Hypothesis

Mitochondrial genome: defects, disease, and evolution

Angus Clarke

Abstract
Defects of mitochondrial function are often caused by defects of the mitochondrial genome. The hypothesis that defective organelles may spread through syncytial tissues as a result of a process of subcellular Darwinian selection is proposed. Tissues are likely to be involved in mitochondrial disease if they are syncytial, are derived from a few embryonic cells only, have little redundancy of function, and are subject to repeated metabolic stress. These effects, together with the random distribution of genetically heterogeneous mitochondria within the fertilised zygote, may account for the varied clinical pictures of mitochondrial disease. Evolution will have favoured the shift of mitochondrial DNA sequences to the nucleus, once the differentiation of tissues had created body compartments in which defective mitochondria could flourish to the detriment of the organism. This model of mitochondrial disease allows the generation of several predictions, testable using currently available laboratory techniques. Avenues of potential therapeutic value are indicated, including the avoidance of hypoglycaemia and the use of selective mitochondrial toxins.

There has been much recent interest in the mitochondrial diseases such as Kearns-Sayre syndrome and Leber’s optic neuropathy. This has developed in parallel with sensitive assays of mitochondrial function, especially of the complexes of the respiratory chain. It has been shown that the mitochondrial DNA (mtDNA) of some patients with mitochondrial myopathy (MtM) has suffered substantial deletions or duplications. The abnormal mtDNA is found principally in the mitochondria of muscle, and to a lesser extent in the mitochondria of leucocytes or fibroblasts, and only affects a subpopulation of the muscle mitochondria in each case. It has been suggested that the defective mitochondria would be subject to negative selection in rapidly dividing cell lineages, such as bone marrow stem cells, and would therefore not be found in such tissues. In contrast, relatively stable tissues, such as muscle, would not benefit from this selection and would therefore be subject to mitochondrial disease (MtD). Another suggestion, relevant to the benign, reversible cytochrome c oxidase deficient mitochondrial myopathy only, is that improvement occurs as a result of the impaired viability of muscle fibres containing large numbers of defective mitochondria. Negative selection against mitochondria with deleted genomes may be more intense than it is against those with duplications.

This paper discusses recent clinical and genetic studies of mitochondria and mitochondrial disease, and proposes an explanation of the clinical features of those cases of MtD produced by defects of the mtDNA. This hypothesis does not apply to those mitochondrial disorders caused by defects in the nuclear genome. However, this model accounts for the tissue distribution of MtD, the great clinical variation of disease both within and between families, and the evolutionary tendency for mitochondrial genes to be transferred to the nuclear genome. In addition, several predictions are made that are potentially falsifiable.

Disease model
The central hypothesis is that some defects of mtDNA give a selective advantage to their mitochondrial lineage. It has been clear for many years that defective mitochondria accumulate in the MtMs, to produce the ‘ragged red fibres’: that they accumulate selectively in the fibres with the lowest cytochrome c oxidase activity, in this class of myopathy, has also been known for some time. Indeed,
ovum to and a only mitochondria would factors
selective drive for the healthy
452 supporting the
defective consequences in instances.
This finding of metabolites
red fibre
muscle syndrome
(2)
The chance: there are few defective mitochondria, if the defective mitochondria in an ovum are clustered, as is possible, then the tissues
mitochondrial metabolism
muscle involvement
syndrome
(3) Positive selection: there are several circumstances in which the defective mitochondria of Kearns-Sayre
Darwinian advantage.

(1) Mitochondrial tissue is regularly subject to metabolic stress that must be at a selective
advantage. This is true whether the defective mitochondria have been found in humans, whereas
no such cases have been found in the human genome. To my knowledge, however, all cases of mitochondrially encoded
mitochondria have been found in humans, whereas a few have been found in animals.

(4) Synaptosomes, a neurocyte in a tissue where cells form synapses,
and seventh cases.

(5) Fetal life: with birth comes the switch to fully aerobic metabolism, as the P/O ratio of fetalmitochondria is less than 2 and hence P/O during
recovery from brief anaerobic intervals. The impact on stroke in normal mitochondrial disorders is
mitochondrial dysfunction, and also during spells of relative hypoglycaemia, later in life, and
masquerading as late mtDNA depletion.

