REVIEW

Transcriptional regulatory circuits controlling mitochondrial biogenesis and function

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We are witnessing a period of renewed interest in the biology of the mitochondrion. The mitochondrion serves a critical function in the maintenance of cellular energy stores, thermogenesis, and apoptosis. Moreover, alterations in mitochondrial function contribute to several inherited and acquired human diseases and the aging process. This review summarizes our understanding of the transcriptional regulatory mechanisms involved in the biogenesis and energy metabolic function of mitochondria in higher organisms.

The mitochondrial genome

A defining feature of eukaryotic cells is that they contain nuclear and mitochondrial genomes sequestered into distinct subcellular compartments. The mitochondrial genetic system is comprised of a circular DNA genome (mtDNA, ~16.5 kb in vertebrates; Fig. 1), the enzymes required for its transcription and replication, and the protein synthetic machinery necessary for the translation of 13 mitochondrial mRNAs (for review, see Garesse and Vallejo 2001). These mRNAs, which account for the entire protein-coding capacity of mtDNA, encode essential subunits of respiratory complexes I, III, IV, and V. The extrusion of protons through complexes I, III, and IV is coupled to the sequential transfer of electrons to a series of carriers of increasing redox potential resulting in an electrochemical proton gradient across the inner membrane. Complex V, comprised of an ATPase coupled to an inner membrane proton channel, can dissipate the proton gradient in the synthesis of ATP or can couple proton pumping to ATP hydrolysis to maintain the gradient. mtDNA also encodes for two ribosomal and 22 transfer RNAs, required for translation by mitoribosomes within the matrix.

The limited coding capacity of mtDNA necessitates that nuclear genes make a major contribution to mitochondrial metabolic systems and molecular architecture (Garesse and Vallejo 2001). One major class of nuclear genes contributes catalytic and auxiliary proteins to the mitochondrial enzyme systems. For example, the majority of the 100 or so subunits of the respiratory apparatus are nucleus-encoded. In addition, nucleus-encoded metabolic enzymes necessary for the oxidation of pyruvate, fatty acids (β -oxidation cycle), and acetyl-CoA (tricarboxylic acid cycle), the biosynthesis of certain amino acids, and the manufacture of heme, among others, are localized to the mitochondrion. A second class of nuclear genes encodes protein import and assembly factors. A third class contributes key proteins that are required for the replication and expression of the mitochondrial genome including nucleic acid polymerases, RNA processing enzymes, transcription and replication factors as well as tRNA-synthetases, translation factors, and ribosomal subunits. Thus, the program regulating mitochondrial biogenesis involves the coordinate actions of nuclear and mitochondrial genes.

Regulatory proteins involved in mitochondrial gene transcription

In yeast, mtDNA transcription is initiated at ~20 transcriptional units throughout the genome (for review, see Poyton and McEwen 1996). In vertebrates, transcription is initiated bidirectionally at two promoters, P_H and P_L for heavy (H) and light strands (L), respectively, within the D-loop regulatory region (Shadel and Clayton 1997; Clayton 2000). The D-loop is the longest noncoding region in vertebrate mtDNA and contains, in addition to P_H and P_L , the H-strand replication origin (O_H ; Fig. 1). In the "strand asymmetric model" of mtDNA replication, the RNA transcript initiated at P_L is cleaved in the vicinity of three evolutionarily conserved sequence blocks (CSB I, II, and III), and H-strand replication is initiated at the sites of these cleavages (Bogenhagen and Clayton 2003). Thus, transcription is coupled to DNA replication

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Figure 1. Human mitochondrial DNA (mtDNA). The genomic organization and structural features of human mtDNA are depicted in a circular genomic map. The D-loop regulatory region is expanded and shown *above*. Protein coding and rRNA genes are interspersed with 22 tRNA genes (denoted by the single-letter amino acid code). The D-loop regulatory region contains the L- and H-strand promoters (P_L and P_H , respectively) along with the origin of H-strand replication (O_H). mtDNA transcription complexes containing mitochondrial RNA polymerase, *Tfam*, and *TFB* are depicted in the expanded D-loop along with the conserved sequence blocks (CSB I, II, and III). The origin of L-strand replication (O_L) is displaced by approximately two-thirds of the genome within a cluster of five tRNA genes. Protein-coding genes include cytochrome oxidase (COX) subunits 1, 2, and 3; NADH dehydrogenase (ND) subunits 1, 2, 3, 4, 4L, 5, and 6; ATP synthase (ATPS) subunits 6 and 8; and cytochrome *b* (*Cytb*). *ND6* and the eight tRNA genes encoded on the L-strand are in bold type and underlined; all other genes are encoded on the H-strand.

and the sites of RNA cleavage are transition sites between RNA and DNA synthesis. A decision must be made to continue transcription through the CSBs or to truncate the nascent RNA to initiate DNA replication. After DNA synthesis begins, the nascent strand is often terminated downstream from a conserved element referred to as a termination-associated sequence (TAS). This event accounts for the triple-stranded D-loop structure and may be important in controlling mtDNA levels (Brown and Clayton 2002).

