Natural and Induced Mitochondrial Phosphate Carrier Loss: DIFFERENTIAL DEPENDENCE OF MITOCHONDRIAL METABOLISM AND DYNAMICS AND CELL SURVIVAL ON THE EXTENT OF DEPLETION.

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Edited by F. Anne Stephenson

The relevance of mitochondrial phosphate carrier (PiC), encoded by SLC25A3, in bioenergetics is well accepted. However, little is known about the mechanisms mediating the cellular impairments induced by pathological SLC25A3 variants. To this end, we investigated the pathogenicity of a novel compound heterozygous mutation in SLC25A3. First, each variant was modeled in yeast, revealing that substituting GSSAS for QIP within the fifth matrix loop is incompatible with survival on non-fermentable substrate, whereas the L200W variant is functionally neutral. Next, using skin fibroblasts from an individual expressing these variants and HeLa cells with varying degrees of PiC depletion, PiC loss of ~60% was still compatible with uncompromised maximal oxidative phosphorylation (oxphos), whereas lower maximal oxphos was evident at ~85% PiC depletion. Furthermore, intact mutant fibroblasts displayed suppressed mitochondrial bioenergetics consistent with a lower substrate availability rather than phosphate limitation. This was accompanied by slowed proliferation in glucose-replete medium; however, proliferation ceased when only mitochondrial substrate was provided. Both mutant fibroblasts and HeLa cells with 60% PiC loss showed a less interconnected mitochondrial network and a mitochondrial fusion defect that is not explained by altered abundance of OPA1 or MFN1/2 or relative amount of different OPA1 forms. Altogether these results indicate that PiC depletion may need to be profound (>85%) to substantially affect maximal oxphos and that pathogenesis associated with PiC depletion or loss of function may be independent of phosphate limitation when ATP requirements are not high.

The mitochondrial phosphate carrier (PiC), encoded by the nuclear gene SLC25A3, catalyzes the transport of inorganic phosphate (Pi) into the mitochondrial matrix (for a review, see Ref. 1); transport is electroneutral and either in symport with H+ or in exchange for OH−. PiC is a typical member of the family of mitochondrial inner membrane carriers with six transmembrane segments, 3-fold symmetry, and N and C termini projecting into the intermembrane space (2). SLC25A3 has alternatively spliced forms of exon 3 (referred to as exon 3A and 3B) giving rise to PiC-A and PiC-B isoforms. PiC-B is expressed ubiquitously, whereas PiC-A is confined to cardiac and skeletal muscle (3, 4). PiC-A has a 3-fold higher transport affinity for Pi, whereas PiC-B has a 3-fold higher maximal transport rate (5). Turnover rate has been estimated at 50,000/s at 25 °C, which is high among carriers (6).

The transport characteristics of PiC favor it as the most important, or only, means of supplying Pi for oxidative phosphorylation (oxphos) as well as for other Pi-dependent processes within the matrix such as Ca2+ buffering. The functional importance of the PiC was first demonstrated in yeast (Saccharomyces cerevisiae) that exhibited delayed growth or failed to grow on non-fermentable substrate when deleted for MIR1, the yeast homologue of PiC (7). That the yeast ΔMIR1 mutant fails to thrive specifically when grown on mitochondrial substrates supports the relevance of MIR1 for oxphos.

Fundamental aspects of Pi transport into mammalian mitochondria and its role in oxphos have been known for decades. However, only very recently has the role of PiC in oxphos and other aspects of mitochondrial function such as in control of the permeability transition pore been directly tested in mammals. This was achieved using mouse models of conditional heart-specific PiC depletion (8, 9) and depletion of PiC in HeLa cells.

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The abbreviations used are: PiC, mitochondrial phosphate carrier; oxphos, oxidative phosphorylation; PiC-A, PiC isoform A; PiC-B, PiC isoform B; Ctrl, control; JO2, O2 consumption; PA-GFP, photoactivatable green fluorescent protein; RPA, region of photoactivation; MFN1, mitofusin 1; MFN2, mitofusin 2; kd, knockdown; FCCP, carbonyl cyanide p-trifluoromethoxyphenylhydrazone; mDsRed, mitochondrial matrix-targeted DsRed1; mPA-GFP, mitochondrial matrix-targeted photoactivatable GFP; PiC mutant, individual expressing the compound heterozygous PiC mutation; MAS, mitochondrial assay solution; Hum, humanized; ANOVA, analysis of variance.

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cells (10) and through the discovery of pathologic variants in the human gene (11–13). These variants are both in exon 3A, specifically c.158–9A→G and c.215G→A, which each cause a frameshift and absence of protein. Functional analyses in skeletal muscle biopsy from homozygous individuals revealed greatly suppressed pyruvate and succinate oxidation rates despite elevated rates of maximal electron transport chain activity (11). Phenotypically, the individuals harboring these exon 3A mutations displayed muscle-related abnormalities soon after birth, namely elevated blood lactate, hypertrophic cardiomyopathy, and skeletal muscle hypotonia.

