Lack of Cytochrome c in Mouse Fibroblasts Disrupts Assembly/Stability of Respiratory Complexes I and IV**

Received for publication, August 1, 2008, and in revised form, December 1, 2008 Published, JBC Papers in Press, December 15, 2008, DOI 10.1074/jbc.M805972200

Uma D. Vempati1, Xianlin Han2, and Carlos T. Moraes*§¶1

From the 1Department of Neurology and 4Department of Cell Biology and Anatomy, University of Miami Miller School of Medicine, Miami, Florida 33136 and the 4Division of Bioorganic Chemistry and Molecular Pharmacology, Department of Internal Medicine, Washington University, St. Louis, Missouri 63110

Cytochrome c (cyt c) is a heme-containing protein that participates in electron transport in the respiratory chain and as a signaling molecule in the apoptotic cascade. Here we addressed the effect of removing mammalian cyt c on the integrity of the respiratory complexes in mammalian cells. Mitochondria from cyt c knockout mouse cells lacked fully assembled complexes I and IV and had reduced levels of complex III. A redox-deficient mutant of cyt c was unable to rescue the levels of complexes I and IV. We found that cyt c is associated with both complex IV and respiratory supercomplexes, providing a potential mechanism for the requirement for cyt c in the assembly/stability of complex IV.

The mitochondrial electron transport chain consists of four multisubunit complexes, namely, NADH-ubiquinone oxidoreductase (complex I),2 succinate-ubiquinone oxidoreductase (complex II), ubiquinone-cytochrome c oxidoreductase (complex III), and cytochrome c oxide (complex IV, COX). Cytochrome c (cyt c) shuttles electrons from oxidative phosphorylation complex III to complex IV. Electrons are transferred from reduced cyt c sequentially to the CuA site, heme a, heme a3, and CuB binuclear center in the complex IV before being finally transferred to molecular oxygen to generate water (1). Respiratory complexes are assembled into supercomplexes (also called respirasomes). These contain complex I bound to dimeric complex III and a variable copy number of complex IV (2).

In Saccharomyces cerevisiae, cyt c is encoded by two genes: CYC1 and CYC7. Mutagenesis studies in yeast have shown that cyt c is required for the assembly of COX (3, 4). In yeast lacking both the cyt c genes (CYC1 and CYC7), COX assembly was absent. It was also shown that cyt c is only structurally required for COX assembly, because a catalytic mutant of cyt c (W65S) was sufficient to bring about near normal levels of COX. However, because yeast lacks complex I, they could not analyze the role of cyt c in the assembly/stability of complex I. Mammals possess two different isoforms of cyt c encoded on different chromosomes: the somatic (cyt cS)- and testis (cyt cT)-specific isoforms. In mouse, the cDNAs bear 74% homology, whereas the proteins possess 86% identity with most dissimilarity in the C terminus.

Cardiolipin (CL) is an anionic phospholipid present almost exclusively in the mitochondrial membranes and constitutes 25% of its total phospholipids (5). Work from several laboratories showed that CL is essential for the membrane anchorage of the respiratory supercomplexes. CL has two main roles in the mitochondrial structure and function, namely, stabilization of mitochondrial membranes and specific interactions with proteins. CL deficiency results in inefficient energy transformation by oxidative phosphorylation, swelling of mitochondria, decreased ATP/oxygen ratio, and reduced membrane potential (6, 7). In accordance, in S. cerevisiae lacking CL synthase, the supercomplex comprising complexes III and IV is unstable (8). Assembly mutants of COX had significantly reduced CL synthase activity, whereas assembly mutants of respiratory complex III and complex V showed less inhibition (9). Subsequently, the proton gradient across the inner mitochondrial membrane was found to be important for CL formation and that CL synthase was stimulated by alkaline pH at the matrix side (10). In this study, we investigated the role of cyt c depletion on CL levels by examining its content and composition in cyt c null cells.

Here we aimed to answer the following questions: What is the role of cyt c in the assembly and maintenance of the different respiratory complexes in mammals? Are there changes in the content/composition of lipids in the cyt c-ablated cells? Analysis of mouse fibroblasts revealed that cyt c is essential for the assembly/stability of COX, and a catalytically mutant form of cyt c cannot rescue the COX defect in the cyt c null cells. CL and triacylglycerols showed significant differences in the cyt c null cells, both in content and composition.