(6) Central nervous system: requires functioning
mitochondrial enzymes to use non-glucose fuels such as lactate and ketone bodies to maintain
the brain between periods of fasting. Defective mitochondria would therefore be at a selective advantage
under unusual circumstances.

devolved. Protein synthesis is increased up to tenfold by mendelian
unusually active mitochondrial replication by different circumstances.
and what selective advantage is possible morphologically?
whether or not the defective mitochondria are capable
of replicating, and what selective advantage is possible
under different circumstances.

The nature of the defect: this will determine
mitochondria within a tissue.
There is another hypothesis, that mitochondria normally use a mixture of different nucleo-
plasmic oxidases, as well as the ATPase, and hence their selective
advantage in fetal life, which is lost from the time of
birth. This present severe neuronal hypoxia but then
recover by the child lives at least a few weeks but then
mitochondria, as in the case of Leber's optic
atrophy.

(3) Positive selection: there are several circumstances in which the defective mitochondria of Kearns-Sayre
Darwinian advantage.

(1) Mitochondrial tissue is regularly subject to metabolic stress that must be at a selective
advantage. This is true whether the defective mitochondria have been found in humans, whereas
no such cases have been found in the human genome. To my knowledge, however, all cases of mitochondrially encoded
mitochondria have been found in humans, whereas a few have been found in animals.

(4) Synaptosomes, a neurocyte in a tissue where cells form synapses,
any replicative advantage enjoyed by defective mitochondria could result in a massive dissemination of the defect and a high proportion of the tissue being affected. Cardiac and skeletal muscle are both put at risk of MtM in this way. Cellular tissues with little mitotic activity after normal growth is complete, for example, kidney and liver, will not suffer from MtD even if they initially contain the same proportion of defective mitochondria. There is not the same metabolic stress, and the cell membrane limits any possible spread of the disorder.

(5) Embryonic origin and bulk of tissue: a small muscle derived from only a few embryonic cells will have a smaller chance of containing defective mitochondria than a large muscle. However, if embryonic cells destined to generate an external ocular muscle happen to contain defective mitochondria, and these are favoured by selection, then this is much more likely to cause symptoms than if the same number of fibres were defective in vastus lateralis. The predilection of MtM for the oculomotor muscles and levator palpebrae superioris can be explained in this way. Similarly, specialised tissues consisting of few cells, such as the cardiac conduction tissue, will be at risk of MtD. This may account for the association of endocrine disease with mitochondrial disease, because endocrine tissue bulk is small, and each type of secretory cell derives from only a few cells in the embryo.

(6) Redundancy of cellular function: if a fraction of cells in liver or kidney functions suboptimally, the organ can compensate up to quite generous limits. The same is not true for the central and peripheral nervous systems and for cardiac conduction tissue. Defective Purkinje fibres and peripheral neurones can cause problems at any age. Failure of specific regions of the CNS after plasticity has been lost will also cause problems corresponding to the function that was served by the malfunctioning cell or clone of cells (which are likely to share both the same mitochondrial defect and much the same function within the CNS). This explains why some tissues are especially susceptible to MtDs.

Discussion

It can be seen that a defect in mtDNA could have a selective advantage, could achieve dominance within a tissue, and cause disease. This is particularly so if the tissue is syncytial, is derived from a few embryonic cells only, has little redundancy of function, and is subject to repeated or continuous metabolic stress. That MtM affects skeletal and cardiac muscle, cardiac conduction tissue, and the peripheral and central nervous systems is comprehensible. Each affected family will have a distinct mutation/deletion of the mtDNA in its defective population of mitochondria, so the clinical pattern will vary with the defect in each case. In addition, random processes will determine the distribution of normal and defective mitochondria within the ovum and hence within the embryo. This will result in considerable variation among the affected members of a family. Others have suggested that the tissue specificity of MtD is the result of defects in tissue specific, nuclear encoded proteins. Such defects may exist, but this explanation will not so easily account for cases with known defects in the mitochondrial genome. The possibility that mitochondrial selection may contribute to the clinical course of MtD has been advanced, but not advocated. The present paper elaborates that suggestion and examines the testable consequences. One important item of evidence in favour of this model is the finding that even a single mitochondrion, if its genome differs from the others in the same cell, can exploit a selective advantage in tissue culture to replace the original mitochondrial population of the cell.