The first variants in SLC25A3 that potentially impact both PiC-A and PiC-B were recently described by one of us in a case report (13). An infant was diagnosed prenatally with cardiac hypertrophy, subsequently required a heart transplant, and is stable. There was no evidence of lactic acidosis or of skeletal muscle dysfunction. Whole-exome sequencing revealed a compound heterozygous mutation in SLC25A3: c.599T→G and c.215G→A, which each cause a frameshift and absence of protein. Functional analyses in skeletal muscle biopsy from homozygous individuals revealed greatly suppressed pyruvate and succinate oxidation rates despite elevated rates of maximal electron transport chain activity (11). Phenotypically, the individuals harboring these exon 3A mutations displayed muscle-related abnormalities soon after birth, namely elevated blood lactate, hypertrophic cardiomyopathy, and skeletal muscle hypotonia.

The identified human PiC mutations were each transformed into the Δmir1 yeast strain (Fig. 1A). Additionally, a humanized version of yeast Pic1p (PIC^human) was included to delineate consequences of the mutation from that of the inherent differences between human and yeast proteins. The PIC^L200W missense variant, which models the point mutation, was similar to wild-type (WT) human PiC in terms of protein amount and functional growth (Fig. 1, A and B). Furthermore, there was no expression or growth defect observed for PIC^L200W even at elevated temperature (37 °C) (Fig. 1, C and D). In contrast, the PIC^QIP variant, which models the GSSAS → QIP mutation, resulted in a slightly decreased protein level (see Fig. 1A, compare QIP with Hum) and the inability to survive in respiratory medium (Fig. 1B), consistent with a defect in ophos. These effects cannot be attributable to the humanized construct because reconstitution with PIC^human, which expresses the wild-type GSSAS sequence, showed comparable expression and functionality to WT yeast at both optimal and higher temperatures (Fig. 1, A–E).

Because these novel PiC mutations were identified in a compound heterozygote, we used the yeast model to ask whether PIC^QIP could act in a dominant negative manner. To this end, pRS316PIC^QIP was transformed into Δmir1(pRS315), Δmir1(pRS315PIC1), Δmir1(pRS315PIC^L200W), or Δmir1(pRS315PIC^human) strains. Growth on fermentable substrate was normal for all strains (Fig. 1D) and was hindered on non-fermentable substrate only when two copies of PIC^QIP were expressed. Thus, PIC^QIP is not sufficient to cause problems in expression and respiratory growth in yeast when a functional
Delayed Proliferation and a Lower Functional Respiratory Capacity in PiC Compound Mutant Fibroblasts—We obtained primary skin fibroblasts from the individual expressing the compound heterozygous PiC mutation (PiC mutant) (13) as well as from four control (Ctrl) individuals. The tendency for a lower expression of PiC protein in the yeast cells transformed with the QIP mutation prompted us to evaluate PiC mRNA and protein expression in the fibroblasts. For quantitative PCR, primers recognizing mRNA for the PiC-A and PiC-B isoforms were validated using human skeletal myoblasts, which are known to predominantly express PiC-B and then to up-regulate PiC-A upon differentiation into myotubes (Fig. 2A). Quantitative PCR analysis using these validated primers revealed unchanged mRNA levels for both isoforms in PiC mutant fibroblasts (Fig. 2A; note that PiC-B is the predominant isoform in fibroblasts). Differently, immunoblot analysis using an antibody raised against the full-length PiC protein showed that PiC expression was ∼40% of the level in Ctrl fibroblasts as determined using fibroblasts from four healthy individuals (Fig. 2B; the representative blot shows two of the control fibroblast lines); the antibody typically revealed two bands, and only the upper band was quantified. Lower molecular weight fragments, which would indicate cleavage products, were not detected. In contrast to this lower PiC protein abundance, the protein expression level of many other mitochondrial proteins was similar between PiC mutant and Ctrl cells (Fig. 2B), pointing to a selective decrease in PiC protein in PiC mutant cells.

To investigate the effect of the PiC compound heterozygous mutation on proliferative capacity in a mammalian system, the primary skin fibroblasts were grown in parallel with Ctrl skin fibroblasts in different media. PiC mutant cells grown in glucose-rich medium appeared healthy but divided more slowly with a doubling time that was approximately twice the Ctrl rate (Fig. 3, upper and middle panels) whether cells were grown in 5 mM glucose or in 25 mM glucose. When cells were challenged with medium free of glucose and containing only mitochondrial substrates (β-hydroxybutyrate and glutamine), Ctrl fibroblasts continued to proliferate although at a much slower rate. In contrast, PiC mutant fibroblasts could not maintain proliferation when glucose was unavailable (Fig. 3, lower panel).

