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## ROS-Triggered Phosphorylation of Complex II by Fgr Kinase Regulates Cellular Adaptation to Fuel Use

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### SUMMARY

Electron flux in the mitochondrial electron transport chain is determined by the superassembly of mitochondrial respiratory complexes. Different superassemblies are dedicated to receive electrons derived from NADH or FADH<sub>2</sub>, allowing cells to adapt to the particular NADH/FADH<sub>2</sub> ratio generated from available fuel sources. When several fuels are available, cells adapt to the fuel best suited to their type or functional status (e.g., quiescent versus proliferative). We show that an appropriate proportion of superassemblies can be achieved by increasing CII activity through phosphorylation of the complex II catalytic subunit FpSDH. This phosphorylation is mediated by the tyrosine-kinase Fgr, which is activated by hydrogen peroxide. Ablation of Fgr or mutation of the FpSDH target tyrosine abolishes the capacity of mitochondria to adjust metabolism upon nutrient restriction, hypoxia/reoxygenation, and T cell activation, demonstrating the physiological relevance of this adaptive response.

### INTRODUCTION

To utilize fuels efficiently, cells must exquisitely integrate the activities of membrane receptors and transporters, the intracellular compartmentalization of molecules, the enzymatic balance of each metabolic step, and the elimination of byproducts (Stanley et al.,

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#### SUPPLEMENTAL INFORMATION

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2013). Appropriate orchestration of all these changes is critical for the cell's ability to adapt to changing functional requirements, such as quiescence, proliferation, and differentiation, and to environmental changes, including survival in response to diverse insults. Factors known to influence this adaptation include the cellular response to oxygen availability (hypoxia-inducible factors HIF1 $\alpha$  and HIF1 $\beta$ ); regulators of energy availability such as mammalian target of rapamycin (mTOR), AMP-activated protein kinase, sirtuin, and forkhead box (FOX)O; and mediators of the response to reactive oxygen species (ROS), such as peroxisome proliferator-activated receptor gamma coactivator-1 alpha (PGC-1 $\alpha$ ). The involvement of these factors illustrates the interconnection between the use of alternate carbon substrates (carbohydrates, amino acids, fatty acids and ketone bodies) and the cellular response to stress, particularly oxidative stress.

At the core of this process are mitochondria. In response to changes in fuel source, mitochondria must modify their location, structure, and metabolite fluxes in order to balance their contribution to anabolism (lipogenesis and antioxidant defenses from citrate, gluconeogenesis, serine and glycine biosynthesis from pyruvate, nucleotide biosynthesis) and catabolism (TCA cycle,  $\beta$ -oxidation, oxidative phosphorylation). Mitochondria are central to ATP synthesis, redox balance, and ROS production, parameters directly dependent on fuel use. All catabolic processes converge on the mitochondrial electron transport chain (mETC) by supplying electrons in the form of NADH<sup>+</sup>H<sup>+</sup> or FADH<sub>2</sub>. The relative proportion of electrons supplied via NADH and FADH<sub>2</sub> varies with the fuel used; for example, oxidative metabolism of glucose generates a NADH/FADH<sub>2</sub> electron ratio of 5, whereas for a typical fatty acid (FA) such as palmitate the ratio is  $\approx$ 2 (Speijer, 2011).

Our recent work on the dynamic architecture of the mETC reveals that supercomplex formation defines specific pools of CIII, CIV, CoQ, and cyt c for the receipt of electrons derived from NADH or FAD (Lapiente-Brun et al., 2013). Since CIII preferentially interacts with CI, the amount of CI determines the relative availability of CIII for FADH<sub>2</sub>- or NADH-derived electrons. The regulation of CI stability is thus central to cellular adaptation to fuel availability. A substrate shift from glucose to FA requires greater flux from FAD, and this is achieved by a reorganization of the mETC superstructure in which CI is degraded, releasing CIII to receive FAD-derived electrons (Lapiente-Brun et al., 2013; Stanley et al., 2013). Failure of this adaptation results in the harmful generation of reactive oxygen species (ROS) (Speijer, 2011). The proportion of supercomplexes dedicated to receiving NADH electrons is further dependent on the structure and dynamics of mitochondrial cristae (Cogliati et al., 2013; Lapiente-Brun et al., 2013), so that reducing the number of cristae favors flux from FAD. In agreement with this, ablation of the mitochondrial protease OMA1, which prevents optic atrophy 1 (OPA1)-specific proteolysis and cristae remodeling, impairs FA degradation in mice, resulting in obesity and impaired temperature control (Quirós et al., 2012).