EXPERIMENTAL PROCEDURES

Genetically Modified Mice and Derived Cell Lines—The crosses performed to obtain mice with the genotype cyt cs−/−.
Lack of cyt c in Mouse Fibroblasts Disrupts Complexes I and IV

cyt ct<sup>−/−</sup> Transgene<sup>lox/lox</sup> and the characterization of lung fibroblasts lacking cyt c were previously described (11).

**Cell Lines**—All cells were grown in Dulbecco’s modified Eagle’s medium with high glucose, supplemented with uridine and pyruvate. Clones L3, L4, and L7 lacked both the somatic and testis isoforms of cyt c and were designated as double knockout (dKO). LF represents the original lung fibroblasts before the deletion of the floxed cyt c. CL1 and CL15 were derived from L3, after the reintroduction of a wild-type cyt c cDNA. CL18 and CL25 were derived from L3 after the reintroduction of a mutant (W60S) cyt c cDNA.

Mouse LM(TK<sup>−</sup>) cells were obtained from ATCC (CCL 1.3), and the mitochondrial DNA (mitochondrial DNA)-less derivative was obtained by ethidium bromide treatment as described previously (12). Somatic cyt c cDNA was subcloned into pRES-puro vector (Clontech, cyt c puro-8) and introduced into cyt c dKO fibroblasts (L3 clone) by stable transfection to generate clones CL1 and CL15. A point mutation (W60S) was introduced into cyt c cDNA by PCR using the QuikChange site-directed mutagenesis kit (Stratagene) on the cyt c puro-8 template using mutant primers (forward: 5’-GCCAACAGAAAGGCGCATCCCCGGAGGAGCATACCTGATGG-3’ and reverse: 5’-CCATCAGGCTATCTCTCCGAGGAGATGATGCCCTTGGTCTTGTTGGC-3’), with the modified nucleotides shown in lowercase. The presence of the mutation was confirmed by sequencing. After stable transfections in L3 (dKO) cells, two clones containing the desired mutation were obtained (CL18 and CL25).

**Immunostaining**—Live cells were stained with MitoTracker red (200 nM, Invitrogen) and subsequently fixed and permeabilized in 4% paraformaldehyde and ice-cold methanol, respectively. Then, cells were incubated with a primary antibody against cyt c (BD Biosciences and Mitosciences) and subsequently with a secondary antibody tagged with fluorescent Alexa-fluor (Molecular Probes) and visualized by confocal microscopy.

**Respiration Measurements**—Cellular respiration was measured by polarography as described (13). The oxygen consumed by cells was measured both before and after addition of the complex III inhibitor, antimycin A, and the subsequent addition of ascorbate and N,N,N',N'-tetramethyl-1,4-phe-nylendiamine dihydrochloride (TMPD).

**Isolation of Mitochondria**—Mitochondria were prepared by the nitrogen cavitation method as described previously (14) with some modifications. In brief, cells were washed in 10 ml of the nitrogen cavitation method as described previously (14) with some modifications. In brief, cells were washed in 10 ml of the nitrogen cavitation method as described previously (14)

**Heme Staining**—250 μg of mitochondria were reduced with dithiothreitol (50 mM) on ice, resolved by SDS-PAGE gels, and blotted onto a PVDF membrane. The heme-associated peroxidase activity was detected on the membrane using SuperSignal chemiluminescence reagent (Pierce) followed by autoradiography, as described (16, 17).

**Electron Microscopy**—Cultured cells were fixed in 2% phosphate-buffered glutaraldehyde (with 100 mM sucrose) overnight at 4 °C, followed by 2% buffered OsO<sub>4</sub> for 1 h at room
temperature. Cultures were further processed for EMbed plastic (Electron Microscopy Sciences) embedding. Areas were chosen for semi-thin sectioning; the 1-/H9262 m sections were stained with toluidine blue/methylene blue/sodium borate. Thin sections (90 nm) were stained with uranyl acetate and lead citrate and examined in a Philips CM-10 electron microscope (FEI Co.). Sections were cut perpendicularly to the plane of the coverslip (18).

