Review: Central nervous system involvement in mitochondrial disease

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Mitochondrial respiratory chain defects are an important cause of inherited disorders affecting approximately 1 in 5000 people in the UK population. Collectively these disorders are termed ‘mitochondrial diseases’ and they result from either mitochondrial DNA mutations or defects in nuclear DNA. Although they are frequently multisystem disorders, neurological deficits are particularly common, wide-ranging and disabling for patients. This review details the manifold neurological impairments associated with mitochondrial disease, and describes the efforts to understand how they arise and progressively worsen in patients with mitochondrial disease. We describe advances in our understanding of disease pathogenesis through detailed neuropathological studies and how this has spurred the development of cellular and animal models of disease. We underscore the importance of continued clinical, molecular genetic, neuropathological and animal model studies to fully characterize mitochondrial diseases and understand mechanisms of neurodegeneration. These studies are instrumental for the next phase of mitochondrial research that has a particular emphasis on finding novel ways to treat mitochondrial disease to improve patient care and quality of life.

Keywords: mitochondrial disease, mitochondrial DNA, neurodegeneration, respiratory chain defects

Mitochondria

Mitochondria are the main source of energy, in the form of adenosine triphosphate (ATP), in neurons, but they also play important roles in initiating apoptosis, iron-sulphur cluster biogenesis and calcium buffering. Mitochondria house the machinery necessary to perform oxidative phosphorylation (OXPHOS) within the inner mitochondrial membrane. This machinery comprises an electron transport chain (ETC) which is made up of four multisubunit complexes (complex I–IV) and two mobile electron carriers (ubiquinone and cytochrome c) that are thought to form supercomplexes [1]. The ETC allows the transfer of electrons through the complexes and the translocation of protons into the intermembrane space. This proton gradient is harnessed by ATP synthase (complex V) to synthesize ATP.

The OXPHOS system is under dual genetic control of the mitochondrial DNA (mtDNA) and nuclear DNA (nDNA). mtDNA exists as a multiple copy circular double-stranded 16.6 kb DNA molecule and encodes for 13 polypeptide proteins for the ETC and the necessary RNA machinery [2 ribosomal RNAs (rRNAs) and 22 transfer RNAs (tRNAs)] required for their synthesis within the mitochondria [2]. The remaining proteins subunits and assembly factors are encoded by the nDNA and imported into mitochondria. In addition to this, the nDNA encodes for proteins involved in mtDNA
maintenance and expression and mitochondrial dynamics, including mitochondrial fission and fusion.

**Mitochondrial genetics and disease**

Mitochondrial disorders represent a common group of human genetic diseases that result from a primary defect of mitochondrial OXPHOS. Impaired energy generation due to a failure of OXPHOS can therefore result from a genetic defect arising in either the mtDNA or nDNA. MtDNA mutations may either be in the form of point mutations, such as the most common mtDNA mutation m.3243A>G which affects mt-tRNA Leu, or large-scale mtDNA rearrangements, including single large-scale mtDNA deletions associated with Kearns–Sayre syndrome (KSS). MtDNA is strictly inherited through the maternal line [3], and can exist in a state of homoplasmy (comprising solely wild type or mutated mtDNA molecules) or heteroplasmy (mixture of wild type and mutated mtDNA). The heteroplasmy level of mutated mtDNA in cell is important for determining whether there is disruption to OXPHOS that results in a detectable biochemical defect.

The heterogeneous nature of mitochondrial disease means that patients may be affected at any age and with multisystem involvement that does not always correlate with genotype, and so this presents a major diagnostic and treatment challenge for patient management. Certain mtDNA mutations are associated with specific syndromes, however, a single mutation can cause several different phenotypes depending on the segregation of the mutation and heteroplasmy level. Perhaps the best example of this is the m.3243A>G MT-TL1 mutation which was first described in relation to the classic mitochondrial encephalopathy with lactic acidosis and stroke-like episodes (MELAS) syndrome [4]. However, the m.3243A>G mutation can also lead to chronic progressive external ophthalmoplegia (CPEO) and maternally inherited deafness and diabetes. Current estimates of the prevalence of mtDNA mutations contributing to human disease in the north east of England affect 1 in 5000 individuals [5,6].