Cells are normally exposed to a mixed supply of fuels, but despite this, cells are often predisposed to preferentially use one source over another, according to their physiological role or status (Stanley et al., 2013). T cells, for example, switch from oxidative to glycolytic metabolism upon activation, coinciding with entry into a proliferative state, and later increase FA oxidation when they differentiate into regulatory T cells. These changes require

remodeling of the mETC NADH/FADH<sub>2</sub> flux capacity, but how cells regulate this choice of carbon source is not understood.

Here, we show that fuel choice is regulated via tyrosine phosphorylation of complex II (CII) subunit FpSDH, mediated by ROS-activation of the tyrosine kinase Fgr. This activation is required to adjust the level of complex I (CI) to optimize NADH/FADH<sub>2</sub> electron use. Our data show this mechanism operating in three physiological situations: upon T lymphocyte activation, in the adaptation of liver and cultured cells to starvation, and in the adaptation of cells to hypoxia/reoxygenation.

## RESULTS

### Above-Normal CII Activity in Cells Expressing Mutant CI

Our laboratory has isolated mouse cell lines carrying different proportions of a null ND6 mutation (Acín-Pérez et al., 2003): EB2615 (30% mutant mtDNA), E23 (66%), E12 (80%), FG12-1 (95%), and FG23-1 (98%). Mitochondria from ND6 mutants showed reductions in CI proportional to the mutation load (Figure 1A). Interestingly, the lines with the highest mutation loads showed elevated activity of CII (Figure 1B). Similar observations have been reported in human cells derived from patients with CI deficiencies (Cardol et al., 2002; Fan et al., 2008; Majander et al., 1991; Pitkanen and Robinson, 1996) and in CI-deficient mice (Kruse et al., 2008). Increased CII activity was proposed to compensate for impaired activity of CI, but the underlying mechanism is unknown. This compensatory phenomenon is specific for CI-deficient cells, since cells lacking CIII or CIV showed no changes in CII activity (Figure S1A).

As reported (Acín-Pérez et al., 2003; Bai and Attardi, 1998), the amount of CI was reduced in cells with a high ND6 mutant load (FG12-1, FG23-1; Figure 1C, upper panel), whereas CII content was unchanged, despite above-normal activity determined spectrophotometrically (Figure 1B) and by in-gel assay (data not shown). G3PDH activity did not differ between WT and ND6 mutants (Figure 1D), indicating that increased CII activity does not reflect a generalized response of enzymes that donate electrons to CIII and is a specific response to CI deficiency.

As a further control, we treated WT FBalb/cJ cells with the specific inhibitor rotenone (200 nM), which blocks CI activity without affecting CI assembly. Rotenone-mediated inhibition of CI was accompanied by a parallel increase in CII activity, after short and long rotenone treatment, indicating that loss of CI function, not its physical absence, is responsible for the high CII activity (Figure 1E).

### CII Activity Is Regulated by Phosphorylation

Several groups have proposed that CII subunit A (FpSDH) is a kinase target (Bykova et al., 2003; Salvi et al., 2007; Schulenberg et al., 2003). To investigate FpSDH phosphorylation, we separated mitochondrial proteins by 2D IEF/SDS-PAGE (isoelectric focusing followed by SDS polyacrylamide gel electrophoresis) and detected FpSDH protein by immunoblot (Figure 1F). The CI subunit NDUFS3 was used as a reference since the stability and migration of this protein are unaffected by the failure of CI assembly (Figure S1B). FpSDH

in FG23-1 samples migrated as multiple spots, running at more acidic positions than samples from WT FBalb/cJ cells. This is compatible with increased phosphorylation of the protein in mutant cells. Treatment of permeabilized FG23-1 mitochondria with calf-intestine phosphatase (CIP) to remove phosphoryl residues restored the WT migration pattern, confirming that the altered pattern is due to phosphorylation (Figure 1F, lower panels). Moreover, blockade of CI activity in FBalb/cJ cells reproduced the FpSDH mobility pattern seen in FG23-1 cells (not shown). These observations indicate that FpSDH is phosphorylated when CI activity is impaired. Consistent with these findings, most FpSDH from FG23-1 mitochondrial samples eluted in the phosphorylated fraction after separation on phosphoprotein-enrichment columns, and this effect was blocked by pretreatment of permeabilized mitochondria with CIP (Figure S1C). In contrast, FpSDH from FBalb/cJ mitochondria was more concentrated in nonphosphorylated fractions (Figure S1C). The activity of CII in mitochondria from FBalb/cJ and FG23-1 cells was sensitive to CIP-mediated dephosphorylation, but this reduction was proportionally more severe in FG23-1 mitochondria, suggesting that phosphorylation increases CII activity (Figure 1G).