Although the yeast models indicated the lack of a dominant negative effect of the GSSAS → QIP mutation, the patient-derived fibroblasts suggested that the presence of the mutation rendered the PiC protein unstable, leading to substantially lower levels of PiC protein, which could suppress oxphos. To determine whether this occurred, O2 consumption (JO2) was measured. Intact PiC mutant fibroblasts grown and studied in low or high glucose (plus 4 mM glutamine) had ∼50% of the basal mitochondrial JO2 (i.e. minus the rate measured after addition of antimycin to inhibit complex III of the electron transport chain) of the Ctrl cells (Fig. 4A shows raw O2 traces from an experiment, and Fig. 4B shows group averages). To indirectly determine whether a Pi limitation on oxphos could account for the lower basal JO2, we evaluated the leak-dependent mitochondrial JO2 (i.e. after addition of oligomycin to inhibit the ATP synthase) and theuncoupler-driven JO2 (using FCCP to evaluate electron transport chain capacity under the prevailing substrate conditions); if a Pi limitation on oxphos was the sole driver of the lower basal JO2 in PiC mutant cells, then leak-dependent and uncoupled JO2 should remain similar to that in Ctrl cells. Differently, both of these were lower in PiC mutant cells, pointing to a broader defect that results in not only lower basal JO2 but also decreased functional capacity of the electron transport chain.
To further investigate the possibility of a \( P_i \) limitation on oxphos in PiC mutant fibroblasts, \( J_{O_2} \) was measured in permeabilized cells supplied with either pyruvate + malate (Fig. 4B) or succinate + rotenone (not shown) and in the presence of saturating ADP (maximal phosphorylating \( J_{O_2} \) state 3), followed by addition of oligomycin and then of FCCP. There were no differences between Ctrl and PiC mutant cells for either of the substrates or any of the conditions. Notably, state 3 \( J_{O_2} \) was not different, indicating the availability of sufficient \( P_i \) to drive maximal oxidative phosphorylation in PiC mutant cells.

**HeLa Cells with Substantial PiC Depletion Can Sustain Oxidative Phosphorylation—**The continued presence in PiC mutant fibroblasts of oxphos, despite the ~60% lower PiC protein abundance, suggested either that the capacity of PiC is far in excess of its requirement for oxphos in these cells or that alternative sources of \( P_i \) are available to the matrix. To further test whether lower PiC can be compatible with substantial oxphos, PiC was acutely depleted in HeLa cells using siRNA; two different siRNA duplexes were investigated (kd1 and kd2; Fig. 5). By 72 h post-siRNA introduction, PiC protein was ~15% of control levels (Fig. 5A). In contrast, the protein expression...
level of other mitochondrial proteins was unchanged 72 h post-siRNA treatment (Fig. 5A). Cell proliferation, measured using cells seeded at 48 h post-siRNA addition, was delayed by PiC depletion (Fig. 5B). Measurements of JO2 in permeabilized cells revealed a 30–40% suppression of state 3 JO2 with either pyruvate/malate (Fig. 5C) or succinate (rotenone; not shown), whereas leak-dependent JO2 and uncoupled JO2 were unchanged by PiC knockdown (Fig. 5C); these results are consistent with a Pi limitation on oxphos as the cause of the lower state 3 JO2. In intact cells, substantial PiC depletion had no effect on basal, leak-dependent, or uncoupler-driven JO2. We further investigated oxphos in HeLa cells that had a lesser acute depletion of PiC; cells were depleted by 60–65% (Fig. 5D), which is similar to the extent of the PiC decrease in PiC mutant fibroblasts (Fig. 2B). In contrast to permeabilized HeLa cells with ~85% decrease in PiC, permeabilized HeLa cells expressing slightly more PiC had the same maximal phosphorylating JO2 as Ctrl HeLa cells. Altogether, the data from intact and permeabilized HeLa cells with different levels of PiC depletion further support that at least some oxphos can be maintained despite substantial PiC depletion.

Altered Mitochondrial Morphology in PiC Compound Mutant Fibroblasts and in HeLa Cells with Acute PiC Depletion—Emerging literature supports a mutual relationship between mitochondrial function and form (15). Indeed, visualization of mitochondrial morphology in fibroblasts expressing mitochondrial matrix-targeted DsRed1 (mtDsRed) revealed fewer elongated mitochondria in PiC mutant than in Ctrl fibroblasts (Fig. 6A and B). Furthermore, in HeLa cells with ~60% decrease in PiC a similar change in the mitochondrial form was obtained (Fig. 5C). A decrease in the fraction of elongated organelles can result from decreased mitochondrial fusion or increased mitochondrial fission.

Less Fusion Activity in PiC Compound Mutant Fibroblasts and in HeLa Cells with Acute PiC Depletion—To evaluate directly whether the distinct mitochondrial morphology in PiC patient and control fibroblasts can result from different levels of interorganellar continuity and fusion events, the cells were co-
PiC in Mitochondrial Metabolism, Dynamics, and Cell Survival

FIGURE 6. Mitochondrial morphology in primary PiC mutant fibroblasts and PiC-deficient HeLa cells. Mitochondrial morphology was scored as follows: fragmented, mainly small and round; partly fragmented, intermediate, mixture of round and shorter tubulated; normal, tubulated, long and higher interconnected; elongated, very long, tubulated. The percentage of cells with the indicated mitochondrial morphologies was determined as a percentage of the total number of DsRed-transfected cells counted (≥18 cells per experiment; n = 4 independent experiments). A, representative figures of the different mitochondrial forms in Ctrl and PiC mutant fibroblasts. B, distribution of the different type of mitochondria in primary fibroblasts. C, distribution of the different types of mitochondria after PiC silencing in HeLa cells. Scr., scrambled siRNA; PiC kd1 and kd2, two different PiC-specific duplexes.

