | 14q24.3    | AR Nephrotic syndrome, deafness                                           |
| COQ7    | 601683       | CoQ10 deficiency                               | 16p12.3    | AR Hypotonia, cardiac hypertrophy                                         |
MELAS disease is diagnosed by the presence of mitochondrial myopathy, encephalopathy, lactic acidosis, and stroke-like episodes. The m.3243A>G mutation in the MT-TL1 gene encoding tRNALEU(UUR) was identified to be the cause of MELAS in 1990. This mutation causes translational defects of the OXPHOS protein assembly and the lack of complex I (CI), which leads to an accumulation of nicotinamide adenine dinucleotide (NAD)+ hydrogen (H) (NADH), increased glycolysis, oxidative stress, and reduced ATP production. Similarly, NARP disease is also associated with energy production due to a mutation in the ATP6 gene (m.9176 T>G). This mutation leads to a heterogeneous and complex disease manifestation, from having adult-onset NARP to fatal infantile subacute necrotizing encephalomyopathy. A study performed by Mordel and colleagues showed that a novel and pathogenic 2-bp microdeletion (m9127-9128 del AT) in the ATP6 gene caused NARP. This heteroplasmy mutation was highly abundant in the muscle (82%), followed by fibroblasts (50%) and blood (10-20%). Functional studies showed that oligomycin-sensitive ATPase hydrolytic activity was reduced at 60%, and ATP synthesis was decreased at 40%

The most common mitochondrial disease in children is Leigh syndrome, with a prevalence of 1:40,000. The m.3243A>G mutation in the MT-TL1 gene encoding tRNALEU(UUR) was identified to be the cause of MELAS in 1990. This mutation causes translational defects of the OXPHOS protein assembly and the lack of complex I (CI), which leads to an accumulation of nicotinamide adenine dinucleotide (NAD)+ hydrogen (H) (NADH), increased glycolysis, oxidative stress, and reduced ATP production. Similarly, NARP disease is also associated with energy production due to a mutation in the ATP6 gene (m.9176 T>G). This mutation leads to a heterogeneous and complex disease manifestation, from having adult-onset NARP to fatal infantile subacute necrotizing encephalomyopathy. A study performed by Mordel and colleagues showed that a novel and pathogenic 2-bp microdeletion (m9127-9128 del AT) in the ATP6 gene caused NARP. This heteroplasmy mutation was highly abundant in the muscle (82%), followed by fibroblasts (50%) and blood (10-20%). Functional studies showed that oligomycin-sensitive ATPase hydrolytic activity was reduced at 60%, and ATP synthesis was decreased at 40%

The most common mitochondrial disease in children is Leigh syndrome, with a prevalence of 1:40,000. The diagnosis is based on neuroimaging and characterized by symmetrical lesions in the basal ganglia, thalamus, and brain stem. Patients with Leigh syndrome usually present with loss of acquired cognitive, visual as well as motor skills. Mutations in both mtDNA and nDNA could lead to mitochondrial dysfunctions in different aspects. For example, mutations in NDUF52 and SURF1 (structural subunits and assembly factors), PHHA1 (Kreb’s cycle components), MTFM (mitochondrial protein translation), and ECHS1 (valine metabolism), are all reported to be associated with Leigh syndrome. Another similar disease is MERRF, which affects 1 in 400,000 individuals and involves the nervous and skeletal muscle systems. MERRF is characterized by myoclonus epilepsy, ataxia, seizures, and myopathy symptoms. The most common mutation is the mtDNA tRNALys A8344G, which accounts for about 80%-90% of all MERRF cases. Another common mutation, causing LHON disease, occurs 1 in 31,000-50,000 people. Clinically, it is characterized by bilateral loss of central vision and some other abnormalities such as movement disorders, dystonia, or multiple sclerosis-like symptoms. Mutations in the NADH dehydrogenase account for 90% of all LHON cases, and the most common mutations are m.11778G>A (ND4), m.14484T>C (ND6) and m.3460G>A (ND1). LHON disease can lead to blindness due to the loss of retinal ganglion cells from the mtDNA mutations affecting the respiratory complex I (CI) subunits, such as m.3460G>A, m.11778G>A and m.14484T>C. There is also a rare mutation such as m.13094T>C reported in LHON disease.
Another PMD with eye disorder is the Kern-Sayre syndrome (KSS), which is characterized by a progressive external ophthalmoplegia, a condition that causes eye muscle paralysis. The patients may also have other symptoms such as cardiac conduction defect, ataxia, and abnormalities in protein levels of the cerebrospinal fluid. The disease affects 1-3 in 100,000 individuals, and most KSS cases are due to mtDNA mutations. Some of the mutations that have been identified in KSS are the deletion of 3,236 bp of mitochondrial chromosome from 10,170 \[35\], mtDNA duplication \[36\], a mtDNA point mutation in the MTTL2 gene \[37\], and a large mtDNA deletion \[38\]. Mitochondrial neurogastrointestinal encephalopathy (MNGIE) disease affects several parts of the body, mostly the digestive and nervous systems \[39\]. It is caused by a mutation in the thymidine phosphorylase gene (TYMP) that leads to decreased enzyme activity and increased thymidine levels \[40\]. MNGIE is a progressive multi-system disorder with various symptoms, including dysphagia, gastroesophageal reflux disease, gastroparesis, intestinal pseudo-obstruction, muscle weakness, ptosis, and ophthalmoparesis. This condition affects intra-mitochondrial nucleoside levels leading to mtDNA depletion and secondary mtDNA mutations \[41\]. Mutations in the POLG and RRM2B genes have also been described in MNGIE-type phenotypes \[42,43\].