### Phosphorylation of FpSDH Is Triggered by ROS

Cells lacking CI contain high levels of ROS (Robinson, 1998), and ND6-deficient cells produce abundant hydrogen peroxide (Moreno-Loshuertos et al., 2006) (Figure 2A), an effect mimicked by treatment with the CI inhibitor rotenone (Dlasková et al., 2008; Radad et al., 2006). Short and long rotenone treatment of WT cells showed an increase in CII activity (Figure 1E) that was accompanied by a parallel increase in ROS production (Figure S2A). ROS modulate several signaling pathways (Dröge, 2002; Hamanaka and Chandel, 2010), prompting us to evaluate the involvement of ROS in CII activation. Culturing cells for 1 week with N-acetyl cysteine (NAC, 5 mM) lowered basal H<sub>2</sub>O<sub>2</sub> levels and decreased CII activity in control and mutant cells (Figures 2A and 2B), but the reduction was proportionally stronger in FG23-1 cells (Figure 2B). Since NAC is not a direct ROS scavenger but rather a precursor of glutathione, which changes the redox status of the cell, we investigated H<sub>2</sub>O<sub>2</sub>-dependent CII activity stimulation in isolated mitochondria. Addition of H<sub>2</sub>O<sub>2</sub> to WT mouse liver mitochondria increased CII activity (Figure 2C) and the phosphorylation of FpSDH (Figure 2D). CII activation was blunted by the presence of catalase, which catalyzes H<sub>2</sub>O<sub>2</sub> decomposition (Figure 2C). As a further control, we analyzed the response to moderate ROS production of fibroblasts lacking supercomplex assembly factor I (SCAFI), which is required for association between CIII and CIV (Lapuente-Brun et al., 2013). The response of WT and SCAFI-deficient fibroblasts to ROS (generated by incubation in the presence of xanthine and xanthine oxidase) was indistinguishable, indicating that ROS-induced upregulation of CII is a general response independent of the assembly of supercomplexes containing CIV (Lapuente-Brun et al., 2013) (Figure 2E). To determine whether CII is activated by superoxide or hydrogen peroxide produced in the mitochondrial matrix, we generated FBalb/cJ- and FG23-1-derived lines expressing mitochondria-targeted catalase (mt-cat) or MnSOD (Figure S2B). ROS production as well as catalase and MnSOD activities measured before and after overexpression confirmed that the cells were expressing functional enzymes (Figures S2C–S2F). Control and CI mutant cells also exhibited basal differences in catalase and MnSOD activities, as previously shown (Moreno-Loshuertos et al., 2006). The expected elevation in

H<sub>2</sub>O<sub>2</sub> production and CII activity in FG23-1 cells and rotenone-treated FBalb/cJ cells was decreased by expression of mt-cat, but not MnSOD (Figures 2F and 2G), suggesting that the key activator of the pathway is H<sub>2</sub>O<sub>2</sub> and not superoxide.

### CII Is Activated by Phosphorylation on FpSDH Mediated by a Src-Type Tyrosine Kinase

We next analyzed immunoprecipitated FpSDH by western blot with anti-phospho-Tyr, anti-phospho-Ser, and anti-phosphoThr antibodies. Only Tyr phosphorylation of FpSDH was detected in FBalb/cJ and FG23-1 cells, but the signal was stronger in FG23-1 cells (Figure 3A). Incubation of isolated mouse liver mitochondria with protein kinase inhibitors revealed that CII activity and FpSDH phosphorylation were reduced by PP2, an inhibitor of Src-family kinases (SFK), whereas no effect was observed with the cAMP/cGMP-dependent protein kinase inhibitor H89 or the PKA agonist 8Br-cAMP (Figures 3B and 3C). These results thus suggest that FpSDH is phosphorylated *in vivo* by a Src-type tyrosine kinase.