A

B

C

MtDsRed

MtPA-GFP

Percentage of cells

Ctrl

PiC mutant

Percentage of cells

Scr.

PiC kd1

PiC kd2

Discussion

The novel compound heterozygous mutations in SLC25A3 described here and previously (13) were associated in vivo with respiratory distress and hypertrophic cardiomyopathy but with unremarkable function of other organs, including skeletal muscle (13). The pathological potential of the novel SLC25A3 mutations was not tested directly and was our aim. Here we report that the point mutation is likely to be functionally neutral, whereas the GSSAS → QIP variant has relevance compatible with a loss of function. Furthermore, the GSSAS → QIP variant does not obligatorily act as a dominant negative. However, in primary fibroblasts harboring the compound heterozygous PiC mutation and thus one potentially functional allele, PiC protein abundance was substantially lower, and the cells were not neotypically normal; the most dramatic phenotype was a proliferation time that was approximately half the control rate and an inability to divide when supplied with only mitochondrial substrates. PiC mRNA was unaffected by the mutations. Thus the compound heterozygous PiC mutation has a functional impact in mammalian cells with instability of the PiC protein as a likely initiating factor. We also found in patient fibroblasts, as well as HeLa cells with different levels of acute PiC loss, that PiC protein could be substantially depleted before oxidative phosphorylation was suppressed. This suggests that the PiC can be expressed in excess of oxphos needs or that other PiC transport mechanisms can compensate. However, PiC mutant fibroblasts could not proliferate in mitochondrial substrate, suggesting that a partial deple-

transfected with cDNA encoding mtDsRed and mitochondrial matrix-targeted photoactivatable GFP (mtPA-GFP). Using confocal microscopy, a time series of fluorescence images was recorded, and 25-μm² square-shaped areas were illuminated by a pulsed laser to photoactivate mtPA-GFP (16, 17). Mitochondrial matrix continuity and connectivity are unveiled by the diffusion of the photoactivated PA-GFP to the regions outside the 2 photon illuminated area. Image time series indicated less mtPA-GFP diffusion in PiC mutant than in Ctrl fibroblasts (Fig. 7A). To quantify connectivity, the time course of the ratio of $F_{mtPA-GFP}$ / $F_{mtDsRed}$ was calculated for the region of photoactivation (RPA) (Fig. 7A). The slower decay of the fluorescence ratio in the RPA indicates a decrease in the combined activity of mitochondrial network formation (40 s) and mitochondrial fusion and mitochondrial movements (500 s) in the PiC mutant fibroblasts (Fig. 7B). To validate fusion events, reciprocal spreading of mtPA-GFP and mtDsRed among mitochondria that were not continuous at the time of mtPA-GFP photoactivation was sought. The fusion events were quantitatively analyzed by the progression of the distribution of mtPA-GFP fluorescence between the images collected within the first 24 s and 8 min after photoactivation. The result of manual counting indicates less mitochondrial fusion activity in the PiC mutant fibroblasts. The total fusion numbers were decreased significantly in the mutant fibroblasts (Fig. 7C). Although Fig. 7C shows fusion event number for only one control fibroblast line that was studied side by side with the PiC mutant, the fusion event numbers were very similar in other control fibroblast lines (7.3 ± 0.3 versus 6.5 ± 0.1, 7.5 ± 0.1, and 6.4 ± 0.3; n = 3–7). Conversely, the average duration of individual fusion events was similar in both PiC mutant fibroblasts and Ctrl (Fig. 7D). Furthermore, the distribution of different fusion types in terms of rapid reversal and orientations (Fig. 7E) did not show any alterations in PiC mutant fibroblasts as compared with the Ctrl. In HeLa cells with ~60% decrease in PiC (attained by two different siRNA duplexes, kd1 and kd2), the mitochondrial connectivity and fusion activity were similarly decreased as in the mutant fibroblasts (Fig. 7, F–I).

ATP produced by oxphos is required to maintain the fusion promoting activity of OPA1, the inner mitochondrial membrane fusion protein (18, 19). To test whether the decreased mitochondrial fusion activity can be attributed to a change in OPA1 or another mitochondrial fusion protein, immunoblotting was performed in whole cell lysates. The total abundance of OPA1 was unaltered in PiC mutant fibroblasts (Fig. 8A). A shift in OPA1 immunoreactivity from the high to the lower molecular weight bands indicates a decrease in the fusion-competent forms (as induced by FCCP; Fig. 8B). However, no change in the relative abundance of the high and low molecular weight bands appeared in the PiC mutant fibroblasts (Fig. 8B). Among the main outer membrane fusion proteins, mitofusin 1 (MFN1) was slightly but significantly decreased, whereas mitofusin 2 (MFN2) and the main fission protein DRP1 were unchanged (Fig. 8A). In HeLa cells with 60% PiC depletion, the abundance of none of these fusion and fission proteins was altered (Fig. 8C). Collectively, the results in two different paradigms of PiC depletion show suppressed mitochondrial fusion activity. However, the decreased fusion activity cannot be attributed to processing of OPA1 to fusion-incompetent forms or to substantial loss of another fusion protein.
tion of PiC can become pathogenic when demand for mitochondrial ATP is high. Finally, PiC mutant fibroblasts, as well as HeLa cells with acute PiC loss, had a more fragmented mitochondrial network and slower rate of inner mitochondrial membrane fusion, raising the possibility that organ pathology might be contributed by dysfunctional mitochondrial processes other than, or in addition to, suppressed oxphos.