The emergence of next generation sequencing has led to a rapid expansion in the number of nDNA mutations being identified [7]. Nuclear encoded DNA defects affecting mtDNA maintenance, OXPHOS assembly and structure and mitochondrial dynamics are predicted to affect approximately 2.9 per 100 000 of the adult general population [5]. One of the most common nDNA defects, are mutations in the POLG gene that encodes for the sole mtDNA polymerase γ and is responsible for replication of the mitochondrial genome. POLG mutations result in a range of different clinical phenotypes either with an early onset, such as in Alpers’ syndrome, or late onset as seen in patients with CPEO, myoclonic epilepsy myopathy sensory ataxia (MEMSA) and ataxia neuropathy spectrum (ANS) and these are associated with either accumulation of multiple mtDNA deletions and/or depletion of mtDNA content within individual neurons [8].

**Neurological symptoms and neuropathology**

Despite only weighing 2% of total body mass, the brain consumes 20% of oxygen and 50% glucose supplied through delivery from the vasculature and is used to drive aerobic respiration within mitochondria [9]. Neurons are heavily dependent on mitochondria for the production of energy for a number of processes underpinning neurotransmission including the regulation of the sodium potassium ATPase pump, regulation of intracellular calcium concentration and exocytosis/recycling of synaptic vesicles [10,11]. Glial cells, such as astrocytes, play important roles in glutamate metabolism, and ion and water homeostasis and are dependent on OXPHOS but also contain stores of glycogen for energy metabolism [12]. It is for these reasons, the neuronal cells are especially vulnerable to deficits in energy generation within mitochondria and neurological deficits are the most common symptoms reported in patients with mitochondrial disease and are probably the largest contributor to morbidity and mortality. We discuss the common neurological features and associated neuropathological findings.

**Cerebellar ataxia**

Cerebellar ataxia is common in both adult and paediatric patients with mitochondrial disease [13–15]. A recent UK cohort study revealed that out of 345 adult patients with mitochondrial disease, 225 were affected by cerebellar ataxia that is progressive in nature. Ataxia is reported in patients harbouring many different mtDNA mutations, including the common tRNA point mutations MT-TL1 and MT-TK and also single
large-scale mtDNA deletions. Cerebellar changes are also reported in patients harbouring nDNA mutations, including RARS2 [16], C10orf2 [17] and POLG mutations [18]. In patients harbouring POLG defects, ataxia may be reported as one of the prevailing neurological features and is frequently reported in patients with MEMSA [19,20] and ANS. The main neuroradiological finding is cerebellar volume loss including progressive cerebellar atrophy and to a lesser extent cerebellar hypoplasia [14,21].

**Neuropathology studies** The major neuropathological findings include cerebellar atrophy, microinfarcts, Purkinje cell loss and morphological changes, including axonal torpedoes and abnormal dendritic arborisations, in remaining neurons [13,22–24]. Patients harbouring MT-TL1 show Purkinje cell vulnerability and a high frequency of microinfarcts (or ischaemic-like lesions) affecting the cerebellar cortex. The aetiology of microinfarcts is not well understood, however, it might be the result of energy deficiency within neurons combined with microvascular abnormalities which lead to localized necrotic cell death [25,26]. The pattern of olivo-cerebellar pathway vulnerability in patients harbouring MT-TK and POLG mutations are very similar. Patients with these genetic defects also demonstrate ischaemic-like lesions that histologically resemble those seen in MT-TL1. Surviving neurons show a profound loss of complex I subunits indicative of complex I deficiency [13,17]. It is not understood if neurons that harbour such a severe complex I defect are capable of normal neurotransmission and why they do not undergo cell death. While cell loss and synaptic changes are detected in patients harbouring a single large-scale mtDNA deletion associated with KSS [13,27,28], the major neurohistopathological hallmark is that of a myelinopathy in the deep white matter of the cerebellum, which is widespread throughout the CNS, and this is attributed to specific changes to the oligodendrocyte populations [29].