H<sub>2</sub>O<sub>2</sub> can promote phosphorylation on Tyr residues by activating Tyr kinases (Balamurugan et al., 2002; Chiarugi, 2008; Minetti et al., 2002) or by inhibiting Tyr phosphatases (Chiarugi, 2008). We incubated isolated mouse liver mitochondria with H<sub>2</sub>O<sub>2</sub> in the presence of either PP2, to inhibit SFKs, or orthovanadate (Ov), a general inhibitor of Tyr-phosphatases. PP2 reduced the activity of CII and prevented the activation induced by H<sub>2</sub>O<sub>2</sub>, whereas Ov had no effect. If we assume that Ov is able to inhibit the phosphatase involved, these results are compatible with a model in which H<sub>2</sub>O<sub>2</sub> promotes FpSDH phosphorylation by activating a PP2-sensitive kinase (Figure 3D). Full demonstration of this mechanism would require the identification of the phosphatase involved.

### The Src-Type Tyrosine Kinase Fgr Interacts with CII In Vivo

The SFKs Lyn, Fyn, Fgr, and Csk have been reported to localize in mitochondria (Augereau et al., 2005; Salvi et al., 2005; Tibaldi et al., 2008). Immunoblot analysis of mitochondrial preparations pretreated with proteinase K detected several tyrosine protein kinases (Src, Lyn, and Fgr), the regulator of Src-type tyrosine kinases Csk (itself a Tyr kinase), and the Ser/Thr kinase PKA (Figure 3E). To establish whether any of these interact with CII, we performed coimmunoprecipitation experiments targeting FpSDH. Mitochondrial membranes were solubilized either with the nonionic detergent dodecyl-maltoside (DDM), to isolate individual respiratory complexes, or with the milder detergent digitonin (DIG), to isolate supercomplexes. In the DDM lysates, none of the probed protein-kinases was coimmunoprecipitated with CII (Figure 3F), whereas in DIG lysates anti-FpSDH specifically coimmunoprecipitated the Src-family kinase Fgr, suggesting physical interaction between Fgr and CII in the mitochondrial inner membrane. To confirm this, we separated DIG and DDM lysates by BN-PAGE followed by denaturing SDS-PAGE (Figure 3G). Fgr kinase was detectable in both preparations but only comigrated with CII in the DIG-lysed samples. None of the other kinases analyzed comigrated with FpSDH, but Src/Csk and PKA both appeared to be associated with high molecular weight complexes (Figure 3G). Interestingly, Src comigrated with its regulator Csk in both preparations, suggesting strong interaction between these kinases. The mitochondrial matrix localization of Fgr kinase was confirmed by subfractionation of pure mouse liver mitochondria (Figure 3H).

## Ablation of Fgr Abolishes the Activation of CII

To demonstrate the role of Fgr in the regulation of CII we examined liver mitochondria from *fgr*<sup>-/-</sup> mice (Lowell et al., 1994), in which Fgr is undetectable but CII content is normal (Figure 4A). Basal CII activity in these mitochondria was below normal and insensitive to H<sub>2</sub>O<sub>2</sub> or PP2 (Figure 4B), and FpSDH phosphorylation was prevented (Figure 4C). CII activity was normal in mitochondria from Lyn and Hck Tyr kinase knock mice (Figure 4D). The absence of Fgr affected CII activity in all analyzed tissues (Figure 4E), whereas activities of the other mitochondrial respiratory complexes remained unchanged or, in the case of CI or combined CI+III, increased (Figures 4F and 4G). We can thus conclude that Fgr is the tyrosine kinase responsible for FpSDH phosphorylation, that it interacts with its substrate, and that FpSDH phosphorylation activates CII.

## Fgr Activates CII through FpSDH Phosphorylation on Y604

Analysis of mouse FpSDH identified six candidate tyrosines for phosphorylation (Figure S3A), three of them in Fgr target motifs predicted by Scansite and Prediction of PK phosphorylation site (PPSP). One residue, Y604, which is conserved among mammals (Figure S3B), was previously identified as an Fgr target in in vitro phosphorylation assays (Salvi et al., 2007). The same report showed phosphorylation of Y543, but in silico predictions indicate this residue is not an Fgr target.

