Human mitochondrial leucyl tRNA synthetase can suppress non cognate pathogenic mt-tRNA mutations

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Abstract

Disorders of the mitochondrial genome cause a wide spectrum of disease, these present mainly as neurological and/or muscle related pathologies. Due to the intractability of the human mitochondrial genome there are currently no effective treatments for these disorders. The majority of the pathogenic mutations lie in the genes encoding mitochondrial tRNAs. Consequently, the biochemical deficiency is due to mitochondrial protein synthesis defects, which manifest as aberrant cellular respiration and ATP synthesis. It has previously been reported that overexpression of mitochondrial aminoacyl tRNA synthetases has been effective, in cell lines, at partially suppressing the defects resulting from mutations in their cognate mt-tRNAs. We now show that leucyl tRNA synthetase is able to partially rescue defects caused by mutations in non-cognate mt-tRNAs. Further, a C terminal peptide alone can enter mitochondria and interact with the same spectrum of mt-tRNAs as the entire synthetase, in intact cells. These data support the possibility that a small peptide could correct at least the biochemical defect associated with many mt-tRNA mutations, inferring a novel therapy for these disorders.

Keywords aminoacyl tRNA synthetase; disease; mitochondria; protein synthesis; therapy

Subject Categories Genetics, Gene Therapy & Genetic Disease; Metabolism

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Introduction

Mitochondria are found ubiquitously in nucleated eukaryotic cells. One of their key functions is to generate ATP via oxidative phosphorylation, in order to provide the cell with a readily usable form of energy. The machinery responsible for this process comprises five complexes, members of which are encoded by either the nuclear or the mitochondrial genome (mtDNA). Energy transcription in the cell is, thus, dependent on the precise and accurate intramitochondrial translation of the 13 mtDNA encoded polypeptides. Defects in mitochondrial metabolism are being increasingly recognized as a cause for disease. Indeed, mitochondrial disease is no longer described as a rare disorder, in part as a result of better awareness and therefore more accurate diagnosis. Studies in the north east of England indicate a clinical incidence of 9.2 in 100 000, with a further 16.5 in 100 000 at risk of developing mtDNA disease before retirement (Schaefe et al, 2008). Pathogenesis can arise from defects either in nuclear encoded proteins that function in the mitochondria (Smits et al, 2010; Chrzanowska-Lightowlers et al, 2011; Rotig, 2011; Keogh & Chinnery, 2013) or from mutations in the mtDNA itself (Chinnery & Turnbull, 2000; Greaves et al, 2012). The ease of sequencing the comparatively small mitochondrial genome (Anderson et al, 1981) has allowed a comprehensive characterization of pathogenic mtDNA mutations, establishing that the majority of these mutations reside in the mitochondrial tRNA genes. These would be predicted to impair mitochondrial protein synthesis and since it is currently not possible to manipulate human mtDNA, researchers have explored other ways to overcome defects of mt-tRNAs. Such approaches have often started with yeast as a model system, in which it has been possible to isolate nuclear suppressor factors that are able to rescue the defective phenotype of mt-tRNA mutants (Rinaldi et al, 1997; Feuermann et al, 2003; De Luca et al, 2006). In addition, in a previous study with human cells lines harbouring a pathogenic mt-tRNAval mutation (m.1624C>T) we showed that overexpression of the cognate aminoacyl tRNA synthetase, VARS2, could partially suppress the consequent molecular defect (Rorbach et al, 2008). This work demonstrated that the mutation caused destabilisation of the uncharged mt-tRNAval, such that the transmitochondrial cybrid line carrying the mutation had severely
depleted steady state levels of mt-tRNA\textsuperscript{val}. Following induction, the higher levels of VARS2 increased the proportion of charged mt-tRNA\textsuperscript{val}, thereby increasing the steady state levels (Rorbach \textit{et al}, 2008). Other examples of suppression via overexpression of a cognate tRNA synthetase in human cell lines now exist. One of these is the leucyl tRNA synthetase (Herbert \textit{et al}, 1988), overexpression of which has been reported to suppress the effect of an m.3243A>G mutation in human cells (Li & Guan, 2010). In contrast to the m.1624C>T mutation that is associated with Leigh disease (McFarland \textit{et al}, 2002), this mutation causes a different mtDNA disease, MELAS (Shaag \textit{et al}, 1997).

Genetically correcting an mtDNA defect is not currently a therapeutic option. Moreover human mtDNA encodes 22 different tRNAs, therefore, it would be ideal if a single approach could be developed that would be equally effective with any pathogenic mt-tRNA mutation. With relevance to this, we were aware of an earlier published observation that the yeast mt-leucyl tRNA synthetase (NAM2) possessed generic RNA binding capability (Labouesse \textit{et al}, 1987; Herbert \textit{et al}, 1988) and subsequently that it could suppress non-cognate mt-tRNA mutations in yeast (Montanari \textit{et al}, 2010). We predicted that the human orthologue, LARS2, might also have the potential to counteract generic mt-tRNA defects in human cells. Here we illustrate that in human cells, overexpression of the non-cognate mitochondrial leucyl tRNA synthetase can indeed have a potential to counteract generic mt-tRNA defects in human cells.