The mitochondrial H- and L-strand transcriptional units differ from most nuclear genes in that they are polygenic. In addition to the RNA primer for H-strand replication, P_L also directs the synthesis of a transcript that is processed to one mRNA and eight of the 22 tRNAs. The polygenic transcript directed by P_H is processed to 14 tRNAs, 12 mRNAs, and the two rRNAs (for review, see Garesse and Vallejo 2001). Both promoters share a critical upstream enhancer that serves as the recognition site for mitochondrial transcription factor A or *Tfam* (previously mtTF-1 and mtTFA), an HMG-box protein that stimulates transcription through specific binding to upstream enhancers. Like other HMG proteins, *Tfam* can bend and unwind DNA, properties potentially linked to its ability to stimulate transcription upon binding DNA (Fisher et al. 1992). In addition to specific promoter recognition, *Tfam* binds nonspecific DNA with high affinity. This property along with its abundance in mitochondria suggests that it plays a role in the stabilization and maintenance of the mitochondrial chromosome through its phased binding to nonpromoter sites (Parisi et al. 1993).

Several lines of evidence indicate that *Tfam* is required for mtDNA replication and maintenance. *Tfam* knock-out mice display embryonic lethality and depletion of mtDNA (Larsson et al. 1998). In addition, *Tfam* levels correlate well with increased mtDNA in ragged-red muscle fibers and decreased mtDNA levels in mtDNA-depleted cells (Larsson et al. 1994; Poulton et al. 1994). ABF2, a related HMG-box yeast factor, is required for mtDNA maintenance and respiratory competence (Diffley and Stillman 1991). Expression of *Tfam* can complement an ABF2 deficiency in yeast, suggesting that the two proteins are functionally homologous (Parisi et al. 1993). Interestingly, despite this functional complementation, ABF2 lacks an activation domain present in *Tfam* and does not stimulate transcription (Dairaghi et al. 1995).

Significant progress has been made in the characterization of the mtDNA transcription initiation machinery. A vertebrate mitochondrial RNA polymerase and a specificity factor that are required for mitochondrial-specific initiation were initially identified and characterized in *Xenopus laevis* (Antoshechkin and Bogenhagen 1995; Bogenhagen 1996). Although purification of the human polymerase has been elusive, a human cDNA that encodes a protein with sequence similarity to yeast mitochondrial and phage polymerases has been identified in database screenings (Tiranti et al. 1997). The encoded protein localizes to mitochondria, suggesting that it is a bona fide mitochondrial polymerase. A human mitochondrial transcription factor B (*h-mtTFB*) cDNA has also been isolated, and the encoded protein has properties consistent with it being a functional homolog of the

yeast specificity factor, sc-mtTFB (McCulloch et al. 2002). The protein is localized to mitochondria and can bind DNA and stimulate transcription from an L-strand promoter in vitro. Subsequently, two isoforms of h-mtTFB, termed TFB1 and 2, were identified (Falkenberg et al. 2002). TFB1 is identical to the initial h-mtTFB isolate. Like the yeast factor, both TFBs share sequence similarities with rRNA dimethyltransferases, although the similarity between TFB2 and this class of enzymes is weaker than that of TFB1. Both TFB isoforms can support specific initiation from mitochondrial promoters in an in vitro system containing purified recombinant proteins. In this system, the TFB-dependent activation of transcription depends on mitochondrial RNA polymerase and Tfam (Fig. 1). Both TFBs interact with mitochondrial RNA polymerase, but *TFB1* has about one-tenth the transcriptional activity of *TFB2*. In addition to binding mitochondrial RNA polymerase, TFB1 also contacts the C-terminal domain of Tfam (McCulloch and Shadel 2003). The region of contact between TFB1 and Tfam is essential for transcriptional activation and corresponds to a 29-amino-acid domain that was previously identified as a Tfam activation domain (Dairaghi et al. 1995). This reinforces the distinction between Tfam and the yeast HMG-box protein ABF2, which, like Tfam, is required for mtDNA maintenance but does not function as a transcription factor.