**Determination of Reactive Oxygen Species**—Reactive oxygen species were detected in live cells using the cell-permeable probe, dichlorofluorescein (Invitrogen) followed by flow cytometry. Cells were trypsinized and counted, and 0.2 million cells were stained with dichlorofluorescein, either untreated or upon treatment with hydrogen peroxide.

**Mass Spectrometric Analysis of Lipids**—Enhanced shotgun lipidomics analyses were performed as described before (19, 20). In brief, lipids were analyzed on a QqQ mass spectrometer (ThermoFisher Scientific, San Jose, CA) equipped with an electrospray ion source. All electrospray ionization mass spectrometric analysis of lipids were conducted by direct infusion employing a Harvard syringe pump at a flow rate of 4 ml/min. Typically a 1-min period of signal averaging was employed for each mass spectrum, and a 2-min period of signal averaging was employed for each tandem mass spectrometric spectrum. A mass resolution of 0.4 Thomson was employed for acquisition of mass spectra with a QqQ instrument (21). The results are presented as the mean from two different cyt c dKO clones (L3 and L4) and two different cyt c reintroduced clones (CL1 and CL15). The error of the method, after standardization to internal controls, is 10% (22).

**RESULTS**

**cyt c−/− Fibroblasts Are Deficient in Respiration**—Lung fibroblasts (LFs) were derived from the conditional cyt c knockout mice as described (11). The cyt c transgene was deleted in cultured lung fibroblasts by the expression of Cre recombinase from a recombinant adenovirus. Because these fibroblasts lack both the cyt c isoforms, they were designated as double knockout (dKO). cyt c dKO cells were used for the reintroduction of wild-type or mutant cyt c. The expression of cyt c was assessed by both immunostaining and Western blots (Figs. 1 and 3A). Respiration was measured in the dKO and cells with reintroduced cyt c. Respiration in the dKO reintroduced with the wt cyt c was close to the values of control LF cells (Fig. 2A). The mutant cyt c cDNA had a single amino acid changed from tryptophan to serine (W60S), as described in *S. cerevisiae* (3, 23). The lines expressing the mutant form are clones CL18 and CL25. In yeast, the W65S mutant was catalytically inactive in electron transfer. As expected, the cyt c dKO cells with the mutant cyt c cDNA reintroduced did not respire. The cells lacked both endogenous and ascorbate plus TMPD-mediated respiration (Fig. 2A). We previously showed that the cyt c dKO cells maintained a mitochondrial membrane potential that could be dissipated with a protonophore (11), a phenomenon associated with a reverse function of complex V in OXPHOS-deficient cells (24). We could not detect differences in reactive oxygen species produc-

![FIGURE 1. Expression and localization of cyt c in fibroblasts. Immunostaining of control (LF), dKO (L3), WT cyt c reintroduced (CL1) and W60S mutant cyt c reintroduced (CL18) cells with cyt c antibodies. Cells were co-stained with MitoTracker red to confirm the mitochondrial localization.](image-url)
Lack of cyt c in Mouse Fibroblasts Disrupts Complexes I and IV

tion between the different cell lines using dichlorofluorescein followed by flow cytometry analysis (data not shown). Other investigators made similar observations in the cyt c null cells (25).

Activity of Respiratory Complexes in cyt c Null Cells—Respiratory complex activities were measured by spectrometric assays on mitochondria isolated from the cells. We observed that the cyt c dKO fibroblasts were completely deficient in complex IV activity (Fig. 2B), whereas the complex III activity was reduced to 50% of the control levels (Fig. 2D). Citrate synthase activity was measured to normalize the other activities and found to be similar in all samples (Fig. 2C). To verify that the mitochondrial DNA content was reduced in the cyt c-ablated cells, we performed Southern hybridization. We found that mitochondrial DNA levels were the same in the control and cyt c null cells (data not shown).

Steady-state Levels of Respiratory Complexes in cyt c Null Cells—Western blot analyses were performed to assess the steady-state levels of different respiratory complex subunits. Antibodies against Ndufa9, SDH, Core2, COXI, and ATPase-β were used (Fig. 3A). Cyt c dKO fibroblasts had undetectable levels of COXI protein. There were no major changes in subunits of other OXPHOS complexes. When we reintroduced either wild-type or mutant cyt c cDNA (W60S) into the cyt c null fibroblasts, COX I protein was observed (Fig. 3A).