Results

As described above, we previously showed that overexpression of VARS2 could partially restore the steady state levels of mt-tRNA\textsuperscript{val} resulting from the m.1624C>T mutation (Rorbach \textit{et al}, 2008). However, this original cybrid population, generated from a fusion of 143B.206 \( \rho^0 \) cells with patient cytoplasts homoplasmic for the m.1624C>T Leigh disease mutation, did not recapitulate the profound biochemical phenotype found in the patient’s skeletal muscle (Rorbach \textit{et al}, 2008). We therefore aimed to exploit the aneuploid nature of the 143B.206 \( \rho^0 \) parental cell line and identify whether isolating clones from the population would expose a stronger metabolic defect in a subset of cells. Clones were derived by serial dilution in glucose. Following expansion, the clones were then independently assessed for growth in galactose, a carbon source that forces cells to use oxidative phosphorylation. Of the 67 clones that were tested only six demonstrated impaired respiratory capacity when challenged to grow on this carbon source. Although a number of these clones were selected for transfection with constructs that would allow inducible expression of aminoacyl tRNA synthetases (aaRS), the data presented here are the characterization of a representative clone (T1) (Fig 1A lanes 2, 6 and 10). Inducibility of aaRS expression was a key factor in the experimental design, as due to the aneuploid nature of 143B.206 cells, it was critical that the behaviour of exactly the same clone could be examined with and without aaRS overexpression. In order to confirm that the respiratory deficiency was attributable to the mt-tRNA\textsuperscript{val} mutation, we transfected the respiratory deficient T1 line with VARS2 (designating it T1V1) and monitored growth on galactose. On induction of VARS2 expression, we observed suppression of the m.1624C>T defect as determined by increased growth on galactose (Fig 1A lanes 7 and 11).

Overexpression of LARS2 can partially rescue the growth phenotype and molecular defects associated with a non-cognate mitochondrial tRNA

The equivalent m.1624C>T MT-TV mutation, as well as the MELAS m.3243A>G MT-TL1 mutation, had previously been introduced into yeast mtDNA and the mitochondrial leucyl or valyl tRNA synthetase were each independently overexpressed. Interestingly, suppression was achieved for the MT-TV mutation using the non-cognate leucyl tRNA synthetase (Montanari \textit{et al}, 2010). More recently it has been shown, again in yeast (Francisci \textit{et al}, 2011), that not only do non-cognate aaRS have the ability to suppress the defects caused by mt-tRNA mutations, but that isolated domains of the aaRS alone could also have a similar effect (Montanari \textit{et al}, 2010; Francisci \textit{et al}, 2011). Since these investigations had been performed in yeast, we decided to use the T1 line harbouring the mt-tRNA\textsuperscript{val} mutation to see if similar suppression by a non-cognate aaRS could be achieved in human cells. In a parallel fashion to the yeast work, we transfected T1 with a FLAG tagged version of the human LARS2, (T1L1). This line together with the T1V1, T1 and 143B.206 \( \rho^+ \) parental cells were routinely propagated on glucose and displayed similar doubling times (Fig 1A lanes 1–4). When transferred to galactose the parental 143B.206 \( \rho^+ \) line continued to grow but with an extended doubling time. In contrast the T1 and aaRS-transfected derivatives were all either unable to grow or showed negligible growth unless cells were induced to express aaRS (Fig 1A cf lanes 7 and 11; 8 and 12). In each cell line the induction of the relevant aaRS was confirmed by western blot analysis (Fig 1B cf lanes 1 and 2; 3 and 4).

Next we measured the steady state levels of respiratory complex proteins to determine whether the partial growth defect suppression truly reflected an improvement in respiratory competence. Following aaRS induction, the levels of mitochondrially encoded COX2 increased. An increase was also observed in the levels of NDUF8, a sensitive marker of Complex 1 (CI) assembly. Nuclear encoded COX4 also appeared to have a modest increase in steady state levels (Fig 1C cf lanes 1 and 2; 3 and 4). Complex II is encoded entirely by the nuclear genome and showed no change when probed for SDHA (Fig 1C). Since the steady state levels are not always a true indicator of complex assembly, Blue Native PAGE was performed with either subsequent western analysis or in gel activity assays. Here again CI appeared unchanged in T1V1 or T1L1 induced cells, however an increase in assembled CI and CIV could be seen, with a more modest increase in CIII (Fig 1D cf lanes 1 and 2; 3 and 4). These increases in complex formation were reflected in the in gel activities for the cell lines expressing either VARS2 or LARS2 (Fig 1E cf lanes 1 and 2; 3 and 4). To directly measure the oxygen consumption we used microscale oxygraphy. Overexpression of either VARS2 or LARS2 resulted in partial recovery of basal and maximal respiration rates, while respiration rates after oligomycin and antimycin inhibition were not altered (Fig 2A). To more accurately assess the
recovery in respiratory chain complex activities, mitochondria were isolated from uninduced and aaRS overexpressing cells, on which spectrophotometric assays were performed for complexes I, II and IV. There was no change in CII activity, whilst induction of aaRS significantly increased CI and CIV activity, where in most cases there was at least a doubling of activity (Fig 2B).