**Extrapyramidal features**

Extrapyramidal features, including dystonia, akathisia (motor restlessness), Parkinsonism and tardive dyskinesia, have been described in patients harbouring POLG mutations. In 2004, Luoma and colleagues reported the first description of Parkinsonism in a patient with autosomal dominant PEO due to mutated POLG [30]. As then parkinsonism has been shown to co-segregate with the POLG defect [30,31], respond to levo-DOPA treatment [30] and the brains of the patients show reduced dopamine-uptake in the striatum following DaT scan [32–34].

Leigh syndrome is a progressive neurodegenerative condition affecting infants owing to mutations affecting mtDNA, including mt-mRNA and mt-tRNA mutations, and nDNA defects [35,36]. Clinical symptoms include developmental delay, hypotonia, dystonia and failure to thrive. The clinical course may be varied, however, patients often present with respiratory abnormalities, nystagmus and ataxia. Characteristic neuroradiological features include bilateral symmetric lesions affecting the brain stem and basal ganglia which are likely to contribute to the dystonia in these patients.

**Neuropathology studies** In idiopathic Parkinson’s disease (iPD), complex I deficiency and accumulation of mtDNA deletions have been detected in substantia nigra pars compacta (SNpc) neurons [37,38]. In a recent study, mtDNA deletions were also found in SNpc neurons from patients with POLG mutations that were very similar to those seen in iPD [39]. Changes in SNpc neurons from patients harbouring POLG and C10orf2 mutations showed neuronal loss without alpha-synuclein or Lewy bodies, whereas remaining neurons demonstrated a preferential complex I deficiency [40]. Another study of patients with POLG mutations demonstrated severe nigrostriatal degeneration with decreased mtDNA in remaining neurons [41]. In both studies, the degree of nigrostriatal degeneration did not correlate with clinical signs of Parkinsonism.

The neuropathological features of Leigh syndrome include symmetrical vasculo-necrotic lesions affecting the substantia nigra, putamen in the basal ganglia and thalamus or sub-thalamic nuclei. These lesions are characterized by spongiform changes, and cytotoxic oedema and also increased capillary prominence [42].

**Peripheral neuropathies**

Peripheral nerve involvement is frequent in mitochondrial disease, and might define the clinical picture in a number of patients with intergenomic signalling defects causing multiple mtDNA deletions and mtDNA
depletion. This includes ANS due to POLG mutations where the neuropathy is mainly sensory with a loss of proprioception and vibration sense and touch and pin-prick sensory changes with a varying degree of distal muscle weakness. Nerve conduction studies show decreased or absent sensory action potentials and varying motor involvement [43]. In almost all patients with mitochondrial neurogastrointestinal encephalopathy, due to mutations in TYMP, the neuropathy is predominantly demyelinating with slowed conduction velocities and axonal degeneration and segmental demyelination in biopsied nerves [44,45]. Charcot-Marie-Tooth hereditary neuropathy Type 2A (CMT2A) disease is due to mutations in MFN2, which encodes a protein involved in mitochondrial dynamics, and this leads to a progressive early onset axonal neuropathy [46]. Peripheral neuropathy might also be present in other disorders due to mtDNA mutations, including point mutations m.8344A>G and m.3243A>G, in conjunction with other neurological features [47,48].

Neuropathology studies Relatively few studies have explored neuropathological features associated with peripheral neuropathy, however, a recent electrophysiological and neuropathological study provided evidence of a sensory neuronopathy in patients with POLG mutations, and confirmed a loss of sensory neurons and respiratory chain deficiency in remaining cells in the dorsal root ganglia. In this patient, respiratory chain deficiency was associated with lowered mtDNA content in remaining neurons [43]. This may explain the common finding of a loss of fibre tracts from the dorsal column in patients with a peripheral neuropathy [43,49].