Another PMD is Alpers disease, which is a childhood progressive neurological disorder that affects 1 in 100,000 individuals. The clinical characteristics of Alpers disease include recurrent seizures, mental loss, movement disabilities, and liver disease with clinical symptoms of spasticity, seizures, and dementia. POLG gene mutations occur in 13% of Alpers cases, resulting in reduced DNA replication, mtDNA depletion, and subsequently reduced ATP production \[44\]. mtDNA depletion syndrome (MDDS) is an autosomal recessive disease that is characterized by severe depletion of mtDNA in tissues. MDDS disease is clinically heterogeneous, in which affected tissues can be in a single organ or multiple organs, including liver, brain, kidney, and muscles. The known cause of MDDS is due to mutations in nuclear genes (nDNA) that affect mtDNA maintenance \[Table 2\]. The known genes are those that encode thymidine kinase 2 (TK2), guanosine diphosphate (GDP)-forming succinyl CoA ligase alpha subunit (SUCLG1), adenosine diphosphate (ADP)-forming succinyl CoA ligase beta subunit (SUCLA2), RRM2B, TYMP and deoxyguanosine kinase (DGUOK). All of these genes are responsible for maintaining the mitochondrial dNTP pools during mtDNA synthesis \[45\]. In some cases, mutations in POLG and Twinkle (TWNK) that are involved in mtDNA replication will also result in insufficient mtDNA synthesis \[45,46\]. Similar to MDDS, ataxia neuropathy spectrum (ANS) is also due to mutations in nuclear genes (nDNA), namely POLG and TWNK genes. Patients with ANS typically have symptoms involving nerve, brain, and muscle dysfunctions. Currently, patients with mitochondrial recessive ataxia syndrome (MIRAS) and sensory ataxia neuropathy dystarthis and ophthalmoplegia (SANDO) are considered having ANS. Mutations in POLG are often inherited via an autosomal recessive pattern, whereas mutations in TWNK are often inherited in an autosomal dominant pattern \[47\].

**Secondary mitochondrial diseases**

Secondary mitochondrial diseases (SMD) constitute a group of complex disease manifestations that can be caused by genetic as well as environmental factors. From a genetic point of view, SMD could be due to pathogenic mutations in other genes that are not related to OXPHOS components, but these mutations disrupt mitochondrial function. Also, adverse environmental effects or other factors such as oxidative stress, aging, drug mitotoxicity, and inflammation are some of the processes that could alter mitochondrial functions. If PMD is inherited, then SMD could be inherited and acquired. Typically, SMD occur after conception, which could result in dysfunction in mitochondrial ATP and also non-ATP (mitochondrial fission and fusion) producing capabilities. Mitochondria are highly dynamic organelles and undergo mitochondrial fission and fusion processes regularly \[119\]. Fission is a process of making short, rod- or sphere-shaped mitochondria controlled by DRP1, which is a large GTPase of the dynamin superfamily \[119\]. In contrast, fusion leads to the formation of long and filamentous mitochondria, involving MFN1, MFN2, and OPA1 \[119\]. In normal conditions, mitochondrial dynamics are well-maintained; however, oxidative stress (intra- and extracellular stresses) could disrupt this balance, generating fragmented mitochondria.
Mitochondrial dynamics maintenance is important to preserve mitochondrial shapes, as well as its functions, inheritance, quality control, and cellular apoptosis. Inefficient mitochondrial dynamics could lead to multifactorial diseases, including diabetes, cancer, and kidney and neurodegenerative diseases. Distinguishing whether mitochondrial dysfunction is inherited or acquired is extremely challenging and still poorly understood. Also, PMD and SMD can have overlapping phenotypes or symptoms; moreover, some patients may not have all the components of mitochondrial disease criteria (MDC). Currently, MDC is used to differentiate between PMD and SMD. MDC uses several criteria, including energy production, biochemical, clinical, tissue, and molecular characterizations. However, there are no universal guidelines to diagnose mitochondrial diseases worldwide. Advances in NGS may assist in the diagnosis of PMD or SMD accurately. Comprehensive molecular profiling could determine which genes and pathways are related to PMD and SMD, thereby improving the diagnosis of PMD or SMD.

Mitochondria and diabetes

Energy production through OXPHOS process in the mitochondria may also lead to oxidative stress conditions by increasing ROS formation in the cells. ROS may activate pro-inflammatory pathways, reduce nitric oxide bioavailability, and could further induce diabetic endothelial dysfunction, subsequently leading to the development of diabetes and diabetic kidney disease (DKD). Diabetes is an endocrine disorder due to multiple factors, including genetics, impaired insulin action, obesity, inflammation, impaired autophagy, increased oxidative stress, and mitochondrial dysfunction. The relationship of diabetes with oxidative stress and mitochondrial dysfunction can partially be explained by the damage-associated molecular pattern (DAMPs) that could initiate inflammatory response via various pathways such as T signaling pathways by interacting with (1) Toll-like receptors (TLRs), (2) nucleotide-binding oligomerization domain (NOD)-like receptor family pyrin domain containing 3 (NLRP3) inflammasome, and (3) cytosolic cyclic GMP-AMP synthase (cGAS)-stimulator of interferon genes (STING) DNA-sensing system. Interestingly, Li and colleagues have identified that m.15897G>A mutation of tRNA\text{Thr} which belongs to the haplogroup D4b1, is present in Type 2 diabetes Chinese patients, and this mutation was maternally inherited. The functional study of this mutation showed that a decreased efficiency of mitochondrial tRNA\text{Thr} leads to reduced efficiency of OXPHOS protein synthesis and assembly and ATP synthesis, and decreased mitochondrial membrane potential.