The analyses thus far were directed at recovery of function rather than any direct consequence on the mutant tRNAval. Steady state levels of mt-tRNAs were therefore assessed by high resolution northern. The level of mt-tRNAval was determined and compared to those of mt-tRNA phe and mtRNAleu(UUR). Over-expression of either VARS2 or LARS2 had no effect on either of the two wild type tRNAs but did increase the level of the mutated tRNAval transcript (Fig 2C cf lanes 2 and 3; 4 and 5). Densitometric analysis indicated that in each case, overexpression of aaRS increased the mt-tRNAval to approximately 150% of the levels in uninduced cells (Fig 2C lower panel).

The impact of aaRS overexpression on de novo synthesis of mitochondrial protein was also assessed. Metabolic labeling was performed on induced and uninduced T1V1 and T1L1 alongside 143B.206 parental cells. Following induction of either aaRS, there was an overall increase in labeled protein of approximately 1.5 fold, which corresponded to the increase seen in steady state levels of the mutated mt-tRNAval under the same conditions. Densitometric analysis of individual products indicated 1.6–1.9 fold increases after VARS2 induction (Fig 2D cf lanes 2 and 3) and 1.3–2 fold increases after LARS2 overexpression (Fig 2D cf lanes 4 and 5).

**LARS2 C-term 67 residues can bind to mt-tRNA in intact cells**

As mentioned previously, there is evidence that a C terminal fragment alone, of the mitochondrial leucyl tRNA-synthetases from either yeast or human, is sufficient to suppress the respiratory

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**Figure 1.** Overexpression of VARS2 or LARS2 relieves defects in growth, steady state levels of OXPHOS proteins and complexes. In each case aaRS overexpression was induced by 3 days tetracycline (Tet) treatment, indicated by + or – symbols. All data and images represent a minimum of 3 independent experiments.

A: Cell growth under glycolytic or respiratory conditions with or without overexpression of aaRS. Equal numbers of 143B.206 Rho+, T1, T1V1 and T1L1 cells were seeded in medium containing either glucose or galactose as the sole sugar source. The extent of growth and viability of each cell line with or without aaRS overexpression was determined at 72 h using the neutral red assay. Statistically significant comparisons for uninduced and induced cells are indicated with * (P = 0.016), ** (P = 0.0086) n = 3.

B: Overexpression of aaRS. Mitochondrial proteins were isolated from uninduced and aaRS overexpressors. Mitochondrial lysates from T1V1 (100 μg) and T1L1 (50 μg) were subjected to western blot analysis using antibodies against VARS2L or FLAG to confirm overexpression. Mitochondrial ribosomal protein DAP3 was used to confirm equal loading.

C: Analysis of steady state levels of OXPHOS proteins following aaRS induction. Cell lysates (15 μg) from 143B.206 Rho+ and T1V1 and T1L1 cells with or without induction were subjected to western blot analysis. Membranes were probed with antibodies directed against OXPHOS proteins and porin as a loading control.

D: Blue Native-PAGE analysis of respiratory chain complex levels. Mitochondria (25 μg) from uninduced and induced T1V1 and T1L1 cells were solubilised for BN-PAGE. Subsequent western blot analysis used antibodies against complex I (NDUFA9), complex IV (COX4), complex III (CORE2) and as a non-mitochondrially encoded control, complex II (SDHA).

E: In gel enzyme activity assay of respiratory chain complexes. BN-PAGE was performed on solubilised mitochondria (50 μg) from uninduced and induced T1V1 and T1L1 cells. Enzyme activities for complexes I, IV and II were examined.
defects caused by mutations in yeast mt-tRNAs Leu, Val or Ile (Francisci et al., 2011). Structural studies indicate that this C terminal region of the tRNA synthetase folds into a compact domain that is flexibly linked to the rest of the structure. It also exhibits tRNA binding activity, recognising the mt-tRNA elbow shape rather than sequence (Tukalo et al., 2005). We therefore determined whether

Figure 2. Overexpression of VARS2 or LARS2 suppresses biochemical defects. In each case aaRS overexpression was induced by 3 days tetracycline (Tet) treatment (indicated by + or −), except for XF24 analysis. All data and images represent a minimum of 3 independent experiments. Statistically significant P values are as indicated.