Epilepsy

Epilepsy may be a presenting feature, such as in myoclonic epilepsy ragged red fibres (MERRF) or MEMSA, or emerge on the background of multiple neurological impairments, such as in MELAS, and seizures may occur in any seismology or classification. A recent study identified that approximately 23.1% of 182 adult patients with genetically-defined mitochondrial disease in a UK cohort develop epilepsy, and that approximately 34.9% of patients with MT-TL1 mutation will develop epilepsy and of these 17.5% also develop stroke-like episodes. In these patients, epilepsy may be classified as focal motor seizures with or without a loss of consciousness. While in patients harbouring MT-TK, epilepsy affects 92% of patients and is usually progressive myoclonus and generalized tonic clonic seizures [50]. Epilepsy occurring in infancy is associated with a particularly poor prognosis with a 50% fatality rate in children 9 months after a diagnosis of epilepsy being made [51]. Alpers’ syndrome affects neonates and infants who may present with refractory seizures, developmental delay and liver involvement. The epilepsy in these patients are particularly devastating as they may develop status epilepticus, which can be focal or generalized, and from which they may never recover [52]. Management of epilepsy in mitochondrial disease is often challenging.

Neuropathology It is difficult to pinpoint neuropathological changes specifically associated with seizure activity in patients with mitochondrial disease as there are often multiple neurological impairments coinciding with the epilepsy and it is difficult to dissect primary changes vs. secondary changes. Often seizures associated with stroke-like episodes reveal foci of necrosis affecting the cerebral cortex, and these will be discussed in further detail under ‘seizure-associated stroke-like episodes’. The importance of neuronal energy depletion in epilepsy is underscored by the high prevalence of epilepsy in mitochondrial disease. A vulnerability of cortical inhibitory interneurons has recently been shown in post-mortem tissues from patients with mitochondrial disease with evidence of combined complex I and IV defects and reduced interneuron densities (Figure 1). The identification of combined respiratory chain defects in this study utilizes a novel immunofluorescent assay to interrogate mitochondrial respiratory chain subunit abundance in conjunction with a mitochondrial mass marker within specific neuronal populations, which allows a more precise and accurate method of quantification [53]. The combined respiratory chain defect in interneurons is proposed to influence neuronal networks which could contribute to lowered threshold for seizure activity and permit neuronal hyperexcitability but this remains to be substantiated [54]. Mitochondrial abnormalities have been detected in both human tissues and animal models of temporal lobe epilepsy with detection of mitochondrial ultrastructural pathology, impairment of complex I in the CA3 region of the hippocampus [55,56].
Seizure-associated stroke-like episodes

Stroke-like episodes are often reported in patients with mitochondrial disease and are best defined as an episode of focal cerebral metabolic crisis. Important triggers that have been recognized include febrile illness, headaches, seizures and dehydration. Stroke-like episodes are one of the main syndromic features of patients with MELAS due to the MT-TL1 mutation, and in these patients the standard diagnostic criteria includes clinical features, onset usually before aged 40 years, neuroradiological imaging evidence and seizures [57]. However, it not only includes patients with the MT-TL1 mutation that are affected by stroke-like episodes but also includes patients harbouring mutations in POLG [58], FBXL4 [59] and other mtDNA point mutations, including MT-TK mutations [60]. In this review, we refer to stroke-like episodes as seizure-associated stroke-like episodes because seizures may signal/be concomitant to the onset of stroke-like episodes in patients and are often detected on electroencephalogram (EEG) [61].

During the prodromal phase, patients might develop positive visual phenomena including phosphenes and moving objects and often associated with a throbbing headache. As the stroke-like episodes evolve, patients can develop negative visual phenomena including homonymous hemianopia and cortical blindness; deficits that may become fixed with time. Stroke-like episodes are detected using magnetic resonance imaging (MRI) as T2-weighted hyperintensities which are typically in posterior brain regions though not associated with any major vascular territories. Recently there have been a number of radiological studies to determine the mechanisms contributing to the onset of a stroke-like episode and also to monitor its progression as they can migrate to other regions of the brain. There is evidence that during preclinical or acute onset of a stroke-like episode, focal hyperperfusion can be detected by either arterial spin labelling or SPECT and can manifest before any structural abnormalities might be seen on MRI [62–67]. In addition to this, there are reports of vasodilation following magnetic resonance angiography and impaired cerebrovascular reactivity.