The Δmir1 yeast strain transformed with one or both of the newly identified human PiC variants was used to evaluate the functional relevance of these variants. Leu-200 is conserved among mammalian species and in yeast and is located within a region that is fully to strongly conserved between human and yeast. Based on the model of PiC determined by Bhoj et al. (13), Leu-200 resides in a small matrix helix that connects trans-

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membrane helices 3 and 4 and interacts with adjacent hydrophobic residues near PX(D/E)XX(K/R), the consensus sequence of all inner mitochondrial membrane carrier proteins. It was predicted that substitution of leucine for tryptophan would not be well tolerated due to tryptophan’s large hydrophobic side chain, which would be exposed to the solvent and could introduce steric clashes (13). However, the L200W substitution did not prevent growth on mitochondrial substrate and could introduce steric clashes (13). Nevertheless, the L200W substitution did not prevent growth on mitochondrial substrate even at elevated temperature, suggesting that this variant is neutral. In contrast, the GSSAS substitution did not prevent growth only on mitochondrial substrate even in unchallenged yeast and despite only slightly decreased PiC protein abundance. This is in line with a loss-of-function mutation and a deleterious effect on oxphos. Based on the nomenclature of the ADP/ATP carrier, GSSAS occurs at matrix loop-connecting transmembrane helices 5 and 6. The variant replaces very small (Gly and Ala), flexible (Gly), and polar (Ser) residues for bulky (Gln), hydrophobic (Ile), and inflexible (Pro) residues. Thus, GSSAS → QIP was predicted to not be well tolerated (13), which is what we found here. Because previously confirmed pathological SLC25A3 variants affected only PiC-A (11, 12), GSSAS → QIP is the first variant that affects both isoforms.

PiC mutant fibroblasts with ~60% less PiC protein had lower basal JO2, maximal leak-dependent JO2, and maximal electron transport chain activity (Fig. 6C) as compared with fibroblasts from four healthy donors. Rather, the decrease in not only basal but also leak-dependent JO2 and maximal electron transport chain activity in intact mutant fibroblasts is consistent with lower provisioning of reducing equivalents to the electron transport chain. The pronounced decrease in JO2 in intact mutant fibroblasts contrasts with minimal decrease in intact HeLa cells with greater PiC depletion. This different response to PiC depletion may arise from long versus short term PiC loss. Another possibility is that mutant PiC protein in fibroblasts induced a stress response that might not occur in HeLa cells that were depleted of PiC through suppression of mRNA rather than mutation. In fact, lower PiC protein was not predicted by either of the mutations which are in-frame; indeed, mRNA expression of both PiC isoforms was unaffected. Lower PiC abundance in the mutation-harboring fibroblasts is also unlikely to reflect natural variability in translation efficiency or turnover because these PiC variant cells deviated similarly in their PiC abundance as compared with fibroblasts from four healthy donors. Rather, depletion of PiC protein in the mutant fibroblasts may reflect an inability of PiC to be properly translated or a tendency for mutant PiC to aggregate, to be misfolded, or to be incorrectly incorporated into complexes (which likely include dimers as well as other components such as the adenine nucleotide translocase (20–22)). Such disruptions in PiC protein folding might induce a mild stress response that is partially responsible for the mitochondrial phenotypes observed in mutant fibroblasts.

The substantial oxphos remaining in permeabilized mutant fibroblasts and PiC-depleted HeLa cells suggests that the abundance of assembled respiratory complexes and ATP synthase as well as their ability to assemble into higher order complexes was unchanged in mutant fibroblasts. Rather, the decrease in not only basal but also leak-dependent JO2 and maximal electron transport chain activity in intact mutant fibroblasts is consistent with lower provisioning of reducing equivalents to the electron transport chain.
are also suggested by observations made in mice with either 
~60 or >95% depletion of PiC in the heart (8, 9). Cardiac PiC 
depletion of ~60% was associated with very mild cardiac hyper-
trophy and no discernible functional defect (9). Even when PiC 
was depleted by >95%, cardiac function was unaffected for at 
least 2 weeks, and ATP in the mitochondrial fraction was 50% of 
normal (8), suggesting compensation at the level of mitochon-
dria or that very little PiC can sustain some level of oxphos. 
Using a fresh skeletal muscle biopsy from an individual with a 
hozygous c.215G→A mutation affecting only PiC-A, sub-
strate oxidation was tested using a variety of conditions and was 
clearly suppressed (11). However, the extent of suppression 
varied considerably, ranging from as low as ~10% of control 
(using the lower extent of the control range) to >30% of con-
trol. Because uncoupler-driven oxidation was within the con-
tral range, the variability in substrate oxidation would not be 
due to differences in the capacity to supply reducing equiva-
lents to the electron transport chain or in the capacity of the 
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lents (using the lower extent of the control range) to 
>30% of control (9). This shift can be the result of the observed decrease in mitochondrial fusion activity. A sensible mechanism to underlie decreased fusion could be a decrease in fusion–competent OPA1 that is 
dependent on oxphos-derived ATP (18, 19). However, neither total OPA1 nor the relative contribution of the different forms 
was altered in mutant fibroblasts or in the PiC-depleted HeLa 
cells. A decrease in the amount of the outer membrane fusion 
mediator MFN1 or MFN2 could also cause suppression of 
mitochondrial fusion (17, 24). Indeed, there was a lower level of 
MFN1 in the mutant fibroblasts, but it was unaltered in the 
PiC-deficient HeLa cells that displayed a similar mitochondrial fusion phenotype to the fibroblasts. Furthermore, MFN2 was 
unchanged in both PiC-deficient paradigms. Activity of the fusion 
proteins can also be affected by posttranslational modifications, 
but no specific mechanism appeared likely here and because of 
that was not tested. Although the specific molecular mechanism 
linking PiC activity to fusion requires further investigation, the 
recent findings might illuminate a common cause of mitochon-
drial fragmentation observed in fixed skin fibroblasts derived from 
patients with various oxphos impairments with a general trend 
that fragmentation increases with severity of the defect (25–28). 
Based on the present results, live cell mitochondrial imaging in 
single cells might detect even a mild oxphos (or mitochondrial) 
defect that remains unnoticeable by JO2 measurements. 