A Oxygen consumption by XF24 analyzer. Representative traces of oxygen consumption rates performed under basal conditions, following the addition of oligomycin (1 μg/ml), uncoupler FCCP (1.5 μM then 3 μM) and antimycin A (2.5 μM) are presented for 143B.206 Rho+, T1, T1V1 and T1L1 cell lines with or without 2 days aaRS overexpression.

B Respiratory chain activity improves on cognate aaRS or LARS2 overexpression. Mitochondria were isolated from 143B.206 Rho+ cells and T1V1 and T1L1 transfectants with and without aaRS induction. Enzyme activities of complexes I, IV and II were measured spectrophotometrically and are presented as ratios against citrate synthase.

C Steady state levels of mt-tRNA\textsuperscript{Val} increase following VARS2 or LARS2 overexpression. Equal amounts of RNA (4 μg) isolated from 143B.206 Rho+ cells and T1V1 and T1L1 transfectants with and without aaRS induction were analysed by high resolution northern blotting. Probes were specific for mitochondrial tRNA\textsuperscript{Leu(UUR)}, tRNA\textsuperscript{Val}, and tRNA\textsuperscript{Phe} or 5S RNA as a loading control. Densitometric measurements of mt-tRNA\textsuperscript{Leu(UUR)}, tRNA\textsuperscript{Val}, and tRNA\textsuperscript{Phe} in uninduced and induced T1V1 and T1L1 transfectants were normalised to the 5S RNA. Data are expressed as percentages of mt-tRNA in induced cells over mean uninduced cell values and presented below *P < 0.02.

D Synthesis of mitochondrially encoded proteins increases upon aaRS overexpression. Mitochondrially encoded proteins in 143B.206 Rho+, T1V1 and T1L1 cells with and without aaRS overexpression, were pulse labeled with \textsuperscript{35}S-methionine and separated by 15% SDS-PAGE. Below the autoradiogram is a western blot of the same membrane probed for β-actin as loading control. Increases in protein synthesis post induction are indicated (right panel) by densitometric quantification of individual RC subunits (designated by *), relative to -Tet and normalized to β-actin.
this fragment alone was able to bind mt-tRNAs other than mt-tRNA-leu(UUR) in human cells. To be sure that we were detecting physiological binding we used the CLIP method. Intact cells are exposed to UV to preserve RNA/protein interactions prior to mitochondrial isolation and immunoprecipitation of the protein of interest. Cells were therefore transfected with various LARS2 constructs that were FLAG tagged to facilitate immunoprecipitation. Although the construct encoding the C-terminal 67 amino acids of LARS2 was FLAG tagged it had no additional N-terminal sequence to promote mitochondrial targeting. However, analysis of the amino acid sequence of the C-terminal peptide using predictive algorithms (including http://www.tcdb.org/progs/helical_wheel.php) does indicate a helix with a clear hydrophobic face but this is only weakly amphipathic. Western blotting of this small fragment confirmed weak but consistent localization to mitochondria after induction. This was performed initially on eluates of FLAG mediated immunoprecipitation from isolated organelles (supplementary Fig S1). We then repeated these studies to identify the exact submitochondrial compartment the C term fragment was able to access. With this aim, mitochondria from cells expressing the C-terminal fragment (Fig 3A lane 1) and mitoplasts were prepared (Fig 3A lane 2) as described in Materials and Methods. These were proteinase K shaved to confirm loss of outer membrane (TOM20), and intermembrane space (AIF) proteins, with retention of the inner membrane (NDUFA9) and matrix (HSP60) proteins (Fig 3A lane 3; 3B lanes 1 and 4). These shaved mitoplasts (Fig 3B lanes 1 and 4) and the postmitochondrial supernatant (PMS; Fig 3B. lanes 3 and 5) were western blotted to confirm that the C-terminal fragment does enter mitochondria. This intramitochondrial matrix localization was further confirmed by the CrossLinking-ImmunoPrecipitation data. CLIP analysis of mRNA-encoded cDNA sequences rescued following crosslinking to physiological RNA substrates and subsequent immunoprecipitation indicated that although the C-terminus showed a significant preference for the two mt-tRNAleu species, it was also able to bind a number of other mt-tRNA species (Fig 3C). Further analysis showed that these binding events were virtually superimposable on the binding pattern of the full length LARS2 (Fig 3D), consistent with suppressive effect seen here and previously in yeast (Francisci et al, 2011). FLAG tagged luciferase was used as a control and showed negligible binding to any mt-RNA (Fig 3E).