To identify the residue involved in CII regulation, we measured CII activity in FBalb/cJ cells transfected with WT FpSDH or one of the mutant FpSDH variants Y543F and Y604F. To ascertain the effect of the specific residues, we also silenced endogenous FpSDH expression (Figure S3C) and analyzed the response to treatment with rotenone (Figure 4H). In cells silenced for endogenous FpSDH, re-expressed WT and Y543F FpSDH restored rotenone-activated CII activity, but this effect was not seen in cells re-expressing the single mutant Y604F (Figure 4H). We next immunocaptured CII from cells expressing WT or Y604F FpSDH and performed Fgr in vitro phosphorylation assays. As proposed for rat (Salvi et al., 2007), mouse FpSDH was phosphorylated by Fgr. In contrast, Fgr was unable to phosphorylate the Y604F FpSDH variant (Figure 4I). No other CII proteins were labeled by this approach. These results confirm the in silico prediction that FpSDH Y604 is the unique Fgr target in CII.

## ROS/Fgr/CII Regulatory Pathway in T Lymphocyte Activation

Activation of resting T cells with anti-CD3/CD28 antibodies (Abs), concanavalin A/IL2 (ConA/IL2), or phorbol-12-myristate-13-acetate (PMA) plus ionomycin triggers ROS production (Kamiski et al., 2010; Nagy et al., 2003). In response to anti-CD3/CD28 Abs or ConA/IL2, T cells isolated from *Fgr*<sup>+/-</sup> and *Fgr*<sup>-/-</sup> produced similar levels of ROS (Figures 5A and 5B). However, whereas CII activity was increased in *Fgr*<sup>+/-</sup> T cells, it remained unaffected in *Fgr*<sup>-/-</sup> cells (Figure 5C). Consistent with this result, restimulation of T lymphoblasts with PMA/ionomycin increased CII activity in *Fgr*<sup>+/-</sup> but not *Fgr*<sup>-/-</sup> cells (not shown).

## ROS/Fgr/CII Pathway in Starvation and Serum Deprivation

One of the better-characterized ways in which liver cells adapt to starvation is through phosphorylation-mediated inhibition of pyruvate dehydrogenase (Huang et al., 2003), which reduces mitochondrial utilization of pyruvate and has been proposed to increase ROS production (Ten and Starkov, 2012). We recently reported that electrons derived from NADH and FADH<sub>2</sub> follow separate routes along the mETC, and that this is important for metabolic adaptation to starvation, with an increase in the rate of FADH<sub>2</sub> oxidization occurring at the expense of the NADH route (Lapiente-Brun et al., 2013). Since Fgr-kinase activates CII (and hence use of FADH<sub>2</sub> electrons), we conducted a more detailed evaluation of the consequences of the lack of Fgr-kinase for liver mitochondria function. The significantly lower maximum CII activity in mitochondrial preparations from Fgr null liver was accompanied by lower-succinate-driven respiration and ATP synthesis (Figure 6A). Maximum CI activity was similar for both genotypes (Figure 6B). Surprisingly, pyruvate-plus-malate- and glutamate-plus-malate-driven respiration rates were below normal in well-fed Fgr null mice, and in the case of pyruvate-plus-malate this was accompanied by significantly lower ATP synthesis (Figure 6B). Lack of Fgr kinase thus appears to have a broader than anticipated effect on liver energy metabolism. No significant differences in citrate synthase activity were detected that could explain this decline in CI dependent respiration (Figure 6A). But although both the CI (NADH route) and the CII (FADH<sub>2</sub> route) are altered in Fgr<sup>-/-</sup> liver mitochondria, the balance of electron supply potential to the mETC from NADH and FADH<sub>2</sub> is shifted, increasing the CI/CII activity ratio (Figure 6C). Starving Fgr<sup>-/-</sup> mice overnight increased CI activity in liver mitochondria and reduced CII activity (Figures 6A, 6B, and S4), sharply increasing the CI/CII activity ratio, whereas the ratio in mitochondria from starved control mice decreased (Figure 6C). Feeding mitochondria with pyruvate or glutamate should generate intramitochondrial NADH, for delivery of electrons to CI. As expected, starvation-induced downregulation of pyruvate dehydrogenase activity reduced pyruvate-plus-malate-driven respiration and ATP synthesis in Fgr<sup>+/-</sup> and Fgr null mice (Figure 6B). In contrast, starvation only decreased respiration and ATP synthesis driven by glutamate-plus-malate in Fgr<sup>-/-</sup> mice, despite the higher maximal CI respiration (Figure 6B). These observations unexpectedly show that lack of Fgr kinase affects liver glutamate utilization. To avoid the complex pleiotropic effects of Fgr-kinase ablation in liver, we mimicked starvation in cultured embryonic fibroblasts (derived from Fgr<sup>+/-</sup> and Fgr<sup>-/-</sup> littermates) by overnight serum deprivation. Serum-deprived Fgr<sup>+/-</sup> fibroblasts showed the typical mitochondrial hyperfusion phenotype required to maintain ATP production upon nutrient deprivation (Gomes et al., 2011), whereas Fgr<sup>-/-</sup> mitochondria were fragmented (Figure 6D). Consistently, mitochondria from serum-deprived Fgr<sup>-/-</sup> cells showed higher processing of OPA1 (Figure 6E). Serum deprivation increased CII activity in Fgr<sup>+/-</sup> but not Fgr<sup>-/-</sup> cells (Figure 6F), recapitulating the lack of CII activation in Fgr<sup>-/-</sup> liver mitochondria from overnight-starved mice. To assess whether starvation responses were due only to Fgr-dependent CII phosphorylation, we analyzed the effects of serum deprivation in fibroblasts silenced for endogenous FpSDH and exogenously re-expressing WT or Y604F FpSDH. Serum deprivation triggered OPA-1 processing in Y604F cells (Figure 6E), mimicking the result in Fgr null fibroblasts, and only cells re-expressing WT FpSDH upregulated CII activity after serum deprivation (Figure 6G). The blunted CII activation in Y604F cells compromised cell survival after serum deprivation,