Insulin resistance (IR) is one of the main risk factors for type 2 diabetes, and mitochondrial dysfunction is related to IR development. Recent in vivo and ex vivo metabolic studies involving humans and rodents showed that mitochondrial dysfunction could lead to ectopic lipid deposition and IR. Pereira and colleagues showed that mtDNA could activate the NLRP3 inflammasome, which subsequently causes endothelial dysfunction and inflammation in diabetes. Diabetes reduces endothelium-dependent vasodilation and escalates vascular ROS generation and caspase-1 and IL-1β activation in streptozotocin (STZ)-induced diabetic C57BL/6 mice, but not in those Nlrp3−/−. Deficiency in NLRP3 could prevent diabetes-associated vascular inflammatory damage and endothelial dysfunction. Another example of how mitochondrial dysfunction is associated with diabetes is via the action of the anti-diabetic drug metformin (MF). The protective effect of MF on regulatory networks and integrated stress responses was observed in the brain tissue of STZ-induced diabetic mice. STZ-induced diabetic mice treated with MF (20 mg/kg) showed a significant decrease in protein carbonylation and oxidation. MF treatment also improved mitochondrial function via the increase of the chaperone proteins (HSP60, HSP70, and LonP1). However, the exact mechanisms of how mtDNA causes diabetes are still not fully understood.

Mitochondria and cardiovascular diseases

The role of mtDNA mutations in cardiovascular diseases (CVD) has been discussed extensively. The MtDNA control region is important for controlling mtDNA gene expression. Umbria and colleagues studied mutations in the mtDNA control region in 154 stroke cases and 211 myocardial infarction (MI) patients.
They found that the m.16145 G>A and m.16311 T>C variants could be risk factors for stroke (conditional logistic regression, $P = 0.038$ & $P = 0.018$, respectively), and that the m.72 T>C and m.73 A>G variants may be protective against MI (conditional logistic regression, $P = 0.001$ & $P = 0.009$, respectively)\[129\]. ROS molecules are very harmful and may damage macromolecules, such as proteins, lipids and DNA\[130\]. One such mechanism for ROS effects is the accumulation of damage-associated molecular patterns (DAMPs), which could activate pathogen recognition receptors (PRRs), triggering chronic inflammation-induced oxidative damage\[129\]. Several pathways associated with DAMPs’ action are the mitochondrial calcium handling ROS production, TLRs, NLRP3 inflammasome, cytosolic cyclic GMP-AMP synthase (cGAS)-stimulator of interferon genes (STING) DNA-sensing system, and nuclear factor kappa B (NF-kB)\[129\].

Another mechanism of how mitochondria could be associated with CVD is via circadian clock regulation\[126\]. Mitochondria are among the organelles that are important in controlling the crosstalk between the circadian clock and metabolic pathways, the intestinal microbiota, and the immune system as well. Mitochondria regulate circadian rhythmicity through NAD+ production, SIRT1/SIRT3 activation, and mitochondrial dynamics\[135\]. Yang and colleagues suggested that interference of the Clock gene could suppress mitochondrial apoptosis pathways by stabilizing mitochondrial membrane potential (MMP) and inhibiting mitochondrial membrane permeabilization. These could be due to reduced BAD and BIM proteins that are essential for apoptosis, as well as lower expression of mitochondrial apoptosis factors, i.e., AIF, CYCS, APAF-1, and SMAC, which suppress the formation of the apoptosome and DNA degradation\[132\]. The circadian clock plays a vital role in transcriptional-translational processes in cellular metabolism and mitochondrial activity\[135\]. Perturbations to circadian rhythm could lead to CVD\[134-137\], for example, the circadian clock genes such as BMAL1 can affect vascular proliferation\[136\], and CLOCK and ARNTL may be related to MI\[136\].

Zhang and colleagues also showed that the circadian rhythm and clock genes are related to acute coronary syndrome (ACS), in which plaque stability was negatively correlated with the expression levels of clock genes. The levels of MMP2 and MMP9 were increased in ST-segment elevation myocardial infarction, non-ST segment elevation myocardial infarction and unstable angina pectoris (UA) compared to the control group ($P < 0.05$)\[140\]. Also, Wang and colleagues showed that increased mtDNA 8-OHdG could increase the odds of having coronary artery disease (CAD) (OR = 1.38), coronary stenosis (OR = 1.29), and higher levels of C-reactive protein\[141\]. Although these findings supported the notion that mitochondrial dysfunction could contribute to the development of CVD, the exact mechanism for that association is still unknown.

**Mitochondria and cancer**

One of the hallmarks of cancer cell development is the metabolic changes known as the “Warburg effect” that shows the role of mitochondria in cancer\[142,143\]. The Warburg effect refers to the situation of cancer cells switching their metabolism and energy production from the oxidative phosphorylation (OXPHOS) to glycolysis with lactic acid production, despite the presence of oxygen (aerobic glycolysis)\[144\]. Although the glycolysis process produces less energy than OXPHOS, the abundance of the glucose influx in the cells can result in more energy production at a faster rate potentially. Initially, the Warburg effect is thought of as a result of the mitochondrial defects that inhibit OXPHOS, eventually causing cancer development\[144\]. However, recent findings showed that many cancer cells have functional mitochondria, in which some exhibit a high level of OXPHOS activity. Whereas some are more glycolytic but still retain their mitochondrial functions\[4\]. Moreover, the discovery of the oncogenes in cancer explains that this switching of metabolism or metabolic re-programming is a complex process and may be due to the activation of oncogenic genes.

Activation of the oncogenic driver mutations in KRAS, PI3K, AKT, mTOR, and MYC, as well as the loss of tumor suppressor expression such as p53, facilitate metabolic switching\[145\]. One of the most known altered pathways is the PI3K/AKT pathway, which can increase glucose uptake and glycolysis in the cells\[146\]. Higher glycolysis leads to more production of pyruvates, which are often converted to acetyl-CoA for ATP production and synthesis of other macromolecules such as lipids and amino acids. Mitochondria serve
as the only source of the acetyl-CoA conversion from pyruvates via its pyruvate dehydrogenase (PDH) complex during the TCA cycle\[^{147}\]. To transport these acetyl-CoAs to cytoplasm requires an additional step of converting acetyl-CoA to citrate by mitochondrial-specific citrate synthase. Transported citrates are then converted back to acetyl-CoA via ATP-citrate lyase (ACL) as the resources for lipid and protein synthesis\[^{147}\]. Activated AKT is required for ACL to work, in which AKT phosphorylates ACL for its activation, thus producing more acetyl-CoA in the cytoplasm to fulfill the higher needs of cancer cells\[^{148,149}\]. This AKT/ACL interaction is supported by the fact that inhibition of ACL enzyme causes a reduction in cell proliferation and tumorigenesis, despite increased glucose uptake\[^{148,149}\]. Thus, these findings imply that metabolic programming via mitochondrial acetyl-CoA and citrate is the main oncogenic action of PI3K/AKT in cancer cells.