A number of cases of MtD are known in which several complexes of the respiratory chain are defective, or in which several RNA or polypeptide chains are lacking. One explanation is that the mtDNA could be deleted for several genes encoding the relevant protein subunits; or possibly tRNA genes could be deleted, which would disrupt the synthesis of several proteins. About 40% of MtM cases have major deletions of mtDNA. Other possibilities, however, do exist. There could be a defect of a (nuclear encoded) structural protein required for the correct attachment and functioning of the respiratory complexes to each other or to the mitochondrial inner membrane. There could be a defect in a mitochondrial receptor for nuclear encoded, cytosol synthesised proteins in whose absence these proteins cannot enter the mitochondrion. All these possibilities will have to be considered as we learn more about the MtDs. In cases where the mtDNA is deleted for tRNA genes, the question arises as to how protein synthesis proceeds at all. Could these defective mitochondria be dependent upon tRNAs produced in other mitochondria, and transported through the cytoplasm or through a mitochondrial web? Such a web of mitochondria may indeed exist, at least in the central mitochondria of muscle, where they form grids around the myofibrils.

CLASSIFICATION

The clinical classification of types of MtD is clearly inadequate, as discussed in a recent, thorough review of such cases. Whereas mtDNA deletions are often associated with ophthalmoplegia, the spectrum of accompanying defects is so wide and variable that a classification in such terms is far too crude to be helpful: it seems that even visceral pathology can result from mtDNA deletions. A genetic classi-
fication may well be appropriate for defects of nucleus encoded genes, since there is likely to be good clinical-molecular correlation. For mtDNA defects, however, there is likely to be only poor clinical-DNA correlation. A dual system of classifying these disorders will be necessary, incorporating both the site, nature, and tissue distribution of the molecular defect, and the clinical presentation.

The one example of good DNA-clinical correlation that might be used to oppose my statement is the case of Leber's optic neuropathy, associated with a single base change at nucleotide 11778. However, this mutation accounts for only a proportion of Leber cases, and the disorder is in any case highly variable even within families, manifesting as a generalised encephalopathy, a movement disorder, cardiac conduction defects, or optic neuropathy. The finding of a biochemical abnormality in the mitochondrial electron transport chain of affected persons is compatible with the intracellular selection model of MtD proposed here.

The accumulation of mtDNA mutations in muscle and other tissues with advancing age has been proposed as an important contributor to ageing and degenerative disease. Certainly, mitochondrial function has been shown to decline with age in 'healthy' subjects. While the advocates of this hypothesis cite the high mutation rate in mtDNA and the lack of a repair system as supporting evidence, I would add that positive selection in favour of new mutation mtDNA defects could occur, rather than their passive accumulation and random segregation between cells. This would naturally accelerate the process of senescence: perhaps mega-doses of ascorbic acid will prove to be beneficial after all!

EVOLUTION
Mitochondria and chloroplasts have most probably evolved from prokaryotic endosymbionts of the early eukaryotes. This helps to explain the resemblance of the transcription and translation systems of these organelles to those of bacteria. The mitochondrial genome of unicellular eukaryotes is larger, and codes for more RNA and protein species than does the mitochondrial genome of man. Thus, yeast mtDNA (78 kb) includes several genes for tRNAs and proteins (some as yet unidentified) that are not found in the smaller (17 kb) human mtDNA. The explanation of this is that the short and long term interests of the yeast mitochondria are identical to those of the whole organism. This is not true of man: the short term interests of defective mitochondria in man can be contrary to their own long term interests, and those of the 'host' organism (the patient with MtD).

Once multicellular organisms developed tissues whose metabolic activities differed substantially, the possibility of mitochondrial disease will have arisen. This will have resulted in selection in favour of the transfer of many functions of the mtDNA into the nucleus. Why this process is not complete, why the mitochondrial genome still exists in man, is not clear. Perhaps the mitochondria have retained little more DNA than is required for them to coordinate their own replication, although we do not know how this is achieved. Alternatively, the existence of multiple copies of the mtDNA dispersed in the cell may be advantageous, and the slight differences between the genetic triplet codes of organelle and nucleus may have made the successful transfer of certain genes to the nucleus into an improbability.