The assembled respiratory complexes were analyzed by BN-PAGE (15) followed by Western analysis. Interestingly, the cyt c dKO cells had undetectable levels of not only complex IV, but also complex I (Fig. 3B and supplemental Fig. S1). Complex III steady-state levels were only mildly reduced in most cells, whereas the relative levels of the complexes II and V were not changed (Fig. 3B and supplemental Fig. S1). Upon reintroduction of cyt c cDNA into the cyt c dKO fibroblasts, we found that fully assembled COX could be restored, and to a lesser extent, also complex I (Fig. 3B).

The W60S mutant clone (CL18), though showing the presence of COX1, lacked detectable levels of both complexes IV and I (Fig. 3B). In-gel activity results were in agreement with the Western blots (supplemental Fig. S2).

Two-dimensional BN-PAGE analysis was performed to address the role of cyt c in the stability of OXPHOS complexes. Digitonin-treated mitochondria from control fibroblasts (LFs) were resolved on a 4–10% BN-PAGE followed by a second dimension SDS-PAGE (12%) and Western analysis. This revealed that a fraction of cyt c was associated with a supercomplex containing complexes I, III, and IV (Fig. 3C). This fraction correlated with the levels of complex IV detected in the respi-ramose. Accordingly, the bulk of the cyt c was associated with complex IV (Fig. 3C).

Newly Synthesized Complex IV Cannot Be Detected in the Absence of cyt c—We wanted to determine whether the lack of complexes I and IV in the cyt c dKO fibroblasts was due to defective synthesis or degradation. Control, dKO, and dKO cells reintroduced with cyt c cDNA were labeled with [S35]methionine and cysteine in the presence of emetine, a cytosolic protein synthesis inhibitor. We observed that all the cell types synthesized COX1, COX2, and COX3 at com-

FIGURE 2. Mouse fibroblasts with disrupted cyt c (somatic and testis) alleles are OXPHOS-deficient. A, control (LF), cyt c dKO (L3, L4, and L7), wild-type cyt c cDNA reintroduced (CL1 and CL15), and mutant cyt c cDNA reintroduced (CL18) intact cells were analyzed for KCN-sensitive oxygen consumption. All cell lines were also analyzed for the ratio of the ascorbate-TMPD respiration (where electrons are donated directly to cyt c) to endogenous respiration (starting at complexes I and II). cyt c dKO cells and those reintroduced with the mutant cyt c cDNA did not respire. B and D, respiratory complexes (III and IV) activities measured by spectrophotometric assays. Citrate synthase activity was measured in all the samples (C) as a measure of mitochondrial levels. cyt c dKO cells and those reintroduced with the W60S mutant cyt c cDNA (CL18 and CL25) lacked complex IV and had reduced complex III activities.
Lack of cyt c in Mouse Fibroblasts Disrupts Complexes I and IV

A Western analysis of mitochondria isolated from the different cell lines using antibodies raised against cyt c, Ndufa9, COX1, SDH, Core2, and ATPase-β. VDAC1 antibody was used as a loading control. cyt c dKO cells (L3 and L4) lacked COX1. BN-PAGE was performed on a 4–13% gel. The proteins were transferred onto a PVDF membrane and probed with antibodies raised against the different complex subunits, namely, Ndufa9 (complex I), Core2 (complex III), COX1 (complex IV), and ATPase-β (complex V). CI, CIII, CIV, and CV are complexes I, III, IV, and V, respectively. cyt c dKO cells (L3) and those reintroduced with the W60S mutant cyt c DNA (L3 and L4) were very reduced in CL1 and CL15. Complex I in the dKO cells with reintroduced cyt c was reintroduced (it was undetectable in the dKO, Fig. 3A). Surprisingly, the levels of complex I in the dKO cells with reintroduced cyt c were still very low. This could be a reflection of the relatively low levels of complex IV in these cells, but the reason for these low levels of complexes IV and I are not clear.