Another known oncogenic action affecting mitochondrial function is the relationship between the cell growth regulators mTOR and hypoxia-inducible factor 1 (HIF-1). mTOR positively influences HIF-1 action during hypoxia\[^{150}\], and HIF-1 is known to increase glycolytic metabolism\[^{155}\]. Another effect of HIF-1 is the activation of pyruvate dehydrogenase kinase 1 (PDK1) expression, which inhibits PDH activity in mitochondria\[^{152-154}\]. This suppression of PDH activity limits the conversion of pyruvates to acetyl-CoA and shifts the conversion of pyruvates to lactate\[^{152-154}\]. Similarly, MYC oncogenic action is through the mitochondria, in which MYC promotes mitochondrial glutamine metabolism by increasing the expression of glutaminase (GLS), an enzyme that deamidates glutamine to glutamate. Glutamate is needed for nucleic acid and amino acid synthesis, which are vital for cancer cell proliferation. Supporting these findings, cancer cells that express oncogenic MYC, cause growth suppression\[^{155}\] and prevent the Rho GTPase-induced cancer cell transformation and proliferation\[^{156}\].

Loss of tumor suppressor expression, p53 for example, can also contribute to mitochondrial energy switching. In healthy cells, p53 suppresses the expression of glucose transporters (GLUT1 and GLUT4) as a mechanism to control glucose uptake by cells\[^{157,158}\]. p53 protein also suppresses the expression of lactate transporter, monocarboxylic acid transporter 1 (MCT1), to inhibit cellular lactate export, and thus controlling the tumor microenvironment\[^{159}\]. p53 also controls the rate of glycolysis by activating the expression of TP53-induced glycolysis and apoptosis regulator enzyme (TIGAR)\[^{160}\] and reducing the expression of the glycolytic enzyme phosphoglycerate mutase (PGM)\[^{181}\]. Loss of p53 expression causes a reduction of OXPHOS activity with evidence of low mitochondrial complex IV activity\[^{182}\]. In another study, reduced p53 expression decreased mitochondrial mass and mtDNA copy numbers\[^{163,164}\]. Importantly, p53 protein is also responsible for inhibiting the oncogenic PI3K/AKT and mTOR pathways\[^{164}\], thus supporting the notion that the loss of p53 expression can initiate metabolic switching via mitochondrial dysfunction.

**Mitochondria and neurodegenerative diseases**

The most common clinical manifestations of mitochondrial diseases are neurological and neuromuscular syndromes\[^{165}\]. There are two theories that can best explain the role of mitochondria in neurodegenerative diseases. First, a decrease in energy production leads to neuronal depolarization that activates the excitatory amino acid receptors and impairs intracellular Ca\(^{2+}\) homeostasis. This situation is followed by protease activation and cell death, which finally leads to neurodegenerative diseases\[^{166}\]. Second is that mitochondria are the source of ROS via the OXPHOS process, particularly from complex I and III of the ETC. Leakage of electrons from complex I and III produce mitochondrial superoxide, which could cause further damage to macromolecules such as proteins, lipids, and DNA, subsequently leading to a reduced ability of mitochondria to perform their functions. ROS could also activate the apoptosis process via the mitochondrial apoptotic pathway by releasing cytochrome C (Cyto-C) from mitochondria to the cytosol. Pro-apoptotic signals such as Bcl-2 family proteins (Bax and Bak) are translocated into the mitochondria leading to mitochondrial transmembrane permeabilization (MMP)\[^{167}\]. Active Bax and Bak are inserted into the outer mitochondrial membrane (OMM), resulting in increased MMP\[^{169}\]. Subsequently, molecules such as Cyto-C, AIF, Smac/
DIABLO (second mitochondria-derived activator of caspase/direct inhibitor of apoptosis protein-binding protein with low pI) and Omi/HtrA2 are released into the cytoplasm. The formation of the caspase-activating complex or apoptosome composed of Cyto-C, Apaf-1, dATP and procaspase-9, thereby induce the activation of the effector caspases required for apoptosis.

Oxidative stress and mitochondrial dysfunction are associated with several neurodegenerative diseases such as Alzheimer’s, Parkinson’s, Friedreich ataxia, and amyotrophic lateral sclerosis (ALS). The first mtDNA disease was identified in 1988 in a patient with LHON, which is caused by mutations in genes encoding OXPHOS subunit I, III, IV, and V. MTND4*LHON11778 is the most common cause of LHON, representing about 40%-60% of all LHON cases. SOD1 gene mutations cause ALS with increased cellular oxidative stress. SOD1 gene encodes SOD1 protein, which acts as an antioxidant defense mechanism for ROS detoxification. Mitochondrial dysfunction such as defective OXPHOS complexes, changes in mitochondrial membrane potential (MMP), high level of oxidative stress, and decreased mitophagy have been associated with idiopathic Parkinson’s disease (PD). Antony and colleagues showed that mitochondria of idiopathic PD undergo morphological changes and increased resistance to depolarization. The basal mitochondrial membrane potential (Ψm) of a skin biopsy from an idiopathic PD patient was higher compared to healthy control. The mitochondrial morphology parameters such as node degree, mean volume, skeleton size, perimeter, form factor, node count, erosion body count, endpoints, and mitochondria count were reduced in idiopathic PD compared to control. Previously, research on the pathogenesis of PD has been focusing on oxidative stress, mitochondrial bioenergetics defects, and apoptosis mechanisms. Currently, it has been postulated that the crosstalk dysregulation between mitochondria and endoplasmic reticulum, as well as lysosome, may lead to mitochondrial dysfunction, including a bioenergetics defect, abnormal protein aggregation, and finally neuronal cell death.