Figure 1. Mitochondrial respiratory chain defects in inhibitory interneurons in a patient with mitochondrial disease. Inhibitory interneurons (GAD65-67; blue) from a control subject show good co-localization of a mitochondrial mass marker (porin; magenta) with a subunit of complex I (NDUFB8; red) and complex IV [cytochrome c oxidase I (COXI); green]. While inhibitory interneurons from a patient harbouring autosomal recessive mutations in polymerase gamma show decreased expression of both complexes I and IV (marked by white asterix) relative to the mitochondrial mass marker. Scale bar = 10 μm.
Stroke-like episodes in mitochondrial disease have a similar manifestation as posterior reversible encephalopathy syndrome (PRES) where headache, encephalopathy, seizures and visual disturbances are common. In PRES, patients typically develop cerebral oedema in the occipital poles, presumed due to cerebral autoregulatory failure [75]. Moreover, it has been conjectured that the vulnerability of the posterior circulation to the development of oedema maybe due to the relative reduced in sympathetic innervation of the posterior circulation [75, 76].

The treatment of seizure-associated stroke-like episodes involves the urgent control of seizures in the acute phase and then continued anticonvulsants for the prevention of further seizures and development of refractory status epilepticus, a widely recognized complication of mitochondrial disease [77, 78]. Additional recent studies advise the implementation of L-arginine therapy in the acute phase and during the interictal phases of stroke-like episodes to prevent or decrease the severity of these episodes in patients with MELAS [79]. The proposed mechanism of action of L-arginine is to allow the conversion of L-arginine into nitric oxide (NO) in the vascular endothelium. NO is a potent mediator of vascular smooth muscle relaxation to improve vasodilation and blood flow in the cerebrovasculature. Recent studies have shown that serum arginine and NO levels are depleted in patients with MELAS and that administration of L-arginine may be beneficial [80, 81].

There is a clear need for standardized, serial neuroimaging studies of patients with stroke-like episodes in order to establish specific imaging criteria to accurately detect and monitor stroke-like episodes, especially when considering therapeutic trials for these patients where accurate and measurable outcomes are crucial for determining drug efficacy. Furthermore, there is an urgent need for effective evidence-based therapeutic strategies using randomized open-label, multicentre studies to confirm the efficacy of anticonvulsant drugs and L-arginine in conjunction with serial neuroimaging to provide insights into the evolution of lesions to the brain and influence the implementation of neuroprotective therapies to prevent the irreversible loss of brain function.

Neuropathology The most common neuropathological findings in patients with stroke-like episodes are the presence of multiple foci of neuronal necrosis (recently termed focal energy-dependent neuronal necrosis) affecting the posterior cerebral cortex, basal ganglia and cerebellum. These areas of chronic necrosis are ischaemic-like in appearance; exhibiting neuronal cell loss, destruction of the neuropil and subcortical white matter accompanied by proliferation of astrocytes and inflammatory cells (Figure 2), but they do not conform to major vascular territories. The degree of necrosis varies according to the severity of the stroke-like episodes, in terms of whether they were recurrent, or timing before death. In some patients, laminar cortical necrosis may be evident affecting cellular layers III and IV–V, whereas in others the entire cortical ribbon may be ablated and might reflect a more chronic process.

While the pathological mechanisms underpinning stroke-like episodes are not clearly defined, unaffected areas of the brain provide clues with evidence of neuronal respiratory chain defects and angiopathic changes. There are a number of hypotheses to explain mechanisms underpinning neurodegeneration occurring during and after a stroke-like event, and these include: (i) Mitochondrial cytopathic theory, (ii) Mitochondrial angiopathic theory and finally and (iii) Neuronal hyperexcitability theory. The mitochondrial cytopathic theory suggests that respiratory chain defects and therefore compromised energy production in neurons or glial cells or a combination of both are sufficient to cause focal necrosis [82]. The mitochondrial angiopathic theory is supported by evidence of aggregated, enlarged mitochondria within smooth muscle and endothelium in the vasculature, and biochemical data revealing strongly succinate dehydrogenase-reactive vessels which show an accumulation of mitochondria, similar to ragged red fibres [83, 84]. This might represent a morphological abnormality of the microvasculature which leads to physical occlusion. There are contradictory reports which suggest that these abnormal mitochondria still retain cytochrome c oxidase (COX) activity, whereas others suggest COX activity is lost and is associated with high mutant load in microvessels [25, 85]. In the latter studies, mitochondrial respiratory chain defects affect pial arterioles and smaller penetrating arterioles which might compromise cerebrovascular autoregulation and therefore blood vessel tone. Another proposed...
CNS involvement in mitochondrial disease