Localization of Mutant cyt c in Mitochondria—Isolated mitochondria and mitoplasts were treated with protease K and subjected to Western analysis. Wild-type and mutant cyt c were detected in both mitochondria and mitoplasts. The wild-type cyt c (CL1 and LF) was resistant to protease K, probably because it was mostly protected within cristae folds (Fig. 5A). However, the mutant cyt c (CL18) was more sensitive to the protease (Fig. 5A, right panel). This increased sensitivity can be due to incorrect folding and/or altered localization in the intermembrane space. Surprisingly, we found that COX I was present at higher levels when the mutant cyt c was reintroduced (it was undetectable in the dKO, Fig. 3A). However, it was also very sensitive to protease K in mitoplasts (Fig. 5A, right panel). This sensitivity has been consistently observed in different experiments. We conclude that a functional cyt c is required for the normal localization of COX I within the inner membrane.

W60S cyt c Has Impaired Heme Addition—We determined the presence of heme groups in cyt c by detecting the peroxidase activity associated with heme. Proteins were resolved on a lithium dodecyl sulfate-PAGE gel and transferred onto a PVDF membrane followed by Western analysis with antibodies against cyt c and complexes I (Ndufa9), III (core), and IV (COX 1).
membrane. The c-type mitochondrial cytochromes were detected on the blot after the addition of a chemiluminescence substrate (Fig. 5B). The signal corresponding to cyt c-heme in W60S mutant was ~30% of the signal in the wild-type cyt c, which is similar to what was reported in the yeast (3).

**cyt c KO Cells Have Abnormal Mitochondria**—As expected from an OXPHOS-deficient cell, cyt dKO fibroblasts showed the presence of enlarged mitochondria, which displayed irregular cristae organization. Abundant lipid droplets (gray) and lipofuscin (darkly stained) were detected in the dKO fibroblasts, but not in the control cells (supplemental Fig. S4). The abnormal lipids, disorganized cristae, and absence of complexes I and IV suggested that the lipid composition of the cyt c null mitochondria was altered.

**cyt c-ablated Cells Have Altered Mitochondrial Lipid Composition**—Shotgun lipidomics analyses on cellular extracts of cyt c null fibroblasts was performed as previously described (19, 20). We demonstrated two marked changes in lipid content in cyt c dKO fibroblasts. First, we found lower CL content in the dKO fibroblasts (Fig. 6). The total CL content in the dKO fibroblasts was 22% lower than in the control. The reduction was mainly present in CL species containing oleic acid (18:1) (supplemental Fig. S5A). Secondly, we found that the total content of triacylglycerols (TAG) in the cyt c null fibroblasts was substantially elevated (over 3-fold), probably reflecting the impaired OXPHOS system (Fig. 6 and supplemental Fig. S5B). The accumulation of TAG content in the dKO fibroblasts was also revealed by the presence of abundant lipid droplets using electron microscopy (supplemental Fig. S4).

After the reintroduction of wt cyt c cDNA, the total amount of CL and TAG, as well as the composition of CL and TAG molecular species were restored to near the levels of control cells (Fig. 6). We also measured the content of lipids in a cell line depleted of mitochondrial DNA (rho zero (ρ0) cells) and its control cell line (LM(TK−)). Mitochondrial DNA deficiency in ρ0 cells resulted in a marked reduction of CL content, and a substantial accumulation of TAG content (Fig. 6). These results suggest that the altered lipid content and/or composition are likely related to the OXPHOS assembly and/or function and not with the specific loss of cyt c.

**DISCUSSION**

The function of cyt c in the assembly of mammalian OXPHOS complexes has not been explored. We found that both the function and the physical presence of cyt c were required for the assembly/stability of respiratory complexes. Pulse-chase experiments revealed that the mitochondrially encoded COX subunits 1, 2, and 3 were reduced in the KO and the mutant cyt c introduced cells as early as 2 h post-synthesis, possibly due to increased turnover of the unassembled subunits. Therefore, our results demonstrate that, in mammals, a functional cyt c is required for the assembly/stability of COX.