The patient harboring the compound heterozygous PiC 
mutation had severe hypertrophic cardiomyopathy that 
required heart transplantation but did not have skeletal myop-
athy, lactic acidosis, or other obvious organ dysfunction (13). In 
contrast, not only hypertrophic cardiomyopathy but also skeletal 
muscle hypotonia and lactic acidosis were symptoms in 
individuals with homozygous mutations leading to loss of 
PiC-A protein in muscle only (11, 12). PiC-B would still be 
expressed; however, there did not appear to be compensation, 
and the levels appeared low (12). Parenthetically, it is also not 
known to what extent PiC-B can functionally compensate for 
PiC-A. The compound heterozygous mutation affects both 
PiC-A and PiC-B, and we have shown here that the GSSAS → 
QIP variant is deleterious and that the variant(s) can destabilize 
PiC protein. However, our data also indicate that some PiC 
protein remains. Furthermore, we show that some oxphos 
capacity is retained even when PiC protein abundance is quite 
low. The latter is also supported by mouse models of PiC deple-
tion in the heart (8, 9). However, we also show that oxphos 
capacity is sensitive to PiC abundance. We propose that these 
findings are consistent with confinement of the phenotype of 
the compound heterozygous mutation to cardiomyopathy 
because of the incessantly high demand for ATP of the heart. 

Our study has revealed that PiC depletion is accompanied 
by defective cell proliferation and mitochondrial fusion and an 
oxphos defect that does not depend on Pi, limitation. Thus, the 
pathogenesis associated with PiC depletion or loss of function 
may, at least in part, reflect these deficiencies. It will also be of 
interest to determine whether these deficiencies are a general 
feature of pathological PiC variants.

Experimental Procedures

Molecular Biology—Yeast MIR1 gene fragment, with 470-bp 
5′-UTR and 50-bp 3′-UTR, was generated by PCR using 
GA74-1A genomic DNA as template and cloned into pRS315. 
The PIIC1 mutants, L200W and QIP (replacing amino acid residues 
247–258 of yeast Pic1p) as well as a humanized version 
(normal human amino acids GSSAS in place of residues 247– 
258 of yeast Pic1p) were created by overlap extension (29) 
and subcloned into pRS315. For the co-expression studies, PICQIP 
was subcloned into pRS316. Construct verification was per-
formed by restriction digest analysis and/or sequencing.

Yeast Strains and Growth Conditions—All yeast strains used 
were derived from GA74-1A (MATa, his3–11,15, leu2, ura3, 
trp1, ade8 (rho″, mit″)). The open reading frame of MIR1 was 
replaced with HIS3MX6 by a PCR-based disruption method. 
The resulting strain was then transformed with pRS316Pic1 
to prevent loss of mtDNA, and the desired strains were gener-
ated by plasmid shuffling. In brief, pRS315, pRS315Pic1, 
and/or pRS315PicQIP were transformed into Δmir1Δ (pRS316Pic1) 
followed by two selections in 5-fluoroorotic acid to generate clones lacking pRS316Pic1. 
Growth studies were performed to evaluate yeast strain func-
tionality. Cultures were inoculated in synthetic complete 
dextrose without leucine (0.17% yeast nitrogen base, 0.5% ammoni-
um sulfate, 0.2% synthetic dropout mixture minus leucine, 2% 
dextrose), grown overnight at either 30 or 37°C, spotted in 
synthetic complete medium without leucine and supplemented 
with dextrose or 3% glycerol and 1% ethanol, and incubated at 
the indicated temperature and duration.
To assess whether the PIC<sup>21P</sup> allele exerts a dominant negative effect on WT PIC1 or the other PIC alleles, pRS316 or pRS316PIC<sup>21P</sup> was transformed into Δmir1 (pRS315), Δmir1 (pRS315PIC1), Δmir1 (pRS315PIC<sup>2100W</sup>), or Δmir1 (pRS315PIC<sup>Hums</sup>). Growth analyses were done as described above except that synthetic complete dextrose without leucine and uracil and the same selection medium supplemented with either dextrose or 3% glycerol and 1% ethanol were utilized for inoculation and spotting, respectively.