Transcriptional regulators of nuclear encoded mitochondrial proteins: the critical role of nuclear respiratory factors 1 and 2

The *cytochrome c* and *cytochrome oxidase* genes have served as the prototypes for identifying regulatory factors that act on nuclear respiratory genes from both yeast and mammalian cells. Early work in yeast demonstrated that transcriptional regulation of the major *cytochrome c* isoform, *CYC1*, was mediated by oxygen and carbon sources through the upstream activation sites, UAS1 and UAS2. This work has been the subject of excellent reviews to which the reader is referred for original citations (Zitomer and Lowry 1992; Poyton and McEwen 1996).

The identification of nucleus-encoded transcription factors required for the expression of the respiratory apparatus in mammalian cells also began with the characterization of the *cytochrome c* gene (for reviews, see Scarpulla 1997, 1999). Interestingly, the mammalian *cytochrome c* promoter has multiple recognition sites for transcription factors that bear no obvious relationship to those identified in yeast (Evans and Scarpulla 1988). A potent *cis*-acting element, localized to the first intron, consists of tandem Sp1 recognition sites that function synergistically to maximize promoter activity. A second *cis*-element binds transcription factors of the ATF/CREB family (Evans and Scarpulla 1989). The *cytochrome c* promoter also contains a recognition site for a transcription factor designated nuclear respiratory factor 1, or NRF-1 (Evans and Scarpulla 1989). NRF-1 is a 68-kD polypeptide with the presence of a C-terminal transcriptional activation domain comprised of glutamine-con-

taining clusters of hydrophobic amino acid residues (Chau et al. 1992; Gugneja et al. 1996). Both endogenous and recombinant proteins bind as a homodimer to palindromic NRF-1 sites through guanine nucleotide contacts over a single turn of the DNA helix (Virbasius et al. 1993a). Serine phosphorylation of the N-terminal domain of NRF-1 enhances both its DNA-binding (Gugneja and Scarpulla 1997) and *trans*-activation functions (Herzig et al. 2000).

NRF-1 has been linked to the transcriptional control of many genes involved in mitochondrial function and biogenesis (Table 1). NRF-1 target genes have been identified by characterization of functional NRF-1-binding sites within their promoters. Many NRF-1 target genes encode subunits of the five respiratory complexes (Virbasius et al. 1993a). However, the regulatory network controlled by NRF-1 extends beyond the respiratory subunits to other classes of genes. These include genes involved in assembly of the respiratory apparatus, constituents of the mtDNA transcription and replication machinery, mitochondrial and cytosolic enzymes of the heme biosynthetic pathway, and components of mitochondrial protein import. Notably, *Tfam* is an NRF-1 target gene consistent with the postulate that NRF-1 plays an integrative role in nucleo-mitochondrial interactions. This hypothesis has been reinforced by the results of several recent studies associating increases in *NRF-1* mRNA levels or DNA-binding activity with mitochondrial biogenesis. NRF-1 and its coactivator PGC-1 (see below) are induced as part of the adaptation of skeletal muscle to exercise training (Murakami et al. 1998; Baar et al. 2002). Similar results were obtained in cultured myotubes in response to elevated calcium, which mimics exercise-induced mitochondrial biogenesis (Ojuka et al. 2003). Likewise, treatment of rats with a creatine analog that induces muscle adaptations analogous to those observed during exercise leads to the activation of *AMP-activated protein kinase* and increased NRF-1-DNA binding activity, cytochrome *c* content, and mitochondrial density (Bergeron et al. 2001). Both *NRF-1* and *Tfam* mRNAs are elevated in cells depleted of mtDNA, presumably as a response to increased oxidative stress (Miranda et al. 1999). Lastly, *NRF-1* and Tfam are up-regulated in response to lipopolysaccharide-induced oxidative damage to mitochondria, presumably to enhance mtDNA levels and OXPHOS activity (Suliman et al. 2003).