---

**FIGURE 4. Complexes I and IV are not synthesized or are extremely unstable in cyt c dKO cells.** Cells were labeled with [35S]methionine and cysteine, followed by a chase for 2, 4, and 24 h, respectively. Lysates from the different cell lines were resolved on an SDS-PAGE (A). Lysates of LF and L3 (dKO) were also analyzed by BN-PAGE (B) and subjected to autoradiography. In C, the same samples shown in B were transferred onto a membrane and probed with antibodies raised against the different complex subunits, namely, Ndufa9 (complex I), COXI (complex IV), and ATPase-β (complex V). VDAC1 and CII(Fp) antibodies were used as loading controls.
In *S. cerevisiae*, cyt c is encoded by two genes: *CYC1*, which encodes Iso-1-cytochrome c that accounts for 95% of the total cyt c in mitochondria, and the homologous and less abundant iso-2-cytochrome c encoded by *CYC7*. Even though iso-2-cytochrome c represents only 5% of the total cyt c, it is sufficient to support respiration and growth, although at a reduced rate on non-fermentable carbon sources. Mutations in both isoforms lead to a respiratory defect (26). Previously, Barrientos et al. (3) reported that yeast lacking both the cyt c alleles, namely, cyc1 and cyc7 failed to assemble COX. The cyc1 null mutant had ~40% of the COX activity of the wild-type cells, even though they had only 12% of the levels of cyt c compared with the control cells. The possibility that cyt c could be required for heme a biosynthesis was excluded (27). It was also shown that cyt c is only structurally required for COX assembly, because a catalytic mutant of cyt c (W65S) was sufficient to bring about near normal levels of COX (3). In contrast, our catalytically inactive mutant was not able to restore complex IV assembly. Work in yeast revealed that COX contains three redox centers: Cox1p contains two redox centers, one formed by heme A, and another by heme a3 and CuB. The third redox center of the enzyme is formed by the two copper ions present in the CuA site of Cox2p. Electrons are transferred from reduced cyt c sequentially to Cox1p, heme a, heme a3, and CuB binuclear center in the complex IV (1). We speculate that the reduction of COX1 by cyt c may be important for its correct insertion in the inner membrane, because COX1 was susceptible to proteinase K digestion when mitoplasts derived from the W60S mutant cyt c-expressing cells were treated with the protease. It is not clear why COX1 was more stable in the mutant than in the null cell line, but the physical presence of cyt c could have a stabilizing effect on unassembled COX1. In addition, based on our observations that cyt c is a part of respirasomes and is associated with complex IV, and the lack of cyt c results in the loss of this complex, we hypothesize that cyt c is required for the stability of the complex IV. Recently it was also reported by others that cyt c is a component of the respirasome (38).

Previous results from our group (15) and others (28) have shown that complex I is not stable in the absence of complex IV. Similar results were observed when complex III was absent (29, 30). Complexes I, III, and IV exist together as supramolecular assemblies called respirasomes. The assembly of a respirasome might have advantages in substrate channeling of quinones and/or cyt c, sequestration of reactive intermediates, and stabilization of individual complexes (2). It is likely that the lack of complex I is related to the lack of complex IV, as previously described (15). However, our results showed that the requirement for complex I assembly/stability is not of a complex IV subunit, but rather the presence of an intact complex IV. In addition, our findings indicate that respirasome formation depends...
not only on OXPHOS complexes, but also on single molecular entities, such as cyt c.

Results from our experiments designed to probe protein localization in mitochondria and heme-association to cyt c revealed indirect evidence that the W60S mutant cyt c is misfolded. Englander’s group did extensive work on cyt c folding using the two-dimensional NMR technique (31, 32). They reported that cyt c is composed of five cooperatively folding units, called foldons, that continually unfold and refold even under native conditions. Foldons are generally coincident with secondary structural elements. Based on the cyt c structure (33), the tryptophan residue at position 60, which we mutated (Trp-60), lies in the C-terminal α-helix. During cyt c folding, the C-terminal α-helix together with the N-terminal α-helix forms the first foldon in a sequential unfolding-refolding pathway (31). Hence, introducing the W60S mutation might impair the subsequent steps in the cyt c folding pathway.