MULTI-OMICS TECHNOLOGIES TO DIAGNOSE MITOCHONDRIAL DISEASE

The main problems in diagnosing mitochondrial disease are the heterogeneity of the individuals in their clinical presentation and the incomplete understanding of the disease pathophysiology. Advances in molecular technologies, particularly the high-throughput omics that can characterize and identify many targets in a single run, have enabled a more in-depth and accurate diagnosis of mitochondrial diseases and their causes. This section intends to summarize the roles of these omic technologies in mitochondrial disease diagnosis.

Genomics

Genomics research for mitochondrial diseases started about 30 years ago when a report of the small sequence of the mitochondrial genome was published. Afterwards, various findings of the novel mtDNA mutations associated with diseases were reported, including the early discoveries of mtDNA mutations in ragged-red fiber myopathies and LHON. During this time, it was clear that the mtDNA mutations were not solely the main culprit, but that other nuclear DNA mutations may play a role in the disease, as in some patients, the lack of mtDNA mutations were observed. This situation was evident in OXPHOS deficiency when a report of the succinate dehydrogenase-complex flavoprotein subunit A (SDHA) mutations in Leigh syndrome (a disease caused by the deficiency of oxidative phosphorylation complex proteins) was published, in which the SDHA gene mutations resulted in the deficiency of complex II protein in the patient. Thus, identification of mtDNA mutations and any related gene mutations is important for unraveling the disease-causing or underlying cause of the mitochondrial disease.

In the early days, there was a limitation in detecting the mutations, as most of the disease-causing mutations were identified using the single candidate gene sequencing techniques, and these methods were time- and resource-consuming. The most popular methods were Sanger sequencing and Affymetrix’s DNA re-sequencing MitoChip. To address the issue with heteroplasmy, several other methods were also...
developed, including PCR-RFLP analysis\textsuperscript{[50]}, allele-specific oligonucleotide dot-blot analysis\textsuperscript{[185]}, real-time amplification refractory mutation system quantitative PCR\textsuperscript{[186]}, and pyrosequencing technique\textsuperscript{[187]}. However, these applications could only detect a single or a few candidate mutations.

The emergence of NGS technologies has rapidly reduced the cost and time spent with a substantial improvement in the detection ability that allows for a wide-scale detection of genome changes\textsuperscript{[180,184]}. The introduction of whole-exome sequencing (WES) and whole-genome sequencing (WGS) technologies increased the mutational detection rate in mitochondrial disease diagnosis. This effect was evidenced by the percentage of disease-causing mutations identified during the pre-NGS era (10%-20%) compared to after NGS era (30%-50% in some cohorts)\textsuperscript{[188-190]}. Typically, there are two workflows for the detection of mtDNA mutations\textsuperscript{[184,192]}: (1) direct analysis (detection of mtDNA sequence from samples that are enriched with mtDNA apart from the cellular DNA; and (2) indirect analysis (the mtDNA mutations are obtained as by-products of the high-throughput sequencing reads).

Direct detection of mtDNA is usually done by adding a technique to purify or isolate the mitochondria before the NGS workflow\textsuperscript{[184,191]}, such as ultracentrifugation (a density gradients isolation) or the biochemical or mechanical isolation of the organelles. Another approach is to use specific probes or primers to isolate mtDNA, such as in microarray hybridization and PCR-based enrichment methods\textsuperscript{[184,192,193]}. However, it is important to note that using a primer-based method often results in large overlapping regions, and these regions must be removed before the variant calling analysis. The main advantage of this direct method is the elimination of the DNA regions homologous with mtDNA sequences or those known as nuclear mitochondrial DNAs (NUMTs), which exist in various sizes as clones of genuine mtDNA and can be specific to some populations\textsuperscript{[194]}. Therefore, the findings from the direct mtDNA analysis are usually more reliable.

A typical workflow for the indirect mtDNA analysis is through the by-product annotation of the sequencing reads from the WES and WGS. From these WES or WGS sequencing reads, the annotation process also includes a step to map the reads to the mitochondrial genome. Since the average coverage of this mtDNA fraction sequence is higher than the normally targeted gene regions due to the high copy number of mtDNA per cells, the mapping results in good quality data\textsuperscript{[195]}. Because of this high-quality data and the cost-effectiveness of the NGS technologies, the indirect mtDNA analysis has become a favorite tool for mitochondrial disease diagnosis, due to a simple workflow. However, this indirect mtDNA technique has one problem with false-positive results due to NUMTs\textsuperscript{[184,191]}. The inability to confirm whether the mtDNA reads from the WES or WGS sequences are from the nuclear or mitochondrial genome can cause ambiguity of the findings. The simple method to eliminate this issue is to align the raw reads first to the mitochondrial genome and filter the non-aligned sequences, though some NUMTs do exist in the mtDNA genome database; thus, false heteroplasmy can be introduced\textsuperscript{[194,191]}. New software such as MitoSeek can help to address this issue, where this program can extract the mtDNA mutation and heteroplasmmy information from WES data\textsuperscript{[196]}. Furthermore, the existence of the databases such as MSeqDR\textsuperscript{[197]}, MITOMAP\textsuperscript{[198]}, HmtVar\textsuperscript{[199]}, HmtDB 2016\textsuperscript{[200]}, Leigh Map\textsuperscript{[201]} and others provide the comprehensive mutation-phenotype relationships to allow the interpretation of the WES and WGS analyses and thereby unravel any novel mutations in the patients. The mitochondrial disease-specific detection kits or panels are already on the market to improve diagnosis\textsuperscript{[202-205]}. By combining the databases and NGS technologies, there is a continuous discovery of many mutations responsible for various mitochondrial diseases.