Perhaps the strongest in vivo link between NRF-1 and the control of mitochondrial function comes from the results of targeted disruption of the *NRF-1* gene in mice (Huo and Scarpulla 2001). Homozygosity of the null allele results in lethality between embryonic days 3.5 and 6.5 (E3.5 and E6.5). The null blastocysts fail to grow in culture despite having a normal morphology. Homozygous null blastocysts are defective in maintaining a mitochondrial membrane potential and have severely reduced mtDNA levels. This is not accompanied by increased apoptosis, making it unlikely that the reduction in mtDNA is associated with a generalized increase in DNA fragmentation. Moreover, the mature oocytes of

Table 1. Nuclear and mitochondrial genes with NRF-1 and NRF-2 recognition sites

	NRF-1 ^a	NRF-2 ^a
Oxidative phosphorylation		
Rat cytochrome <i>c</i>	+	
Human cytochrome <i>c</i>	+	
Complex I:		
Human NADH dehydrogenase subunit 8 (TYKY)	+	
Complex II:		
Human succinate dehydrogenase subunit B	+	+
Human succinate dehydrogenase subunit C	+	+
Human succinate dehydrogenase subunit D	+	+
Complex III:		
Human ubiquinone-binding protein	+	
Human core protein I	+	
Complex IV:		
Rat cytochrome oxidase subunit IV		+
Mouse cytochrome oxidase subunit IV		+
Mouse cytochrome oxidase subunit Vb	+	+
Rat cytochrome oxidase subunit Vb	+	+
Human/primate cytochrome oxidase subunit Vb	+	+
Rat cytochrome oxidase subunit VIc	+	
Human cytochrome oxidase subunit VIaL	+	+
Bovine cytochrome oxidase subunit VIIaL	+	+
Human cytochrome oxidase subunit VIIaL		+
Bovine cytochrome oxidase subunit VIIc		+
Complex V:		
Bovine ATP synthase γ subunit	+	
Human ATP synthase c subunit	+	
Human ATP synthase β subunit		+
mtDNA transcription and replication		
Human Tfam	+	+
Mouse Tfam		+
Rat Tfam		+
Mouse MRP RNA	+	
Human MRP RNA	+	
Human TFB1	+	+
Mouse TFB1		+
Human TFB2	+	+
Mouse TFB2		+
HEME biosynthesis		
Rat 5-aminolivulinate synthase	+	
Mouse uroporphyrinogen III synthase	+	+
Protein import and assembly		
Human Tom 20	+	+
Human Tom 70		+
Mouse chaperonin 10	+	
Human SURF-1		+
Mouse COX17	+	+

(continued)

Table 1. (continued)

	NRF-1 ^a	NRF-2 ^a
Ion channels		
Human VDAC3	+	
Mouse VDAC3	+	
Human VDAC1		+
Shuttles		
Human glycerol phosphate dehydrogenase		+
Translation		
Human mitochondrial ribosomal S12	+	+

^aOriginal references for the majority of the indicated NRF-1 and/or NRF-2 target genes that are related to mitochondrial function have been cited elsewhere (Scarpulla 1997, 2002). New additions include human *Tomm 70* (Blesa et al. 2003) and mouse *COX 17* (Takahashi et al. 2002).

heterozygous mothers have a normal complement of mtDNA, supporting the argument against a defect in mtDNA amplification during oogenesis. Therefore, the mtDNA depletion occurs between fertilization and the blastocyst stage and most likely results from the loss of a NRF-1-dependent pathway of mtDNA maintenance. Interestingly, *Tfam*-null embryos also exhibit severely depleted levels of mtDNA but survive to E8.5–E10.5 (Larsson et al. 1998). Thus, it is likely that the early mortality of *NRF-1*-null embryos results from the combined effects of reduced levels of mtDNA and disruption of other NRF-1-dependent functions.

Characterization of cytochrome oxidase genes led to the identification of a second regulatory factor designated as NRF-2 (for review, see Scarpulla 1997, 1999). A series of directly repeated NRF-2 sites within the mouse *COXIV* promoter overlaps multiple transcription initiation sites and contains additional binding sites for the ETS-domain family of transcription factors (Virbasius and Scarpulla 1991; Carter et al. 1992). The complex binding the NRF-2 sites was purified to homogeneity from HeLa cell nuclear extracts and is comprised of five subunits. These include a DNA-binding α subunit and four others (β_1 , β_2 , γ_1 , and γ_2) that complex with α but alone do not bind DNA. The NRF-2 complexes activate transcription through four directly repeated ETS-domain-binding sites in the *COXVb* promoter, suggesting that NRF-2 may also act on multiple respiratory promoters (Virbasius et al. 1993b).