Import of apocytochrome c into the mitochondrial intermembrane space is distinct from that of proteins containing a presequence. Using proteoliposomes, apocytochrome c was shown to require the protease-resistant part of the TOM complex (translocase in outer membrane), involving TOM40 and cytochrome c heme lyase (CCHL) (34). cyt c import does not require ATP hydrolysis and is driven by the interaction of cyt c with its “trans-side receptor” CCHL (35, 36). The W60S apocytochrome c-CCHL interaction might be weak, instead of the normal strong and stable binding. Hence, the covalent attachment of heme by CCHL was found to occur only in a low percentage of the mutant cyt c apoprotein. The heme addition renders cyt c translocation process irreversible, which is followed by folding to the native conformation and dissociation from CCHL. In addition, the cyt c structure reveals that tryptophan at position 60 is one of the residues that interact with the heme, mutation of which could result in improper folding.

CL is an anionic phospholipid present exclusively in the mitochondrial membrane and constitutes 25% of its total phospholipids (5). Work from several laboratories showed that CL is not only essential for membrane anchorage of the respiratory supercomplexes but also possesses multiple other roles in the mitochondrial structure and function, including stabilization of mitochondrial membranes, contribution to the very dynamic mitochondrial fusion and fission, and facilitating specific interactions with mitochondrial membrane proteins. In S. cerevisiae lacking cardiolipin synthase, the supercomplex comprising complexes III and IV was unstable (8). Additionally, assembly mutants of complex IV have significantly reduced CL synthase activity, whereas assembly mutants of respiratory complex III and complex V are less inhibitory (9). Subsequently, the proton gradient across the inner mitochondrial membrane was shown to be important for CL formation, and the authors conclude that CL synthase is stimulated by alkaline pH at the matrix side (10).

In the current study, we found both cyt c ablation and mitochondrial DNA depletion lead to a deficiency in CL content. Because the proton gradient is important for CL synthesis, the disruption of the OXPHOS respiration chain, or complex IV, could be sufficient to explain our findings of CL deficiency in the cyt c dKO fibroblasts and ρ0 cells. Although not a specific effect, the changes in cardiolipin may also affect complex and supercomplex assembly/stability.

In conclusion, cyt c is essential for the assembly/stability of complex IV in mammals. In the absence of cyt c, complex IV cannot assemble or is extremely unstable. We also showed that the W60S cyt c mutant cannot restore COX assembly, even though it can somehow stabilize COX1, enough to protect it from degradation from endogenous proteases. The presence of cyt c in association with complex IV and the respirasome suggests a structural function for this small protein. In summary, our findings show that, in mammalian cells, not only OXPHOS multisubunit complexes are interdependent, but also small single polypeptide carriers, such as cyt c.

Acknowledgments—We are grateful to Dayami Hernandez for technical assistance. We are also grateful to the input from Dr. Antoni Barrientos during the course of this work.