revealed by a higher proportion of apoptotic annexin V positive Y604F cells (Figures 6H and S5C).

### ROS/Fgr/CII Pathway in Reoxygenation-Induced Metabolic Reprogramming

A drop in O<sub>2</sub> availability triggers several adaptive mechanisms, including reduction in the activities and protein levels of OXPHOS components and in ROS production (Ali et al., 2012; Heather et al., 2012; Papandreou et al., 2006), and a notable accumulation of succinate (Casarano et al., 1976). However, sudden reoxygenation, as occurs in reperfusion after ischemia, is accompanied by a sharp increase in ROS production as the electron transport chain readapts to oxygen availability. To evaluate the role of ROS-mediated phosphorylation of FpSDH in this adaptation, we cultured Fgr<sup>+/-</sup> and Fgr<sup>-/-</sup> fibroblasts for 48 hr at 21% O<sub>2</sub> (normoxia), 1% O<sub>2</sub> (hypoxia), or 1% O<sub>2</sub> followed by reoxygenation at 21% O<sub>2</sub> for an additional 48 hr. Immunostaining and western blot analysis indicated that ROS-mediated mitochondrial biogenesis upon reoxygenation was impaired in Fgr<sup>-/-</sup> cells, with only Fgr<sup>+/-</sup> cells recovering normoxic mitochondrial numbers and shape (Figures 7A and 7B). In both genotypes, hypoxia reduced the amount of the mitochondrial proteins Tom20 and FpSDH (Figure 7B), consistent with the reported loss of mitochondria upon hypoxia (Kim et al., 2011). In the FpSDH re-expression model, hypoxia reduced mitochondrial content (measured as the FpSDH:actin and Tom20:actin ratios) in FpSDH-silenced fibroblasts re-expressing WT or Y604F FpSDH. As predicted, reoxygenation restored or increased mitochondrial protein content in cells re-expressing WT FpSDH, and this recovery was impaired in Y604F cells; however, Y604F cells did show partial mitochondrial recovery, differing from the more severe phenotype in Fgr<sup>-/-</sup> fibroblasts (Figure 7B). The reason for this difference is likely that germline lack of Fgr affects targets other than CII required for full recovery.

To test the metabolic effect of reoxygenation, we measured CII activity in Fgr<sup>+/-</sup> and Fgr<sup>-/-</sup> fibroblasts cultured with 25 mM glucose or the more physiological 10 mM. Hypoxia did not alter CII activity under any conditions, and reoxygenation increased CII activity only in Fgr<sup>+/-</sup> cells (Figures 7C and S5A). Likewise, in re-expression assays only WT FpSDH fibroblasts upregulated CII activity upon posthypoxia reoxygenation (Figures 7C and S5B). CII has been proposed to trigger apoptosis, depending on its attachment to the inner mitochondrial membrane (reviewed in Grimm, 2013). When detached, CII is not assembled as a holocomplex, and its succinate ubiquinol reductase (SQR or CII) activity, which involves coenzyme Q reduction, is decreased. However, succinate dehydrogenase (SDH) activity is unaltered, resulting in superoxide leakage that leads to apoptosis (Albayrak et al., 2003; Lemarie et al., 2011). SDH activity was slightly higher in Y604F-expressing cells in normoxia, but the proportion of apoptotic cells (annexin V positive) was unaffected. Upon reoxygenation, the balance between CII and SDH activity in cells re-expressing WT FpSDH shifted toward CII, whereas in cells re-expressing Y604F it shifted more toward SDH (Figure 7D). This was reflected in significantly more severe apoptosis after hypoxia/reoxygenation in Y604F-re-expressing cells, in which CII activation is blunted, but not in cells re-expressing WT FpSDH (Figures 7D and S5C).