Purification and molecular cloning of all five NRF-2 subunits established that *NRF-2* is the human homolog of mouse *GABP* (LaMarco and McKnight 1989) and that the two additional human subunits, β_1 and γ_1 , were minor splice variants of *GABP* subunits β_1 and β_2 (Gugneja et al. 1995). The function of the non-DNA-binding subunits is twofold. First, the *GABP* β_1 subunit, corresponding to NRF-2 β_1 and NRF-2 β_2 (Gugneja et al. 1995), has a dimerization domain that facilitates cooperative binding of a heterotetrameric complex to tandem binding sites (Thompson et al. 1991). In solution, *GABP* exists as

an $\alpha\beta$ heterodimer but is induced to form the heterotetramer $\alpha_2\beta_2$ by DNA containing two or more binding sites (Chinenov et al. 2000). The crystal structure of the heterotetramer bound to DNA has been determined (Batchelor et al. 1998). The second function of the non-DNA-binding subunits is to contribute a transcriptional activation domain. This domain resembles that found in NRF-1 and has been localized to a region upstream from the homodimerization domain (Gugneja et al. 1996).

Functional NRF-2 sites have now been identified in several COX promoters as well as in many other genes related to respiratory chain expression (for review, see Scarpulla 2002). As with NRF-1, the list of respiratory genes containing NRF-2 sites has expanded in recent years (Table 1). These include genes for *Tfam* (Larsson et al. 1998; Rantanen et al. 2001) and the newly discovered TFB factors (Falkenberg et al. 2002; McCulloch et al. 2002) involved in mitochondrial transcription and DNA replication (Rantanen et al. 2001). Genes encoding three of the four human succinate dehydrogenase (complex II) subunits also have both NRF-1 and NRF-2 sites in their promoters (Au and Scheffler 1998; Elbehti-Green et al. 1998; Hirawake et al. 1999). In many cases, NRF-1 sites are also present in NRF-2-dependent promoters, but this is not a general rule. For example, several COX promoters and the rodent *Tfam* (Choi et al. 2002) and TFB (Rantanen et al. 2003) promoters do not have obvious NRF-1 consensus sites (Table 1). This contrasts with the human *Tfam* (Virbasius and Scarpulla 1994) and TFB (R.C. Scarpulla, unpubl.) promoters, which rely on functional NRF-1 and NRF-2 recognition sites for their activities.

A subset of respiratory genes does not appear to be regulated by NRF-1 or NRF-2. Other well-characterized regulatory factors have been implicated in the expression of these genes. The transcription factor Sp1 is associated with the activation and/or repression of *cytochrome c₁* (Li et al. 1996b) and *adenine nucleotide translocase 2* genes (Li et al. 1996a), both of which lack NRF sites (Zaid et al. 1999). Sp1 sites are also common to many GC-rich promoters including those that are NRF-dependent. The muscle-specific COX subunits, *COXVIaH* and *COXVIII*, are also lacking NRF sites but depend on MEF-2 and/or E-box consensus elements for their expression (Wan and Moreadith 1995). Thus, the same or similar factors required for the expression of other muscle-specific genes are linked to the regulation of these tissue-specific COX subunits. In contrast, the promoter of the ubiquitously expressed liver isoform, *COXVIaL*, depends on NRF-1 and NRF-2 as well as Sp1 for full activity (Seelan et al. 1996). This is consistent with the observation that in gene pairs encoding ubiquitous and tissue-specific isoforms of a given protein, the NRF-1 site, when present, is associated with the ubiquitously expressed gene (Virbasius et al. 1993a). Finally, the initiator element transcription factor YY1 participates in the expression of certain COX genes. Functional YY1-binding sites have been detected in the promoters of genes encoding *COXVb* (Basu et al. 1997) and *COXVIIc* (Seelan and Grossman 1997). Multiple YY1 sites in the *COXVb* promoter bind YY1 and possibly other factors, and at least one of these

sites helps confer a negative regulatory effect on *COXVb* promoter activity (Basu et al. 1997). In the *COXVIIc* promoter, two YY1 sites in conjunction with an NRF-2 site act as positive regulators of promoter activity (Seelan and Grossman 1997). It is also important to note that regulation of most nuclear genes encoding mitochondrial enzymes upstream of the respiratory chain is NRF-1/NRF-2-independent. For example, genes encoding mitochondrial fatty acid oxidation enzymes are regulated by the peroxisome proliferator-activated receptor alpha (PPAR α) and other NRF-1-independent regulatory pathways (Gulick et al. 1994). Thus, any unifying transcriptional model of mitochondrial biogenesis needs to account for the expression of genes that are NRF-independent.