**Mammalian Cell Culture**—Primary skin fibroblasts were obtained from the PIC mutant (13) as well as two age-matched controls and two controls from older individuals, resulting in fibroblasts from four healthy individuals that were used as controls. Fibroblasts were cultured in DMEM containing 4 mM glutamine and either 5 mM glucose or 25 mM glucose. For some growth curve experiments, cells were cultured in glucose-free DMEM containing 4 mM glutamine and supplemented with 5 mM β-hydroxybutyrate; this medium forced cells to rely on mitochondria for ATP production (30, 31). PIC mutant and Ctrl cells were only used for 10 passages and were always compared at the same passage number. HeLa cells (ATCC) were cultured in DMEM containing 25 mM glucose, 4 mM glutamine, and 1 mM pyruvate; HeLa cells were not used beyond passage 14. All cells were maintained in a 5% CO<sub>2</sub>, 21% O<sub>2</sub> incubator at 37 °C.

**Transfections with siRNA and Plasmid DNA**—Before silencing, HeLa cells were plated in antibiotic-free medium, and the samples were silenced in serum-free culture medium with PIC-specific (Invitrogen and OriGene; two different siRNA duplexes) or, as a negative control, scrambled siRNAs (100 nM) (1–5). The silencing was validated with Western blotting. For evaluation was confirmed by Western blotting.

**Growth Curves**—Cell growth curves were performed in fibroblasts and HeLa cells. For all conditions, cells were counted in duplicate wells for each time point. Cell counting was performed using a hemocytometer. Each growth curve was performed in cells from independent passages. For HeLa cells, growth curves were started 72 h after treatment with siRNA; continued knockdown for another 48 h was confirmed by Western blotting.

**Mitochondrial Bioenergetics**—JO<sub>2</sub> was measured in adherent cells using the Seahorse XF24. Fibroblasts or HeLa cells were seeded at 40,000/well the day before the experiment. For experiments in intact cells, cells were washed two times in bicarbonate-free DMEM containing the amount of glucose in which the cells had been cultured (i.e. 5 or 25 mM) without or with 4 mM glutamine, pH 7.4 at 37 °C, and then 650 μl of that medium/well was added as the final assay medium. Injections of oligomycin (0.5 μg/ml final concentration), FCCP (see figure legend for final concentrations), and antimycin (1 μM final concentration) were prepared in the respective assay media. To equilibrate cells and degas the culture plate, cells were preincubated at 37 °C in environmental CO<sub>2</sub> for 45 min.

For experiments in permeabilized cells, cells were washed and preincubated as for intact cells. After the 45-min preincubation, cells were washed again with mitochondrial assay solution (MAS) (32) (70 mM sucrose, 220 mM mannitol, 10 mM KH<sub>2</sub>PO<sub>4</sub>, 5 mM MgCl<sub>2</sub>, 2 mM HEPES, 1 mM EGTA and 0.2% (w/v) BSA, pH 7.2 at 37 °C), and then 450 μl/well MAS containing 10 mM pyruvate, 2 mM malate, 4 mM ADP (from a 1 M stock that had been adjusted to pH 7.2), and permeabilizing reagent (1 mM after optimization; Seahorse Biosciences) were added as the final assay medium. The final concentrations of ADP and substrates are expected to be saturating based on experiments in 5–10 μg of isolated mitochondria (32, 33). Oligomycin (2.5 μg/ml final) and FCCP (4 μM final) were injected sequentially. Some wells only received a final injection of oligomycin + FCCP to record phosphorylating JO<sub>2</sub> for a longer time to monitor the stability of JO<sub>2</sub>; after an increase in JO<sub>2</sub> over the first ~5 min of the experiment, JO<sub>2</sub> remained stable for at least 10 min prior to adding oligomycin and uncoupler. The mix-wait-measure protocol used was that recommended for isolated mitochondria (32) that has worked well in our hands (33). To verify that cells were permeabilized, in some experiments, a subset of wells contained succinate (5 mM) and rotenone (1 μM) instead of pyruvate and malate.

For both intact cell and permeabilized cell experiments, JO<sub>2</sub> values were normalized to total cellular protein. Plates used for both types of experiment were always plated at the same time, allowing us to use the protein amount obtained from the intact cell experiment for the permeabilized cell experiment (measurement of protein from the latter was confounded by the presence of BSA in the MAS).

**Mitochondrial Fusion Dynamics**—Imaging measurements were performed as described previously (16, 17). Briefly, cells were incubated in a 0.25% BSA-containing extracellular medium consisting of 121 mM NaCl, 5 mM NaHCO<sub>3</sub>, 4.7 mM KCl, 1.2 mM KH<sub>2</sub>PO<sub>4</sub>, 1.2 mM MgSO<sub>4</sub>, 2 mM CaCl<sub>2</sub>, 10 mM glucose, and 10 mM Na-Hepes, pH 7.4, at 37 °C. Recordings of mtPA-GFP and mtDsRed (512 × 512 pixels) were performed using 488 and 561 nm laser lines at 0.25 s<sup>−1</sup> data acquisition frequency using an LSM780 microscope and 63×/1.4 numerical aperture Apoplan oil objective. Experiments were run at 37 °C. To photoactivate PA-GFP (and photobleach Ds-Red) in 2P mode in 5 × 5-μm areas, a pulsed laser system (760 nm; Chameleon, Coherent, Inc.) was applied. Image analysis was done using either Spectralyzer (custom designed) or Zen2010 (Carl Zeiss).