BNGE revealed a hypoxia-induced generalized decrease in the content of assembled OXPHOS complexes and Tom20 (consistent with loss of mitochondrial proteins evident in Figure 7B), with no alteration in the proportion of CIII dedicated to each coenzyme Q pool (Figure 7E). Posthypoxia reoxygenation of WT re-expressing cells restored OXPHOS complexes and supercomplex assembly while maintaining these proportions. In contrast, reoxygenated Y604F-expressing cells restored the normoxic level of CIII dedicated to NADH but not the amount dedicated to FADH<sub>2</sub> (CIII+IV and free CIII) (Figure 7E), indicating a higher-than-normal dedication to processing electrons from the NADH-CoQ pool than from the FADH<sub>2</sub>-CoQ pool (Lapiente-Brun et al., 2013).

## DISCUSSION

The data presented here demonstrate that Tyr phosphorylation of FpSDH increases CII activity in vivo, and that this mechanism triggers remodeling of the mETC to reset its capacity for processing NADH- versus FADH<sub>2</sub>-derived electrons. This Tyr phosphorylation is H<sub>2</sub>O<sub>2</sub> mediated, is catalyzed by the Src-family kinase Fgr, and specifically targets Y604 in FpSDH. The finding that the catalytic subunit of CII can be phosphorylated is consistent with earlier observations (Bykova et al., 2003; Schulenberg et al., 2003). Moreover, our proposal that Fgr is the tyrosine kinase responsible for this phosphorylation concurs with a previous report demonstrating that Fgr, but not Lyn, is able to promote the in vitro phosphorylation of Y535 and Y596 of rat FpSDH, which correspond to Y543 and Y604 in mouse (Salvi et al., 2007).

H<sub>2</sub>O<sub>2</sub>-triggered activation of CII provides a mechanism for the association of increased CII activity and defective CI, observed in human patients and in a range of organisms from *Chlamydomonas reinhardtii* (Cardol et al., 2002) and *Rhodobacter capsulatus* (Dupuis et al., 1998) to mouse and humans (Esteitie et al., 2005; Fan et al., 2008; Majander et al., 1991). Another feature of ROS-driven activation of CII is that since hydrogen peroxide can permeate through cell membranes, any extramitochondrial source of H<sub>2</sub>O<sub>2</sub> can potentially activate CII, suggesting a mechanism to promote metabolic adaptation in response to signals that increase H<sub>2</sub>O<sub>2</sub>. Our results thus show that activation of CII by Fgr kinase in response to a primary wave of extramitochondrial ROS can trigger a secondary wave of ROS production as a consequence of CII activation. This pathway provides a mechanism for amplifying ROS signals within the cell. The increase in CII activity triggered by H<sub>2</sub>O<sub>2</sub> is a quick response mechanism, independent of gene expression and therefore not involving any increase in mitochondrial biogenesis or regulation through PGC-1 $\alpha$ , a common feature of mitochondrial disease (Moreno-Loshuertos et al., 2006; Acín-Pérez et al., 2009; Srivastava et al., 2009; Wenz et al., 2008).

FpSDH activity is regulated by acetylation on several lysine residues, and deacetylation mediated by sirtuin 3 (Cimen et al., 2010; Finley et al., 2011) increases CII activity independently of ROS. The regulation of CII acetylation is incompletely understood, but fuel availability is likely to play a part, probably in a complex tissue-specific pattern (Boyle et al., 2013; Finley et al., 2011). The convergence of multiple posttranslational modifications on the catalytic subunit of CII highlights the importance of fine-tuning CII activity to ensure correct cell metabolism. This role has remained unappreciated despite the considerable

knowledge accumulated on the function of the tricarboxylic acid (TCA) cycle. Two other TCA cycle enzymes, aconitase and KGDH, are known to be reversibly downregulated by physiological increases in ROS levels (Bulteau et al., 2003; Moreno-Loshuertos et al., 2006). Our current results show that CII should be included among the TCA cycle enzymes regulated by ROS.