There are several reports suggesting that nuclear and mitochondrial genes are controlled by common *cis*-acting elements that are the targets of the same or similar transcription factors. Sequence similarities to the OXBOX/REBOX (Haraguchi et al. 1994) and Mt (Suzuki et al. 1995) elements have been localized to the mitochondrial D-loop. The ability of these elements and their nuclear gene counterparts to bind proteins from crude extracts with the same specificity has been taken as evidence for shared regulatory factors between the two genetic systems (Haraguchi et al. 1994). Similarly, other nuclear factors, such as thyroid hormone receptors, have been implicated in mitochondrial gene expression (for review, see Wrutniak-Cabello et al. 2001). However, there is no evidence that these proteins can use the mitochondrial transcriptional machinery to direct mitochondrial gene expression.

The critical role of transcriptional coactivators in the mitochondrial biogenic regulatory cascade: The PPAR γ coactivator-1 (PGC-1) family

As described above, the mitochondrial biogenic program involves the integration of multiple transcriptional regulatory pathways controlling the expression of both nuclear and mitochondrial genes. This highlights a mechanistic enigma fundamental to the control of mitochondrial biogenesis. How is the activity of multiple transcription factors (e.g., NRF-1, NRF-2, PPAR α , mtTFA) coordinately regulated during the mitochondrial biogenic process? Moreover, in the context of such complex integration, how is cell- and tissue-specific function achieved? For example, mitochondria within the brown adipocyte are poised for uncoupled mitochondrial respiration, whereas in other tissues such as heart, mitochondrial respiration is largely coupled for high-level ATP production. To add to the complexity, skeletal muscle is capable of supporting both coupled and uncoupled respiration. New insight into this problem was provided by the discovery of the transcriptional coactivator PPAR γ coactivator 1 α (PGC-1 α) by Spiegelman and colleagues (Puigserver et al. 1998). PGC-1 α was cloned in a yeast two-hybrid screen for brown adipose-specific factors that interacted with the adipogenic nuclear receptor PPAR γ (Puigserver et al. 1998). PGC-1 α serves as a direct transcriptional coacti-

vator of PPAR γ and is a member of a growing list of proteins that coactivate transcription factors through direct protein–protein interactions (for review, see Knutti and Kralli 2001; Puigserver and Spiegelman 2003).

Transcriptional coactivators serve multiple functions including modification of chromatin through posttranslational histone acetylation, direct interaction with the RNA polymerase II complex, mRNA processing, and recruitment of other transcriptional coactivators (for review, see Robyr et al. 2000; Belandia and Parker 2003). Present evidence indicates that PGC-1 α coactivates its targets via recruitment of additional coactivators with histone acetylase activity, such as SRC-1 (Puigserver et al. 1999). In addition, the PGC-1 α molecule contains domains capable of interacting with and processing pre-mRNA (Monsalve et al. 2000). PGC-1 α also interacts directly with the TRAP/Mediator complex (Wallberg et al. 2003). Unlike most known transcriptional coactivators, PGC-1 α is unique in that it exhibits a tissue-enriched expression pattern and is highly inducible (Puigserver et al. 1998; Knutti and Kralli 2001; Puigserver and Spiegelman 2003). PGC-1 α is enriched in brown adipose, heart, slow-twitch skeletal muscle, and kidney—tissues with high-capacity mitochondrial systems. The expression of the *PGC-1 α* gene is rapidly induced by cold exposure, short-term exercise, and fasting; physiologic conditions known to increase the demand on mitochondria to produce heat or ATP (Puigserver et al. 1998; Wu et al. 1999; Goto et al. 2000; Lehman et al. 2000; Baar et al. 2002; Terada et al. 2002; Irrcher et al. 2003; Pilegaard et al. 2003; Terada and Tabata 2003). These latter observations suggested that PGC-1 α is involved in the physiologic control of mitochondrial function.