Not all mitochondrial tRNA synthetases can rescue the m.1624C→T mutation

Finally, to determine whether suppression of the M·TV mutation could be elicited by overexpression of other mitochondrial aaRS, we transfected T1 with constructs allowing inducible expression of two mitochondrial class II synthetases, either alanyl (AARS2; T1A2) or phenylalanyl (FARS2; T1F2) tRNA synthetase. Overexpression of AARS2 or FARS2 protein had no effect on galactose growth (Fig 4A). Analysis of the steady state level of the mtDNA encoded COXII was performed and confirmed that there was no change following overexpression of either of these aaRS proteins (Fig 4B). This lack of suppression was further confirmed by microscale oxygraphy measurements, where overexpression of non-cognate aaRS, other than LARS2, could not rescue the biochemical deficiency of the T1 mt-tRNAleu mutation (Fig 4C).

Discussion

The data presented here indicate that human mt-leucyl tRNA synthetase (LARS2) has the ability to suppress the biochemical defects seen in mitochondrial metabolism that can result from an mt-tRNA-va

val mutation. It has been previously documented that in cell lines harbouring mitochondrial tRNA mutations, partial or full suppression can be achieved by overexpression of the cognate aminoacyl tRNA synthetase (Park et al, 2008; Rorbach et al, 2008; Li & Guan, 2010). Moreover, there is a naturally occurring example of suppression of a homoplasmic m.4277T→C MT·TV mutation by overexpression of the cognate tRNA synthetase. Here, a family carrying this homoplasmic mutation showed varied levels of penetrance of the clinical and biochemical defect. Tissue specific investigations of the index case and the clinically unaffected mother demonstrated that the mother had naturally elevated levels of iso-leucyl-tRNA synthetase (LARS2) that suppressed the phenotype resulting from the otherwise pathogenic mutation (Perli et al, 2012). Why then should LARS2 have the ability to suppress mutations in non-cognate tRNAs? It appears that LARS2 may have developed a bifunctional role in the cell. Aminoacyl tRNA synthetases are evolutionarily critical proteins that were probably involved in the transition from the RNA world, and appear to have recruited new cell functions (Rho et al, 2002; Sarkar et al, 2012; Yao et al, 2013). Indeed, the yeast LARS2 equivalent, LeuRS was not originally identified by its ability to aminoacylate its cognate tRNA, consistent with it having developed multiple functions. It was identified through a screen originally devised to isolate proteins that when mutated, could compensate for mitochondrial splicing defects in yeast (Dujardin et al, 1980). This screen revealed a number of proteins that showed ‘nuclear accommodation of mitochondria’ and their genes were accordingly classified as NAM1, NAM2 and so forth (Dujardin et al, 1980). It was only following subsequent characterisation that NAM2 was confirmed as encoding the mitochondrial leucyl-tRNA synthetase (LeuRS) (Herbert et al, 1988). Further inspection of the protein domains was strongly suggestive of an RNA binding activity (Labouesse et al, 1985, 1987) that would be consistent with a role in splicing.

The importance of this observation to the work presented here is that splicing factors often act by stabilizing the structure of the catalytically active core of self-splicing RNA transcripts. Importantly, the group I intronic sequences that LeuRS recognizes and is directly involved in splicing in yeast, are highly structured (Michel et al, 1982; Labouesse, 1990) and are distinct from the structures of group II introns (Bonen & Vogel, 2001). It is therefore possible that LeuRS may recognize common structures in mt-tRNAs that are similar to those conserved in group I introns (Michel et al, 1982; Labouesse, 1990). Such an hypothesis has been previously proposed by Myers et al (Myers et al, 2002) to explain the recognition of group I introns by another type 1 mitochondrial aminoacyl tRNA synthetase, tyrosyl-tRNA synthetase (Akins & Lambowitz, 1987; Majumder et al, 1989; Kittle et al, 1991). Clearly, additional sequence or structural specificity will be required for splicing, but at least for LeuRS the result may be a generic ability to bind to non-cognate mt-tRNAs.

The splicing function and thus part of the RNA binding activity was initially identified in the yeast LeuRS by deletion analysis, showing that this activity resided in the C terminal region of the protein, and that these changes essentially had no impact on the
aminacylation activity (Li et al., 1996). This C-terminal domain is highly conserved, in amino acid sequence and in structure, from Archaea to human mitochondria, but has been lost in the mammalian cytosolic forms (Cusack et al., 2000; Tukalo et al., 2005; Hsu et al., 2006). Further, although there is no requirement for intron splicing in human mtDNA expression, the human LARS2 is still able to restore splicing in a yeast NAM2 delete strain (Houman et al., 2006). More recent work has highlighted how this C-terminal is working. Although the ~60 amino acids in this C-terminal extension form a unique domain that appears to have evolved to optimize its multiple roles in aminocytosine and splicing (Hsu et al., 2006), the C-terminal domain of LeuRS does not interact with the anticodon of its cognate tRNA (Larkin et al., 2002). This region interacts directly with the elbow of the L-shaped tRNA and is based more on structural recognition of the tRNA shape rather than through specific base-base interactions (Tukalo et al., 2005; Hsu et al., 2006).