Mitochondrial dynamics was evaluated by two different approaches. Spread of PA-GFP from the area of photoinactivation was evaluated by masking the 5×5-μm areas and quantifying the time-dependent decay in the fluorescence ratio of PA-GFP and DsRed. The ratio values in each region were normalized to the peak reached during 2P excitation. Fusion events were also counted manually (34). Image analysis was performed in Spectralyzer or Zen2010 imaging software.

**Immunoblotting**—For yeast studies, protein extraction from yeast cells, SDS-PAGE, and immunoblotting were carried out as described (35, 36). The antibodies used for the yeast studies have been described previously (37–39). Images were captured using the Fluorchem Q system (Cell Biosciences, Inc.), and relative protein levels were quantified using ImageJ. Statistical analyses were performed with StatPlus for mammalian cell
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studies, cells were lysed in radioimmune precipitation assay buffer, and then lysates were clarified and assayed for protein content using the bicinchoninic acid assay. Immunoblotting was carried out as described previously (33, 14). The following antibodies were used: SLC25A3 (1:500; Sigma SAB1400208; rabbit), complex I 37-kDa subunit (1:1000; Abcam MS111; mouse), complex II subunit A (succinate dehydrogenase subunit A; 1:1000; Abcam MS204; mouse), complex III core 2 protein (1:1000; Abcam MS304; mouse), MFN1 (1:1000; rabbit polyclonal provided by Richard Youle, National Institutes of Health, Bethesda, MD), MFN2 (1:1000; mouse monoclonal generated against MFN2(710–757)-GST and N-terminal peptide CNSIVTVKKNRIIM-OC provided by Heidi McBride, McGill University, Montreal, Canada), OPA1 (1:1000; BD Biosciences 611738), and GAPDH (1:1000; rabbit), complex I 37-kDa subunit (1:1000; Abcam MS111; mouse), prohibitin (1:1000; Abcam 28172; rabbit), α-tubulin (1:1000; Cell Signaling Technology 2125; rabbit), cytochrome c (1:1000; BD Pharmingen 556433), and GAPDH (1:10000; Advanced Immunochemicals 8-GAPDH-r; rabbit). Preliminary experiments, not shown, using different amounts of fibroblast and HeLa cell lysate were used to determine the amount of protein that provided a signal that was within the linear range of detection. In addition, proteins that were used as loading controls were chosen because they did not vary with PiC depletion.

Quantitative PCR—Total RNA was extracted from cells using Trizol® (Invitrogen). Purified RNA was treated with RQ1 DNase (Promega, Madison, WI) for 30 min at 37 °C. Total RNA concentration was measured with a Qubit® fluorometer (Invitrogen). RNA was reverse transcribed using oligo(dT)20 primers and SuperScript III (Invitrogen). Quantitative PCRs were done using iTaq SYBR Green Supermix with ROX™ Reference Dye (Bio-Rad) in 20-μL reactions (20 ng of cDNA/reaction) using an Eppendorf Mastercycler® ep realplex. Primers were designed using Eurofins Primer Design Tool: isoform A: forward primer, 5’-CGCCGTTGGAAGACGAGC-3’; reverse primer, 5’-CAAATGGCTATGATGTGGTGC-3’; isoform B: forward primer, 5’-CCGTGGAAAGATCAGTGTTG-3’; reverse primer, 5’-AGCTGTTGTGTAGACC-3’. Custom oligos were purchased from Eurofins MGW Operon (Huntsville, AL).

Statistics—The data are shown as the mean ± S.E. of cells recorded for at least three independent cultures unless it is specified differently. Significance of differences was determined by unpaired Student’s t test or one- or two-way analysis of variance with Tukey post hoc tests as appropriate (see figure legends). p < 0.05 was taken as significant.

Author Contributions—E. L. S. and G. H. developed the project and designed the experiments in mammalian cells. S. M. C. designed experiments in yeast. A. G. conducted the mitochondrial morphology and fusion dynamics studies and performed related Western blotting and analyzed the corresponding results. M. G. A. conducted and analyzed experiments in yeast models. Q. L., L. A.-P., T. G., C. M., and E. L. S. conducted bioenergetics and cell proliferation experiments in mammalian cells, performed Western blotting of mammalian cell samples, and analyzed the corresponding results. N. S. procured and provided the human fibroblasts. E. L. S. and G. H. wrote the paper.

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References

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Natural and Induced Mitochondrial Phosphate Carrier Loss: DIFFERENTIAL DEPENDENCE OF MITOCHONDRIAL METABOLISM AND DYNAMICS AND CELL SURVIVAL ON THE EXTENT OF DEPLETION

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