Several lines of evidence indicate that the transcriptional coactivator PGC-1 α serves as a key regulator of mitochondrial biogenesis in mammals. First, studies focused on the biologic function of PGC-1 α revealed that it activates the transcription of mitochondrial uncoupling protein-1 (*UCP-1*) through interactions with the nuclear hormone receptors PPAR γ and thyroid hormone receptor (Puigserver et al. 1998). These findings further supported a role for PGC-1 α in the process of mitochondrial uncoupled respiration and thermogenesis in brown adipose tissue. Second, forced expression studies in adipogenic and myogenic mammalian cell lines demonstrated that PGC-1 α markedly induces the expression of *NRF-1*, *NRF-2*, and *Tfam* (Wu et al. 1999). PGC-1 α can also interact directly with and coactivate NRF-1 on the *Tfam* gene promoter. Third, studies in primary cardiac myocytes in culture and in the hearts of transgenic mice have demonstrated that overexpression of *PGC-1 α* up-regulates the expression of genes involved in mitochondrial fatty acid oxidation, most of which are PPAR α targets, in addition to NRF-1 targets (Lehman et al. 2000). Cardiac-specific overexpression of *PGC-1 α* in transgenic mice leads to massive mitochondrial proliferation, ultimately resulting in cardiomyopathy and death (Lehman et al. 2000). Interestingly, in neonatal cardiac myocytes in culture, PGC-1 α induces mitochondria that support largely coupled respiration consistent with the known ATP-gen-

erating function of this organelle in heart (Lehman et al. 2000). Lastly, forced expression of *PGC-1 α* in skeletal muscle of transgenic mice triggers mitochondrial proliferation and the formation of mitochondrial-rich type I, oxidative (“slow-twitch”) muscle fibers (Lin et al. 2002b). Collectively, these results indicate that PGC-1 α is capable of promoting mitochondrial biogenesis through its coactivating effects on key factors such as NRF-1.

The gain-of-function studies described above provide compelling evidence that PGC-1 α serves as a transcriptional coactivator to promote mitochondrial biogenesis in postnatal mammalian tissues. Although *NRF-1* is a key target of PGC-1 α , it is clear that this transcription factor does not control all of the components of the mitochondrial biogenic response. Multiple PGC-1 α targets have now been identified, indicating that this coactivator serves as a pleiotropic regulator of multiple pathways involved in cellular energy metabolism within and outside of the mitochondrion (Knutti and Kralli 2001; Puigserver and Spiegelman 2003). Following the identification of PPAR γ as the initial PGC-1 α transcription factor target, a variety of additional members of the nuclear receptor superfamily have been shown to interact with PGC-1 α . This list includes PPAR α (Vega et al. 2000), thyroid hormone receptor (Puigserver et al. 1998), retinoid receptors (Puigserver et al. 1998), glucocorticoid receptor (Knutti et al. 2000), estrogen receptor (Puigserver et al. 1998; Knutti et al. 2000; Tcherepanova et al. 2000), HNF-4 (Rhee et al. 2003), and estrogen-related receptors (ERRs; Huss et al. 2002; Schreiber et al. 2003). In addition, several non-nuclear-receptor PGC-1 α partners have been identified, in addition to NRF-1, including myocyte-enhancing factor-2 (MEF-2; Michael et al. 2001) and FOX-01 (Puigserver et al. 2003). Although several of the PGC-1 α partners serve functions outside of the mitochondrion such as HNF-4 and FOX-01 (gluconeogenesis; Rhee et al. 2003; Puigserver et al. 2003) and MEF-2 (glucose transport; Michael et al. 2001), others are linked to the mitochondrial biogenic transcriptional regulatory program. For example, PGC-1 α coactivates the nuclear receptor PPAR α , a key regulator of nuclear genes encoding mitochondrial fatty acid oxidation enzymes (Vega et al. 2000). More recently, PGC-1 α was found to coactivate the orphan nuclear receptors ERR α and ERR γ (Huss et al. 2002; Schreiber et al. 2003). Although the exact biologic function of ERRs has not been delineated, ERR α and ERR γ are enriched in tissues with high mitochondrial oxidative capacity including brown adipose tissue and heart. In addition, *medium-chain acyl-CoA dehydrogenase* (MCAD), a known PPAR α target that catalyzes the initial step in mitochondrial fatty acid β -oxidation, is also regulated by ERR α (Sladek et al. 1997; Vega and Kelly 1997; Huss et al. 2002). These results suggest that ERR α and PPAR α may drive distinct but overlapping mitochondrial pathways downstream of PGC-1 α .