Why should an aaRS, which should have tRNA sequence specificity, develop recognition of shape over sequence? The answer may be that despite their predicted stable ‘cloverleaf’ structure, organelar tRNAs appear to have greater structural relaxation than bacterial tRNAs (Fender et al., 2012) so there is arguably a greater need for interacting proteins to stabilize their shape. In support of this is the observation that a number of in vitro aminacylation assays show greater activity if a polyamine is preincubated together with the substrate tRNA (Bullard et al., 1999). In particular, the addition of spermine is believed to stabilize the cloverleaf structure of certain tRNAs (Pegg, 1988). Curiously this enhancement of aminocytosine was not observed in assays with the human mt-leucyl tRNA synthetase (Bullard et al., 2000). This would support our hypothesis that LARS2 itself has a stabilizing effect on the tRNA substrate, which in turn will promote more efficient aminacylation, particularly in the case of destabilizing mutations.

Finally, is this non-cognate mt-tRNA stabilizing function unique to the C-terminal of LARS2? The data in this paper only minimally address this issue. We find that representatives of the class II mitochondrial aaRSs are unable to even partially rescue growth of the MT-TV mutation. However, the accompanying paper shows that at least one other type I tRNA synthetase, VARS2 is capable of cross rescuing non-cognate mt-tRNA defects in human cell lines. This may also be the case for other mitochondrial aaRS, but we are unable to comment directly on this. However, it is interesting to consider that leucyl tRNA synthetase belongs to a subfamily of large monomeric synthetases that includes valyl- and isoleucyl-tRNA synthetases. Further, the yeast mitochondrial orthologues of each of

![Figure 3. LARS2 C-terminus binds mitochondrial tRNA species in vivo.](image)

In each case overexpression of the LARS2 FLAG tagged C-terminus was induced by 3 days tetracycline (Tet) treatment. Samples presented in panels A and B were separated on a single 16% Tricine PAG. Post transfer the membrane was cut to allow probing with different antibodies, positions of the molecular weight markers (MWM) are indicated. Samples in C, D and E represent RNA binding determined by crosslinking/immunoprecipitation (CLIP). Derived data is presented as Ion Torrent reads aligned against the mtRNA genes, which are depicted as in circular mtDNA. Concentric circles indicate a log scale of number of reads / site. A Proteinase K (PK) shaving of mitoplasts confirms loss of OMM and IMS but retention of IMM and matrix fractions. Mitochondria (50μg, lane 1), mitoplast (75μg, lane 2) and shaved mitoplast (50μg, lane 3) fractions were analysed by western with the antibodies indicated. B LARS2 C-terminal peptide is localized to the inner mitochondrial compartment. The shaved mitoplasts (~1mg of preparation presented in panel A; lanes 1 and 4) and TCA precipitated postmitochondrial supernatant (PMS; lanes 3 and 5) were probed for the presence of the LARS2 C-terminus. COXI was used as a marker to confirm presence of inner mitochondrial proteins that were absent from the PMS fraction. The poor resolution is due to the large amount of protein loaded in this well. Left Panel directly visualized by BioRad Chemidoc MP, right panel visualized by autoradiography. C–E CLIP derived mt-RNA binding capacity. (C) Physiological RNA binding by the LARS2 C-terminal peptide; (D) Physiological RNA binding by the full length LARS2; (E) As a control, RNA binding by the mitochondrially targeted FLAG tagged luciferase was also determined.
these proteins has been shown to suppress non-cognate mt-tRNA mutations (Francisci et al., 2011). Perhaps this subfamily of aminoacyl tRNA synthetases has retained an RNA binding feature that can be exploited therapeutically. Preliminary analysis of hydropathy profiles or primary amino acids sequences showed no obvious motifs but more detailed investigation of whether there are any characteristics shared by members of this family will form part of our ongoing studies.

In summary, we have shown that overexpression of the human mitochondrial leucyl tRNA synthetase can be used to suppress the defects caused by a destabilizing mutation in mt-tRNAval, a non-cognate mt-tRNA. Further, CLIP data of RNA binding derived for the C-terminus of LARS2 demonstrated an almost identical binding spectrum to mitochondrial tRNAs as the full length LARS2. We believe that this provides good evidence for the potential therapeutic use of the C-terminus of LARS2 for most pathogenic human mt-tRNA mutations.