**TRANSCRIPTOMICS**

Although genomic NGS techniques are powerful enough to diagnose mitochondrial diseases, the rate of detection for disease-causing mutations are only 25%-50% of cases\textsuperscript{[180,184,192]}. To improve this detection rate, an approach to employ the whole transcriptome sequencing technologies such as the RNA-seq by prioritizing the candidate genes (i.e., those genes that are involved in the oxidative phosphorylation pathways, etc.) to
complement the DNA sequencing, can unravel hidden or deep intronic mutations which usually are missed from interpretation of WES and WGS data. An example of this RNA-seq approach was in the primary muscle samples of the genetic myopathy patients, in which the RNA-seq was able to identify disease-causing mutations in 21% of cases. It is important to note that the DNA sequencing technique was unable to detect the mutations in these patients. However, the challenges of using RNA-seq are attributed to the transcriptomic profiling issues, such as the batch effects, and the requirement of robust filtering pipelines to confirm the results. Nevertheless, the fact remains that the RNA-seq technique can detect mutations in the patients who are not detected from WES and WGS sequencing.

PROTEOMICS

mtDNA encodes 13 proteins, including the mitochondrial respiratory chain proteins, ribosomal RNAs and transfer RNAs. The remaining mitochondrial proteins, which include the TCA cycle components, β-oxidation, protein transports, and the other respiratory chain subunits, are from nuclear DNA. Therefore, to characterize the proteome profile of mitochondrial diseases can be very challenging. Up until now, the number of the mammalian mitochondrial proteins discovered is about 1,100 to 1,900, based on the classifications in each database. One of the earliest databases for the mitochondrial proteome is MITOP, which was released in 1999 and followed by the first comprehensive human mitochondrial proteome database, the MitoProteome Project. Currently, MitoProteome contains about 1,705 genes and 3,625 proteins that are associated with mitochondria. After that, various databases with their analysis tools have been released, including the MitoP2, MitoMiner, and MitoCarta databases.

An example of the mitochondrial dysfunction study using the proteome analysis is the identification of C17orf89 (NDUFAF8) mutation in Leigh syndrome, in which mass spectrometry (MS) crosslinking interactome analysis was able to show C17orf89/NDUFAF8 as a new candidate for the unresolved cases of isolated complex I deficiency. Another study of proteome profiling of the mitochondrial ribosomes revealed that in the small ribosomal subunit, MRPS34 mutations were responsible for the destabilization of the subunit and impaired monosome assembly in the fibroblasts of Leigh syndrome patients. Importantly, the findings were after WES sequencing in those patients, indicating that proteome profiling could also complement WES sequencing to improve the diagnostic detection of mitochondrial disease.

METABOLICOMICS

Due to limited publications, the potential of metabolomics tools to diagnose mitochondrial disease is uncertain. Lactate and pyruvate have been used as biomarkers for mitochondrial dysfunction, though these biomarkers have low sensitivity and specificity. One example is that the lactate stress test was used in the diagnosis of mitochondrial myopathy. However, the sensitivity of the lactate stress test was 69%, but it can complement the other clinical tests to confirm the diagnosis. Advances in technologies allows for the application of mass spectrometry-based metabolomics to profile thousands of small metabolites. In a study of the specific subgroup of the Leigh syndrome patients with mutations in the LRPPRC gene, analysis of the blood and urine metabolites revealed that there were 45 distinct metabolites, including ketones, lipids, kynurenine, lactate, and pyruvates. These findings were important in highlighting the role of metabolomics in unraveling the physiology of mitochondrial disease. However, whether these 45 signature metabolites are specific to the subgroup of Leigh syndrome or applicable to all forms of mitochondrial diseases is unknown. Therefore, further works are needed to confirm these findings, especially in a large cohort, to establish the relationship and diagnostic capacity of the metabolomic approach in mitochondrial disease.

FUNCTIONAL GENOMICS

Following the WES or WGS analysis, the presence of the rare variants or variants of unknown clinical significance (VUS) is challenging to interpret for definitive mitochondrial disease diagnosis. Functional
genomics is an area of research of using multiple molecular data to interpret or understanding the impact of a DNA sequence variant on complex biological processes in a cell. Typically, functional genomic utilizes high-throughput data of multiple omics from a single patient or a disease, to evaluate the impact of these genetic variants in transcription and protein translation\(^\text{[220,221]}\). After that, the validation of these variants is achieved by the functional cell- and tissue-based assays, along with animal models to establish the genotype-phenotypes association as evidence for the pathogenicity.

Typically, the first step of validation is via the confirmation of VUS pathogenicity in the mitochondrial OXPHOS system, which can be done via tissue-based assays (e.g., skeletal muscle biopsy) or cell-based assays (e.g., primary fibroblast culture)\(^\text{[222]}\). An example is the study of genetic myopathy patients, in which the RNA-seq approach in the primary muscle samples was used to complement the DNA sequencing technology to improve the identification of disease-causing mutations\(^\text{[207]}\). Importantly, in this study\(^\text{[226]}\), the pathogenicity of these two splice site VUS were confirmed by analyzing the results with the tissue expression database (Genotype-Tissue Expression (GTEx) Consortium\(^\text{[223]}\)). They found that these splice site VUS are observable in the muscles but have very little presence in the cultured dermal fibroblasts\(^\text{[207]}\), thus indicating that a correct sampling tissue type does matter to discover the relevant genetic defects. Since the GTEx database is freely accessible online, for any mitochondrial study that lacked the tissue biopsy samples, a comparative analysis via the proxy tissue sample data could further refine the findings. The confirmation of the protein product expression and changes in the tissue and primary cells, via the biochemical assays and protein assays such as SDS-PAGE or BN-PAGE, are also used to complement the DNA sequencing analysis\(^\text{[222]}\).