PGC-1 α is now known to be a member of a family of transcriptional coactivators. The first PGC-1 α relative, *PGC-1-related coactivator* (PRC), was identified through a database search (Andersson and Scarpulla 2001). PRC

The differences in regulation and tissue-expression patterns of PGC-1 family members suggest that each confers distinct biologic responses. In support of this idea, recent work by the Spiegelman laboratory has provided evidence that PGC-1 α and PGC-1 β isoforms exert coactivator-specific bioenergetic effects (St-Pierre et al. 2003). Specifically, overexpression studies in C₂C₁₂ myotubes demonstrated that although both PGC-1 α and

Upstream signaling events involved in the control of mitochondrial biogenesis: PGC-1 as an integrative coactivator

Signal transduction pathways play a major role in the physiologic regulation of mitochondrial function and biogenesis; therefore, it is not surprising that *PGC-1 α* activity and expression are regulated by similar signaling pathways. In tissues poised for mitochondrial thermogenesis, such as brown adipose, the β -adrenergic/cAMP pathway is upstream of the PGC-1 α -mediated regulation of targets such as *UCP-1* [Puigserver et al. 1998]. A sig-

nificant body of evidence focused largely on skeletal muscle indicates that in response to contractile activity, calcium-dependent signaling pathways trigger a cascade of regulatory events leading to increased formation of oxidative fiber types and a marked increase in mitochondrial number and function (Holloszy and Coyle 1984; Chin et al. 1998). Several important gain-of-function studies have now provided evidence for regulatory links between calcineurin A, calcium/calmodulin-dependent protein kinase (CaMK), PGC-1 α , and skeletal muscle mitochondrial biogenesis. First, overexpression of *CaMK* in the skeletal muscle of transgenic mice triggers a robust mitochondrial biogenesis associated with an induction of *PGC-1 α* expression (Wu et al. 2002). Second, overexpression of *PGC-1 α* in the skeletal muscle of transgenic mice leads to the formation of slow-twitch skeletal muscle fibers and an induction of genes involved in mitochondrial oxidative metabolism (Lin et al. 2002b). Third, studies performed in myogenic cell lines indicate that both calcineurin A and CaMK are capable of activating *PGC-1 α* gene transcription (Handschin et al. 2003). The calcineurin A-mediated activation of *PGC-1 α* transcription is dependent on MEF2 response elements, whereas CaMK-mediated regulation requires CREB-binding sites.

Several other signal transduction pathways have been implicated in the control of *PGC-1* expression and activity. p38 MAPK activates *PGC-1 α* by releasing repression of an unidentified factor and by increasing PGC-1 α protein stability (Knutti et al. 2001; Puigserver et al. 2001). p38 MAPK can also activate the PGC-1 α partner, *PPAR α* , suggesting that activation of this signaling pathway influences mitochondrial fatty acid oxidation (Barger et al. 2001). However, the role of the p38 MAPK pathway in regulating mitochondrial biogenesis is not known. More recently, evidence has emerged that nitric

oxide (NO) activates mitochondrial biogenesis in a variety of cell types including adipocytes, and HeLa cells (Nisoli et al. 2003). The mitochondrial thermogenic response is significantly altered in mice lacking *eNOS*. This NO effect is dependent on cGMP and linked to *PGC-1 α* activation. These results raise the intriguing possibility that mitochondrial biogenesis is one of the important effects of NO activation. Given the known role of NO as a vasodilator, it is tempting to speculate that this key upstream regulatory pathway coordinately regulates downstream events including an increase in the capacity to use oxygen in mitochondria.

Summary

Over the past decade, significant new insight has been gained into the circuitry of molecular regulatory cascades controlling mitochondrial biogenesis and function (Fig. 3). The interdependence of nuclear and mitochondrial genomes has evolved with the emergence of the mitochondrion as a eukaryotic organelle. It is likely that the complexity of the mammalian organism mandates a complex regulatory network that provides for the dynamic coordinate control of nuclear and mitochondrial genes during development and in the adult. This regulatory circuitry not only triggers mitochondrial biogenesis in response to developmental and physiologic cues, but also confers cell- and tissue-specific features. New insight into the dynamic control of mitochondrial function and biogenesis has been provided by the identification of relevant transcription factors, transcriptional coactivators, and upstream signaling events. However, the mechanisms involved in the control of cell-specific mitochondrial phenotypes and the full cast of transcriptional regulatory factors comprise an exciting investigative frontier. New experimental approaches such as the

Figure 3. PGC-1 serves a central integrative role in the transcriptional regulatory cascade upstream of the mitochondrial biogenic response. A schematic representation of the mitochondrial biogenic regulatory cascade, including known PGC-1 partners and putative upstream signaling pathways.

delineation of tissue-specific mitochondrial proteomes (Mootha et al. 2003) should provide an excellent framework for future studies aimed at understanding the molecular events involved in defining the mitochondrial phenotype.

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