Another evolutionary consequence of the transfer of mtDNA sequences to the nucleus is that populations and species will differ in the functions remaining to the mitochondrial genome. Interspecies hybrids may then be viable, because at least one copy of each gene will be present in the nucleus, but may also be infertile. This would arise if the two species' mtDNAs had lost different genes to the nucleus, because many of the offspring would then inherit some such genes from neither parent. There could, on this model, be differential infertility, depending upon which species had provided the hybrid's mother. Could this effect be important in determining the infertility of interspecies hybrids?

PREDICTIONS AND POSSIBLE THERAPIES
This model of mitochondrial evolution and disease permits certain extrapolations to be made (concerning cases in which the defect is in the mtDNA).

(1) The behaviour of mitochondria in tissue culture from cases of MtM should correlate with the clinical picture. For the severe but reversible neonatal myopathy, the defective mitochondria should have a demonstrable advantage under the metabolic conditions found in utero, and this should be reversed under 'normal' metabolic conditions. The metabolic conditions required to elicit a selection in favour of defective mitochondria in other cases of MtM will probably be more stringent, representing a significant lactic acidosis, or at least a substantial perturbation of intracellular energy metabolism. Such defects should be demonstrable in the myoblasts found in tissue culture of muscle from cases of MtM. Skin fibroblasts from cases of lactic acidosis in infancy have been found to have defects of the mitochondrial respiratory chain; it might be possible to seek evidence of mitochondrial selection in fibroblasts from these patients as well as in muscle tissue. It may also be possible to use the technique of King and Attardi for demonstrating mitochondrial selection with mitochondria from cases of MtD injected into human cells in culture. Their demonstration in vitro that exogenous mitochondria can repopulate human cells lacking mtDNA has established a methodology that
could address the issues raised in this paper, and could be used to test the possible efficacy of potential therapies for patient specific mtDNA defects. These techniques will provide an understanding of how the replication of mitochondria in various MtDs is influenced by the surrounding metabolic milieu, and may provide a means to examine the normal control of mitochondrial replication within different tissues of the higher organisms. There have already been suggestions that variation in mitochondrial function may be partitioned into additive effects of the nucleus and the mtDNA, and into an element accounted for by nucleus-mtDNA interactions.

(2) Another prediction would be that the severe, reversible, neonatal MtM should particularly affect muscles whose blood supply derives from the descending aorta more severely than muscles of the right upper limb and the brain, because these tissues are privileged in utero through receiving blood of higher oxygen content than elsewhere. The defective mitochondria should be less evident in these tissues. This effect could be sought with nuclear magnetic resonance spectroscopy, or through taking muscle biopsy samples from several sites.

(3) According to this model, selection in favour of the mtDNA defect in CNS and retina will be exacerbated by hypoglycaemia. It may therefore be helpful to prevent even trivial hypoglycaemia, and the associated switch to alternative fuels for the brain. To achieve this would necessitate the use of frequent starchy feeds by day and gastric drip feeds by night in children with mitochondrial disease of any sort, as in the management of glucose-6-phosphatase deficiency. Similarly, those with MtM should be encouraged to perform gentle exercise only: strenuous exertion could assist the selection of the defective mitochondria.

(4) Another therapy of possible value would be chloramphenicol: this well known mitochondrial poison may be found to discriminate against the defective mitochondria in cases of MtM. This will have to be tested in vitro, possibly by the techniques of King and Attardi before it is used clinically. Perhaps a continuous, low dose regimen would suppress the disease without causing a ‘grey baby syndrome’. Other antimicrobial drugs, inhibitors of bacterial protein synthesis or replication, could also be considered for such use.

(5) Redox agents that will assist the tissues by circumventing the need for the full oxidative phosphorylation pathway may also be helpful, as previously suggested and practised.

Finally, this proposal provides an example of selfish DNA that could be of interest to evolutionary theorists. As with meiotic drive, the intracellular Darwinian selection of defective mitochondria allows experiments to be devised that will test our models of evolution as well as of disease.

I would like to acknowledge the helpful comments of Drs M Johnson, D Turnbull, and H S A Sherratt.

26 Parker WD, Oley CA, Parks JK. A defect in mitochondrial