**Materials and Methods**

**Cell culture**

Human 143B.206 Rho⁺, cybrid derivatives and Flp-InT-Rex-293 cells (HEK293T; Invitrogen Life Technologies Ltd, Paisley, UK) cells were cultured (37°C, humidified 5% CO₂) in DMEM (Sigma-Aldrich Company Ltd., Dorset, UK) supplemented with 10% (v/v) foetal calf serum (FCS), 1× non-essential amino acids (NEAA) and 2 mM L-glutamine and where appropriate with the addition of 10 μg/ml BlasticidinS and HygromycinB (100 μg/ml) post transfection to select for successful integration. For growth on respiratory substrates, the medium contained glucose-free DMEM (Gibco Life Technologies Ltd, Paisley, UK), 0.9 mg/ml galactose, 1 mM sodium pyruvate, 10% (v/v) FCS, NEAA and 2 mM L-glutamine and 50 μg/ml uridine. In each case induction of aaRS in each cell line was achieved by addition of 1 μg/ml tetracycline for 3 days.

**Neutral red cell counting assay**

The number of living cells was evaluated by automated counting and neutral red assay as described in Repetto et al. (2008).

**Cell lysate, westerns and antibodies**

SDS–PAGE analysis was performed on cell lysate extracted from cultured cells as described previously (Soleimanpour-Lichaei et al., 2007). After electrophoresis, gels were transferred to a PVDF membrane (GE Healthcare, Amersham, UK) and processed for immunoblotting. The following commercially available antibodies were used: NDUFB8, NDUFA9, Core2, COXI, COXII, COXIV, SDH, Complex V α subunit, porin (MitoSciences, Eugene, OR, USA); cytochrome c (BD Biosciences, Oxford, UK); β-actin, FLAG (Sigma Life Science); DAP3 (Abcam, Cambridge, UK); TOM20 (Santa Cruz, Heidelberg, Germany); AIF (NEB, Hitchin, UK); HSP60 (BD Biosciences); VARS2 custom synthesized; all secondaries were HRP conjugated (Dako, Stockport, UK).

**Spectrophotometric activity assays**

The activities of complex I, II and CIV and citrate synthase, as a mitochondrial matrix marker, were determined in isolated mitochondria as previously described (Kirby et al., 2007).
Oxygen consumption measurements

Measurement of intact cellular respiration was performed using the XF24 analyzer (Seahorse Bioscience, Saint Marcell, France). Cells were plated at a density of 10 000 cells/well on XF24 tissue culture plate. After 4 h cells were induced by adding tetracycline (1 µg/ml), which was retained in the culture media for 2 days. Prior to the respiration assay, cells were rinsed and cultured in assay medium supplemented with 5 mM glucose 10 mM sodium pyruvate (Sigma) according to manufacturer’s protocol. Cells were incubated at 37°C in a CO2 free incubator for 1 h prior to measurement. Oxygen consumption was measured under basal conditions, and in the presence of oligomycin, a complex V inhibitor (1 µg/ml), the complex III inhibitor antimycin (2.5 µM), and mitochondrial uncoupler FCCP (first addition at 1.5 µM, second addition at 3 µM, Sigma) to assess maximal oxidative capacity. To normalize respiration rates to cell number, cells were fixed with 4% paraformaldehyde and counted after nuclear staining with Hoechst 33258 (1 µg/ml, Invitrogen).

Mitochondrial preparation

Mitochondria were isolated using superparamagnetic microbeads as described previously (Hornig-Do et al., 2009). For the confirmation of intramitochondrial localization of the C-terminal fragment of LARS2, preparations were as follows. HEK-293T cells were homogenized (glass:Teflon dounce homogeniser, 20 passes) on ice in isolation buffer (10 mM Tris-HCl, pH 7.4, 0.6 M mannitol, 1 mM EGTA and 0.1% BSA). Aggregates were removed by centrifugation at 400 g for 10 min at 4°C. Mitochondria were pelleted at 11 000 g for 10 min at 4°C. The post-mitochondrial supernatant was TCA precipitated before resus- pension in dissociation buffer. Pelleted mitochondria were washed (in isolation buffer) and mitoplasts generated by hypotonic shock in ice cold 5 mM Tris-HCl pH 7.4, 1 mM EDTA. Mitoplasts were Proteinase K treated (5 µg/mg of mitochondria) in isolation buffer lacking BSA, for 30 min at 4°C. Reactions were then stopped by the addition by 1 mM PMSF and washed in isolation buffer lacking BSA.

Metabolic labelling of mitochondrial translation products

Essentially as described in Chomyn (1996). Cells were labeled with 35S-methionine (0.05 mCi/25 cm² flask) for 1.5 h in methionine- and cysteine-free DMEM supplemented with 100 µg/ml metemine and 5% dialyzed FBS (Chomyn, 1996). Aggregates were removed by centrifugation at 400 g for 10 min at 4°C. Mitochon- dria were pelleted at 11 000 g for 10 min at 4°C. The post-mitochondrial supernatant was TCA precipitated before resuspension in dissociation buffer. Pelleted mitochondria were washed (in isolation buffer) and mitoplasts generated by hypotonic shock in ice cold 5 mM Tris-HCl pH 7.4, 1 mM EDTA. Mitoplasts were Proteinase K treated (5 µg/mg of mitochondria) in isolation buffer lacking BSA, for 30 min at 4°C. Reactions were then stopped by the addition by 1 mM PMSF and washed in isolation buffer lacking BSA.