Cognitive decline and behavioural disturbance

Cognitive impairment and dementia have been frequently in case reports of patients with mitochondrial disease due to mtDNA mutations, however, there are very few large-scale longitudinal studies and standardised criteria to define cognitive impairments. A neuropsychological study conducted on 16 adult patients with mitochondrial disease (either due to MT-TK point mutation, single large-scale mtDNA deletion or multiple mtDNA deletions) revealed non-verbal cognitive impairments and verbal short-term memory deficits, associated with working memory, with hypometabolism in the temporal lobes on neuroradiological imaging [89]. In a separate study, patients with either a single large-scale deletion or MT-TL1 point mutation identified focal deficits in visual construction, attention and abstraction [90]. In patients harbouring the MT-TL1 mutation, neuropsychological testing provided evidence of a global deterioration in cognition affecting executive function, attention, language, memory, visuospatial and motor function [91]. Cerebral lactic acidosis in patients harbouring MT-TL1 mutations correlates with neuropsychological scores in patients with or without stroke-like episodes implying dysfunction in the frontal domains which is not detected by neuroradiological imaging [92].

Neuropathology

There is a paucity of studies correlating neuropathological changes with cognitive impairments in mitochondrial disease. Although it is likely that deterioration in cognition might be attributed to the presence of necrotic foci which impinge on multiple circuits involved in cognitive processing, certainly cognitive decline has been observed in patients without stroke-like episodes implying that the frontal domains are affected. Neuropathological studies detect changes in neuronal populations in patients affected by either mtDNA or nDNA mutations which could contribute to cognitive dysfunction in mitochondrial disease. A loss of calbindin expression in hippocampal neurons without neuronal loss has been described, and also loss of cortical inhibitory interneurons and respiratory chain deficiencies which could certainly contribute to impaired network activity in the brain [54,93].

Modelling the neurology of mitochondrial disease

Clinical, molecular genetic and neuropathological studies have given important insights into the disease pathogenesis. The application of the knowledge gained from these studies is now driving the development of cell and animal model systems to test hypotheses and

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Figure 2. Focal energy-dependent neuronal necrosis affecting the occipital cortex of a patient harbouring a MT-TK mutation. Focal neuronal necrosis affects the cortical ribbon (A: cresyl fast violet) causing neuronal cell loss and spongiform degeneration (B: cresyl fast violet) accompanied by astrogliosis (C: weak expression of GFAP) and inflammation (D: HLA-DP, DQ, DR). Remaining neurons within the lesion show both complex I expression (E: NDUFB8 – black arrow) and complex I deficiency (E: NDUFB8 – red arrow), whereas mitochondrial density is maintained in all neurons (F: porin – black arrow). Subcortical white matter shows a loss of myelin (G: Loyez) and axonal loss (H: phosphorylated neurofilaments), whereas in unaffected cortical regions myelin (G:ii; Loyez) and axonal density are maintained (H: phosphorylated neurofilaments). Scale bar = 100 μm.
adopt a functional approach to understanding mechanisms and evaluate the efficacy of treatment.

Transgenic mouse models

Modelling mtDNA disease has been impeded by the inability to generate stable mtDNA point mutations in mice without undergoing a bottleneck effect and loss of the mutation through successive generations of mice [94]. Many of the models generated to date have utilized Cre/Lox technology to selectively knockout (KO) nuclear genes involved in: (i) mtDNA maintenance, replication, transcription, translation (ii) expression and assembly of OXPHOS complexes and (iii) mitochondrial dynamics to evaluate the effect on specific populations of neurons. A comprehensive description of these models is beyond the scope of the current review and we refer readers to a three part review miniseries published last year which discusses each model in detail [95–97].