REFERENCES

1. Moraes, C. T., Diaz, F., and Barrientos, A. (2004) Biochim. Biophys. Acta 1659, 153–159
2. Schagger, H., and Pfeiffer, K. (2000) EMBO J. 19, 1777–1783
3. Barrientos, A., Pierre, D., Lee, J., and Tzagoloff, A. (2003) J. Biol. Chem. 278, 8881–8887
4. Pearce, D. A., and Sherman, F. (2003) J. Biol. Chem. 278, 20879–20882
5. Robinson, B. H. (2000) Pediatr. Res. 48, 581–585
6. Koshkin, V., and Greenberg, M. L. (2000) Biochem. J. 347, 687–691
7. Zhao, M., Slachevsky, A. M., and Greenberg, M. L. (1998) J. Biol. Chem. 273, 2402–2408
8. Koshkin, V., and Greenberg, M. L. (2000) Biochem. J. 347, 687–691
9. Zhao, M., Slachevsky, A. M., and Greenberg, M. L. (1998) J. Biol. Chem. 273, 2402–2408
10. Gohil, V. M., Hayes, P., Matsuyama, S., Schagger, H., Slachevsky, A. M., and Greenberg, M. L. (2004) J. Biol. Chem. 279, 42612–42618
11. Vempati, U. D., Diaz, F., Barrientos, A., Narisawa, S., Mian, A., Millan, J. L., Boise, L. H., and Moraes, C. T. (2007) Mol. Cell Cell Biol. 27, 1771–1783
12. Dey, R., and Moraes, C. T. (2000) J. Biol. Chem. 275, 7087–7094
13. Barrientos, A., Kenyon, L., and Moraes, C. T. (1998) J. Biol. Chem. 273, 14210–14217
14. Gottlieb, R. A., and Adachi, S. (2000) Methods Enzymol. 322, 213–221
15. Dey, R., Moraes, C. T. (2000) Mol. Cell Cell Biol. 26, 4872–4881
16. Bernard, D. G., Gabbily, S. T., Dujardin, G., Merchant, S., and Hamel, P. P. (2003) J. Biol. Chem. 278, 49732–49742
17. Dutta, C., and Henry, H. L. (1990) Anal. Biochem. 184, 96–99
18. Plant, G. W., Currier, P. F., Cuervo, E. P., Bates, M. L., Pressman, Y., Bunge, M. B., and Wood, P. M. (2002) J. Neurosci. 22, 6083–6091
19. Han, X., and Gross, R. W. (2005) Mass Spectrom. Rev. 24, 367–412
20. Han, X., Yang, Y., Yang, J., Cheng, H., and Gross, R. W. (2006) J. Lipid Res. 47, 864–879
21. Mancuso, D. J., Sims, H. F., Han, X., Jenkins, C. M., Gu, S. P., Yang, K., Moon, S. H., Pietka, T., Abumrad, N. A., Schlesinger, P. H., and Gross, R. W. (2007) J. Biol. Chem. 282, 34611–34622
22. Cheng, H., Mancuso, D. J., Jiang, X., Gu, S., Yang, J., Yang, K., Sun, G., Gross, R. W., and Han, X. (2008) Biochemistry 47, 5869–5880
23. Schweingruber, M. E., Stewert, J. W., and Sherman, F. (1979) J. Biol. Chem. 254, 4132–4143
24. Buchet, K., and Godinot, C. (1998) J. Biol. Chem. 273, 22983–22989
25. Mansfield, K. D., Guzy, R. D., Pan, Y., Young, R. M., Cash, T. P., Schumacker, P. T., and Simon, M. C. (2005) Cell Metab. 1, 393–399
26. Downie, J. A., Stewart, J. W., Brockman, N., Schweingruber, A. M., and Sherman, F. (1977) J. Mol. Biol. 113, 369–384
27. Barros, M. H., and Tzagoloff, A. (2002) FEBS Lett. 516, 119–123
Lack of cyt c in Mouse Fibroblasts Disrupts Complexes I and IV

28. Li, Y., D’Aurelio, M., Deng, J. H., Park, J. S., Manfredi, G., Hu, P., Lu, J., and Bai, Y. (2007) J. Biol. Chem. 282, 17557–17562
29. Acin-Perez, R., Bayona-Bafaluy, M. P., Fernandez-Silva, P., Moreno-Loshuertos, R., Perez-Martos, A., Bruno, C., Moraes, C. T., and Enriquez, J. A. (2004) Mol. Cell 13, 805–815
30. Schagger, H., de Coo, R., Bauer, M. F., Hofmann, S., Godinot, C., and Brandt, U. (2004) J. Biol. Chem. 279, 36349–36353
31. Krishna, M. M., Maity, H., Rumbley, J. N., Lin, Y., and Englander, S. W. (2006) J. Mol. Biol. 359, 1410–1419
32. Maity, H., Maity, M., and Englander, S. W. (2004) J. Mol. Biol. 343, 223–233
33. Rumbley, J., Hoang, L., Mayne, L., and Englander, S. W. (2001) Proc. Natl. Acad. Sci. U. S. A. 98, 105–112
34. Diekert, K., de Kroon, A. I., Ahting, U., Niggemeyer, B., Neupert, W., de Kruijff, B., and Lill, R. (2001) EMBO J. 20, 5626–5635
35. Dumont, M. E., Cardillo, T. S., Hayes, M. K., and Sherman, F. (1991) Mol. Cell Biol. 11, 5487–5496
36. Mayer, A., Driessen, A., Neupert, W., and Lill, R. (1995) Methods Enzymol. 260, 252–263
37. Deleted in proof
38. Acin-Perez, R., Fernandez-Silva, P., Peleato, M. L., Perez-Martos, A., and Enriquez, J. A. (2008) Mol. Cell 32, 529–539