BN-PAGE and enzymatic in-gel-activity

BN-PAGE analysis was performed as previously described (Hornig-Do et al., 2012). Briefly, mitochondria were solubilised by dodecyl-maltoside (DDM, Sigma) at 2 g/g protein and incubated for 20 min on ice. The supernatant was collected after 20 min centrifugation at 25 000 g. To resolve individual complexes and smaller supercomplexes, 25 µg of DDM treated mitochondrial membrane proteins were separated on 4.5–16% gels (Wittig et al., 2006). Post electrophoresis, complexes were transferred to PVDF membranes and sequentially probed with indicated antibodies. In-gel complex activ- ity assays were carried out to estimate the activity of CI, CII and CIV as described (Calvaruso et al., 2008).
The paper explained

Problem
Mitochondrial DNA (mtDNA) disease has been recognized for over 20 years. As diagnostic tools have been refined, it is clear that mtDNA disease should no longer be regarded as a rare disorder. The majority of pathogenic mutations occur in the mtDNA-encoded mt-tRNAs, with more than 200 independent mutations having been described. These result in defects of mitochondrial protein synthesis, causing aberrant oxidative phosphorylation and a panoply of clinical presentations. To date, there is no effective cure for these disorders.

Results
Each aminoacyl tRNA synthetase has a specificity for its cognate tRNA. This process of recognition ensures that tRNAs are charged with the correct amino acid and that the fidelity of DNA sequence to protein composition is maintained. In contrast to this we show that overexpression of a non-cognate mitochondrial aminoacyl tRNA synthetase can overcome the respiratory defect caused by an mt-tRNA mutation. The pathogenic mt-tRNA\textsuperscript{Val} mutation, which causes a loss of transcript for non-cognate mt-tRNAs is weaker than its affinity for its cognate mt-tRNA\textsuperscript{Val} but is sufficient to be detected by cross linking and immunoprecipitation. Moreover, we have identified that this same mt-tRNA binding pattern is achieved with a short peptide corresponding to the C-terminus of LARS2.

Impact
Gene therapy with individual mitochondrial aminoacyl tRNA synthetases is theoretically a possibility but not an excessively practical one. Having a single therapeutic agent that could ameliorate mtDNA disease caused by any mt-tRNA mutation would be a huge step forward in terms of treatment. The possibility of using a short peptide that can intrinsically target mitochondria and effect therapeutic potential on all 22 mt-tRNA substrates is worthy of further investigation.

Identification of oligoribonucleotides bound in vivo
To identify physiological interactions between proteins and mitochondrial RNAs cross linking immunoprecipitation (CLIP) was carried out essentially as previously described (Dennerlein et al., 2010). To generate a more comprehensive data set, the final PCR products were prepared for Ion Torrent sequencing, following manufacturer’s instructions, rather than subcloning into pCR4-TOPO. Sequence data for 100 000–190 000 reads was collected, aligned to mtDNA as a reference sequence using the Torrent Suite software on the Ion Torrent server. The alignments were then viewed using IGV (integrative genomics viewer) and presented against a circular depiction of human mtDNA, using a log scale to indicate the number of reads per RNA site.

Supplementary information for this article is available online: http://embom.m.embopress.org/

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Author contributions
ZCL and RNL were involved in hypothesis derivation, experimental design and execution, interpretation and funding project. HTHD made substantive contributions to acquisition, presentation and analysis of data. AM constructed plasmids and assisted in CLIP experiment. AR performed LARS2 CLIP experiments and preparation of these samples for ion torrent sequencing. HT performed Ion Torrent sequencing of barcoded cDNA libraries. AA constructed plasmids. DPA-K generated and cloned the initial hybrid population and identified the respiratory deficient clones. LF and SF made contributions to the design of the experiment, interpretation and funding project. AM contributed to acquisition, presentation and analysis of data. SF studied the C-terminal domain sequences to be cloned. All authors critically discussed results and approved final version of the manuscript.

Conflict of interest
The authors declare that they have no conflict of interest.

References
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Francisci S, Montanari A, De Luca C, Frontali L (2011) Peptides from aminoacyl-tRNA synthetases can cure the defects due to mutations in mt tRNA genes. Mitochondrion 11: 919 – 923


Labouesse M (1990) The yeast mitochondrial leucyl-tRNA synthetase is a splicing factor for the excision of several group I introns. Mol Gen Genet 204: 209 – 221


Li GY, Becam AM, Slonimski PP, Herbert CJ (1996) In vitro mutagenesis of the mitochondrial leucyl tRNA synthetase of Saccharomyces cerevisiae shows that the suppressor activity of the mutant proteins is related to the splicing function of the wild-type protein. Mol Gen Genet 252: 667 – 675