Here, we provide a brief overview and examples of models associated with defects in: (i) mtDNA maintenance and (ii) OXPHOS in certain neuronal populations. A number of mouse models have been generated by targeting TFAM which is a nuclear encoded transcriptional activator which has a number of roles within the mitochondrion including binding to the mtDNA promoter to activate mitochondrial transcription, provides RNA primers to facilitate mtDNA replication and also plays a histone-like role by coating the mtDNA. There are two models which have used neuronal-specific KO of TFAM, including forebrain neurons using a CaMKIIα-Cre [Mitochondrial Late Onset Neurodegeneration (MILON) mice] and dopaminergic neurons using a DAT-Cre (MitoPark mice) which lead mtDNA depletion in the targeted neurons. The MILON mice show a reduction in mtDNA copy number in neocortex at 2 months of age and develop a phenotype at age 5–6 months whereby they severely deteriorate and die within 2–3 weeks after onset [99]. Neuropathologically the neocortex undergoes severe degeneration and loss of organization, whereas in the hippocampus, CA1 is absent with a profound loss of pyramidal cells, CA2 is intact and CA3 is slightly affected. Chimeric MILON mice were generated to study the effects of mosaic pattern of wild type and mtDNA depletion in forebrain neurons. They revealed that low levels (20%) of respiratory chain deficiency were sufficient to produce behavioural abnormalities, whereas higher levels (80%) of respiratory chain deficiency were sufficient to induce trans-neuronal degeneration events [100].

The MitoPark mice harbour dopaminergic neuron-specific KO of TFAM and develop a progressive Parkinson-like phenotype at 12 weeks of age with motor deficits, dopamine depletion and SNpc degeneration [101]. In asymptomatic mice aged 6–8 weeks, electrophysiological experiments reveal subtle changes in nigrostriatal function with an absence of pacemaker activity in SNpc neurons and impaired dopamine release from their axonal terminals [102].

The second group of transgenic mouse models we discuss are those associated with defects in OXPHOS either due to abolished complex subunit or assembly unit expression. There are a number of model which target subunits associated with complex I as complex I deficiencies are frequently detected in mitochondrial disease. A neuronal and glial-specific KO (using a nestin-Cre) of NDUFS4 (NADH:ubiquinone oxido-reductase subunit S4), which encodes for a nuclear-encoded accessory subunit of complex I, has been generated.

Figure 3. A working hypothesis to dissect out mechanisms underpinning formation of focal neuronal necrotic lesions during stroke-like episodes. Mitochondrial DNA defects are segregated throughout individual mitochondria and clonally expand to high mutant loads to cause mitochondrial respiratory chain defects defined by a loss of respiratory chain complex activity. Respiratory chain deficiency compromises mitochondrial function leading to reduced adenosine triphosphate (ATP) generation, and impaired calcium buffering. Respiratory chain deficiencies have been documented in multiple cell types within the CNS with neurons, glia and blood vessels (smooth muscle and endothelial cells) and while the impact on cellular function is not yet fully elucidated, the potential mechanisms are described here. Respiratory chain deficient inhibitory interneurons could contribute to modified neuronal and synaptic function leading to neuronal network instability and neuronal hyperexcitability which could in turn lower the threshold for seizure generation. Seizures are commonly reported in the early phase of a stroke-like episodes and can be detected on electroencephalogram. Respiratory chain deficiencies affecting glial cells, including astrocytes (deficient – red arrow, intact – black arrow), are also observed which could affect synaptic transmission (e.g. glutamate metabolism, tonic homeostasis) and contribute to cortical spreading depression. While respiratory chain deficiencies affecting the CNS vasculature might leading to cerebral auto regulatory failure leading to a mismatch between neuronal activity and appropriate blood flow. This further worsens the ongoing neuronal insult leading to cytotoxic and vascular oedema, hypertensions on magnetic resonance imaging and irreversible cell loss in necrotic lesions.

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This mouse shows rapidly progressive gait impairments, breathing difficulties and death by 7 weeks of age and therefore resembles Leigh syndrome. Neuropathological features include complex I deficiency and progressive degeneration in the form of spongiosis and lesions within the olfactory bulb, cerebellum and vestibular nuclei accompanied by progressive glial activation and inflammation [103,104]. A recent study has investigated hypoxia as a possible treatment strategy for ameliorating disease progression: NDUFS4 KO mice were exposed to chronic hypoxia (11% oxygen) environment showed extended lifespan with improved motor coordination and reduced neuroinflammation upon neurohistopathological assessment [105].