Once the pathogenicity is confirmed, most studies performed the additional assays to discover the disease mechanism or molecular effects of the VUS in cell lines or animal models\(^\text{[222]}\). In most cases, the selected genetic variant is introduced into a cell or animal model via the cell-directed mutagenesis or CRISPR-Cas9 (clustered regularly interspaced short palindromic repeats) technology\(^\text{[220,221]}\). In a study of patients with mitochondrial respiratory chain complex deficiencies, the comprehensive analyses of genetic screening and fibroblast biochemical analysis together with functional cell line assays were able to identify multiple three novel causative variants, in which all of them were pathogenic based on functional cell-based investigation assays\(^\text{[189]}\). This comprehensive evidence for the pathogenicity of the genetic variants is important to elucidate the disease mechanism. Thus, most disease-specific genomic databases will have reported evidence of pathogenicity to support the clinical significance of the variants found.

**IMAGING TECHNOLOGIES**

Another strategy is to complement the NGS data with magnetic resonance imaging (MRI) of the brain or muscles to confirm the changes in the proteins and structures\(^\text{[224]}\). Most mitochondrial diseases are heterogeneous in clinical presentation and symptoms, which are often mixed between diseases. Since many of the patients exhibit neurological symptoms, the application of MRI can detect these changes. One example is to use magnetic resonance spectroscopy (MRS) to evaluate brain chemistry for the detection of metabolic and oxidative defects\(^\text{[224]}\). MRS is a non-invasive in vivo brain imaging to detect biochemical metabolites such as N-acetyl aspartate (NAA), lactate, choline, creatine, and myoinositol\(^\text{[224]}\). Increased lactate levels are a common feature in mitochondrial disease patients, and such lactate elevation has been observed in the brain\(^\text{[225-227]}\), and muscle\(^\text{[228]}\) of patients. However, the findings in the brain are more consistent compared to the muscle\(^\text{[224]}\). Furthermore, this lactate elevation was evident in the early stage before any abnormalities or lesion could be detected in the brain of the mitochondrial disease animal model\(^\text{[229]}\), indicating the usefulness of this MRS technique to improve the diagnosis. Interestingly, the phenotypic changes at the brain structures can complement the genetic screening analysis. In Leigh syndrome, the MRI scans of patients with confirmed SURF1 and COX mutations (nuclear mutations) have T2-abnormalities in the brainstem nuclei, whereas the caudate and putamen lesions are seen in patients with mtDNA mutations\(^\text{[230-232]}\). These findings suggest that MRI scans can confirm the genotype-phenotype changes occurring in mitochondrial disease and thereby improve diagnostic yield.
CHALLENGES IN DIAGNOSING MITOCHONDRIAL DISEASES

Even though quite significant publications of disease-causing mutations are available, challenges remain on how to improve the diagnosis of these mitochondrial diseases in clinical settings, as the rate of detection for disease-causing mutations is only 25%-50% of cases\textsuperscript{[180,184,192]}. Most of the diagnosis approaches are using the NGS technologies, in which the first step is to use the WES approach, followed by a muscle biopsy if more confirmation is needed for the pathogenicity\textsuperscript{[180]}. Various reasons can explain the failure to detect mtDNA mutations in some patients, such as the existence of the difficult-to-detect mutations, including the recurrent \textit{de novo} mutations\textsuperscript{[233]}, splice site defects, mutations in deep intronic or repeated sequences, and others\textsuperscript{[180,184,192]}. One way to address such limitation is to use trio sequencing of parents and child to allow for accurate detections of these difficult-to-detect mutations, as used by the Deciphering Developmental Disorders Project\textsuperscript{[234]}, and the Genomics England 100,000 Genomes (100K) Project\textsuperscript{[235]}.

With the problems of heteroplasmic mtDNA mutations, many recommend that sequencing of muscle DNA is needed to complement the WES findings, especially with the low mutant load. Since most patients with mitochondrial diseases are usually carrying a mixture of wild-type and mutated mtDNA (heteroplasmic), their clinical manifestations of the disease also depend on the ratio of the mutated to wild-type mtDNA\textsuperscript{[236]}. Some of these low-frequency heteroplasmy variants can turn into deleterious high heteroplasmy variants\textsuperscript{[237]}, and could thereby further complicate the diagnosis. Integrated analysis of the omics can also help to improve the diagnosis, as multiple omic findings could verify the accuracy of the results. An example is a cohort study of adult mitochondrial disease patients with the mtDNA mutation m.3243 A \(\rightarrow\) G, in which the combined analysis of proteomics and metabolomics of their urine samples showed very distinct alterations in lysosomal proteins, calcium-binding proteins, and antioxidant defenses\textsuperscript{[238]}. Importantly, these changes were evident in the asymptomatic carriers of m.3243A \(\rightarrow\) G\textsuperscript{[238]}, therefore suggesting the plausibility of a new and early screening strategy of this type of mutation in the patients and their families.

Another issue is the presence of the NUMTs that could interfere with WES or WGS data interpretation and analysis\textsuperscript{[194]}. The indirect method using the WES/WGS data to identify the mitochondrial mutations is a favorable approach due to its cost-effectiveness and high reproducibility. However, the presence of the NUMTs gives some ambiguity to the results\textsuperscript{[184,192]}. Thus, some studies have opted for an addition of the mitochondrial isolation step in the workflow before RNA extraction and sequencing steps to eliminate the NUMTs. However, the resources used are enormous and labor-intensive\textsuperscript{[181]}. Similarly, the proteomic approach for the mitochondrial study also faces the challenges of getting pure mitochondrial proteins\textsuperscript{[204]}. To enrich these mitochondria, many methods have been developed, including the mechanical or chemical disruption method, the differential centrifugation method, and recently introduced magnetic device method\textsuperscript{[239-241]}. However, mitochondrial proteins have dynamic ranges; thus, the samples usually undergo fractionation to reduce their complexity before the analysis\textsuperscript{[208]}, which could increase the cost and time spent for each additional procedure. Most studies use sodium dodecyl sulfate (SDS) polyacrylamide gel electrophoresis (PAGE), and gel slicing to separate the proteins, followed by high-performance LC-MS analysis\textsuperscript{[100,239,242-244]}. Despite the vast potential of these proteomic applications to diagnose mitochondrial disease, the problems lie within the diversity and tissue-specific expression of these mitochondrial proteins. Currently, only indirect measurements are available to detect them\textsuperscript{[208]}. Moreover, the lack of methods to differentiate between the mitochondrial and cytoplasmic functions of these proteins\textsuperscript{[208]} also contribute to the problems. In addition, there are also the issues of technical expertise to use the proteome interactome analysis tools, and the expensive cost to run the comprehensive proteome profiling\textsuperscript{[208]}. Therefore, innovative approaches and advancement of the proteomic applications in the future are needed to solve these issues, and hopefully to increase the potential of these proteomic applications in diagnosing mitochondrial diseases.

Another improvement for the diagnosis of mitochondrial disease using the genetic data is to perform periodic reanalysis of WES/WGS data of the patients, using various or newly improved bioinformatic
pipelines, until the phenotypes changes\textsuperscript{[245]}. In disease with heterogenous genetic and phenotypes involved, the diagnostic yield of using the WES or WGS data ranges from 15\% to 50\% depending on the options used during the inclusion, selection, and stringency criteria of the analysis. The results also depend on the status of the knowledge and genetic databases on the time of sampling, as this research area is continually expanding\textsuperscript{[245,246]}. Previous studies showed a significant improvement in the diagnostic yield if the patient's genomic data is re-analyzed again with an improved bioinformatics pipeline\textsuperscript{[247,248]}. Therefore, a periodic re-analysis of the WES or WSG data in one or two years will improve the identification of the disease-causing mutations.

The most important issue is the plausibility and effectiveness of these applications for routine diagnostics of mitochondrial disease. To run all these high-throughput technologies needs high expertise and advanced machine, which are simply not possible for most of the community clinics. Until there is some development on the cutting-edge technologies to deliver the same comprehensive results as the NGS, but with a far less expensive machine, this issue remains a challenge. Another associated problem is the clinical interpretation of the variants or mutations of unknown significant found from the WES or WGS, which can be confusing to the clinicians even with a genetic counseling background. Thus, the existence of the Clinical Genome Resource\textsuperscript{[249]} and ClinVar\textsuperscript{[250]} online databases help in terms of the interpretation of the genomic variants for the clinicians, which are based on the functional characterization of those variants in various disease models. Despite these challenges, NGS tools are already being used to diagnose mitochondrial diseases in many laboratories, as the potential for screening and diagnosis outweigh the downsides of these applications.

**MITOCHONDRIAL DNA GENETIC TESTING AVAILABLE ON THE MARKET**

Several genetic testing approaches are available on the market nowadays. Here, we discuss the availability of genetic testing for mitochondrial diseases. Centogene is one of the rare disease companies that offer testing for 6,500 genes, which is facilitated by highly innovative analytical platforms based on genomics, proteomics, and metabolomics (https://www.centogene.com). Another company, GeneDX, founded in 2000 by two researchers from the National Institute of Health (NIH), USA, was known as a world genomic leader due to their expertise in a rare and ultra-rare genetic disorder analysis. They offer services for the identification of 20,000 genes and hundreds of rare diseases. Also, they offer the chromosomal abnormalities test (https://www.genedx.com). Family Tree DNA is based in Houston, Texas. They provide services for mitochondrial diseases, for example, the autosomal DNA testing, Y DNA testing, and mtDNA testing (http://www.familytreedna.com). Genebase is one of the established direct-to-consumer (DTC) personal genomics company founded in 2005 based in Vancouver, Canada. Genebase received the world’s prestigious DNA laboratory accreditations, such as the American Association of Blood Banks (AABB), College of American Pathology (CAP), Clinical Laboratory Improvement Amendments (CLIA), and ISO17025 for their services (https://www.genebase.com). 23andMe is also a DTC company, and they offer a simple DNA test from saliva, which can be done at home. They provide the test kit for analysis with all the DNA information directly to the customers (https://www.23andme.com). YSEQ company was established in 2013 by Thomas and Astrid Krahn, which offer several platforms for ancestry study and testing, such as Y-SNPs, Y-STRs, and whole-genome testing with an application for rare disease (https://www.yseq.net/). FullGenomes is one of the DTC company founded in 2013 to make NGS technology possible for the public. They offer products such as whole-genome sequencing and interpretation (SNP, STR, and phylogeny) (https://www.fullgenomes.com). Living DNA was founded in 2016 by David and Hannah Nicholson to provide a DTC NGS screening kit (https://livingdna.com). Oxford Ancestors was founded in 2000 by Prof. Bryan Sykes, and they provide the test for mtDNA maternal and yDNA for paternal ancestry (http://www.oxfordancestors.com) [Table 3].

**CONCLUSION**

Mitochondrial diseases are complex and clinically heterogeneous, making early diagnosis difficult. Advances in omics technology, particularly genomics, NGS via targeted mtDNA genome of WES and WGS enable
a comprehensive detection strategy with high sensitivity and specificity. Early identification of affected individuals is important for the patient’s outcome, as early treatment will result in a better prognosis. Importantly, with a simple blood sampling, the cascade screening of the family members can be done with the same strategy.

DECLARATIONS

Authors’ contributions
Wrote the manuscript and revised the manuscript for intellectual content: Sulaiman SA
Wrote a few parts of the manuscript: Mohd Rani Z, Radin FZM
Planned and wrote the manuscript and also revised the manuscript for intellectual content: Abdul Murad NA

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