Mice harbouring NDUFA5 (NADH:ubiquinone oxidoreductase subunit A5) KO, a nuclear-encoded structural subunit of complex I, in forebrain neurons (using CaMKIIα-Cre) were healthy until 10–11 months of age when they became lethargic, demonstrated a loss of motor coordination and hindlimb clasping. These mice showed only a mild encephalopathy with evidence of complex I deficient neurons without evidence of neurodegeneration which the authors suggest might be related to compensatory processes [106].

Recently, forebrain neuron-specific (using CaMKIIα-Cre) KOs for RISP (encoding for a catalytic subunit of complex III; Reiske iron-sulphur cluster protein) and COX10 (encoding an accessory protein involved in assembly of COX) have been created to understand more about the selective brain involvement in mitochondrial disease. These studies revealed that the RISP-KO mice developed a rapid phenotype aged 2 months with poor rotarod performance with a rapid disease progression and died aged 3–3.5 months. Evaluation of CNS tissues confirmed complex III deficiency and high levels of oxidative stress in remaining neurons, particularly affecting the piriform cortex. While in the COX10-KO mice, rotarod performance was poor aged 3 months but they survived until aged 8–12 months with a vulnerability of the cingulate cortex and oxidative stress in remaining neurons [107].

**Induced pluripotent stem cells**

Induced pluripotent stem cells (iPSC) hold the potential for in vitro disease modelling, drug screening and cell-replacement therapies. iPSCs constitute cells which are capable of self-renewal and can differentiate into any cell type and are generated by reprogramming patient fibroblasts with pluripotency associated transcription factors. In recent years, a number of studies have utilized patient-derived iPSC to understand mechanisms of mitochondrial disease exploiting the main advantages of (i) maintaining the nuclear genetic background of the patient, and (ii) differentiation into disease-specific tissue- or cell-type, e.g. postmitotic neurons. There are some caveats associated with the study of iPSCs including the acquisition of genome instability during culture [108], the alteration of mitochondrial morphology with rounded appearance and poor cristae structure, low mtDNA copy number and respiratory capacity to facilitate the metabolic switch to glycolysis. In addition, impaired mtDNA integrity may be a by-product of cellular reprogramming [109]. There have been a number of studies utilizing iPSCs to understand pathogenetic mechanisms in mitochondrial disease, and these are discussed in more detail in a recent review by Hatakeyama and colleagues [110].

Induced pluripotent stem cells have been derived from fibroblasts from patients with MELAS due to the m.3243A>G mutation and two cell lines were generated: one harbouring high heteroplasmic levels (>80%) and the other which was low heteroplasmic levels (undetectable). Following differentiation into neurons, the high heteroplasmasy line showed downregulation of complex I and an increase in mitophagy [111]. Another study that investigated fibroblasts from a patient with MELAS created iPSC harbouring the m.13513G>A mutation at heteroplasmic levels (50–60%) and a mutation-free cell line. Heteroplasmic cell lines showed a decline in heteroplasmic levels with long-term maintenance in culture [112]. In a recent study, cellular homoplasy for the m.8993T>A mutation associated with Leigh syndrome was genetically corrected using somatic cell nuclear transfer [113].

**Future directions/studies**

Unlike other neurodegenerative diseases, brains from patients with mitochondrial disease show atrophy and severe neuronal cell loss that this is not associated with any extra- or intracellular accumulation of misfolded proteins. The salient neuropathological features are the decrease or absence of mitochondrial respiratory chain subunits, particularly involving subunits comprising complex I, and high ratio mutated mtDNA to wild type
mtDNA in surviving neurons. The future of mitochondrial research should aim to prevent the irreversible brain loss and neurological disability in these patients. Post-mortem human brain studies are invaluable for characterizing the chronic neurodegenerative changes, and new techniques, such as CLARITY [114,115], will dramatically improve our understanding of disease pathogenesis. However, our neuropathological studies can be strengthened by linking to patient iPSCs and transgenic mouse models to interrogate functional systems and enhance our understanding of progressive mechanisms of degeneration and facilitate the development of better-directed therapeutic intervention.

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