Our results also show that Src-family kinase signaling operates within mitochondria, regulating the fundamental metabolic processes of the TCA cycle and oxidative phosphorylation. SFKs are implicated in a wide variety of signaling pathways, regulating cell growth, differentiation, cell shape, migration, and survival (Ingley, 2008; Parsons and Parsons, 2004). Moreover, the physiological role of SFK activation by H<sub>2</sub>O<sub>2</sub> has also been shown in another model system: the role of Lyn kinase as a redox sensor activated by H<sub>2</sub>O<sub>2</sub> in leukocyte wound attraction (Yoo et al., 2011). The role of reversible protein phosphorylation in mitochondria is an emerging field (Pagliarini and Dixon, 2006; Pagliarini et al., 2005), and several mitochondrial proteins have been proposed as Tyr phosphorylation targets (Augereau et al., 2005; Salvi et al., 2005, 2007; Tibaldi et al., 2008). Moreover, a number of SFKs have been proposed to localize in mitochondria (Pagliarini and Dixon, 2006; Tibaldi et al., 2008). One previous report has described the function of a mitochondrial SFK, demonstrating that c-Src enhances the activity of subunit II of cytochrome *c* oxidase, the terminal mETC enzyme (Miyazaki et al., 2003). Thus the TCA cycle and the mETC are both positively regulated by the action of SFKs: Fgr for the TCA cycle and c-Src for the mETC. By bringing the regulation of mitochondrial energetic metabolism within the remit of cellular signaling pathways regulated by SFKs, these findings have broad implications for the integration of cell adaptation and energy regulation. The results presented here highlight the importance of this regulation in the adaptation to loss of CI, regulation of CII activity during the metabolic switch upon activation of naive T cells, and metabolic adaptation of mitochondria to starvation and hypoxia/reoxygenation. CII has also been proposed to act as universal oxygen sensor (Baysal, 2006), suggesting that this pathway is also critically important in situations where cells need to respond rapidly to changes in fuel availability or O<sub>2</sub> concentration.

## EXPERIMENTAL PROCEDURES

Further details of methods are provided in Supplemental Experimental Procedures.

### Cell Culture

Cell lines were grown in Dulbecco's modified Eagle's medium (DMEM; GIBCOBRL) supplemented with 5% fetal bovine serum (FBS; GIBCOBRL). Fibroblasts from Fgr<sup>+/-</sup> and Fgr null mice were isolated from mouse ear and immortalized by transfection with pLOX-Ttag-iresTK (Addgene) and grown in DMEM with 10% FBS. FBalb/cJ and FG23-1 cells expressing MnSOD or HA-tagged mt-catalase and FBalb/cJ cells overexpressing different forms of FpSDH and silenced for the endogenous FpSDH were generated by viral infection using polybrene (8 µg/ml).

### **Isolation of Mitochondria and Mitochondrial Fractions from Mouse Liver and Cell Lines**

Mitochondria were isolated from cell lines as described (Schägger and von Jagow, 1991) with some modifications, and from mouse liver as described (Fernández-Vizarra et al., 2002).

### **OXPHOS Function and Enzyme Activities**

O<sub>2</sub> consumption was measured in mouse liver mitochondria (100 µg) as described (Hofhaus et al., 1996). ATP synthesis in isolated mitochondria (15–25 µg mitochondrial protein) was measured using a kinetic luminescence assay (Vives-Bauza et al., 2007). Mitochondrial fractions were prepared and the activities of individual complexes measured spectrophotometrically (Birch-Machin and Turnbull, 2001). Catalase activity was measured in total cell lysates (300 µg) (Moreno-Loshuertos et al., 2006). Total SOD and MnSOD (KCN insensitive) activities were assessed in total cell lysates (50 mg) using the SOD Assay Kit (Sigma-Aldrich).

### **Blue-Native Gel Electrophoresis**

Cell-culture-derived mitochondria (50–75 µg) were separated on 5%–13% gradient blue native gels (Schägger and von Jagow, 1991).

### **Isoelectric Focusing and 2D SDS-PAGE**

Mitochondrial preparations (100 µg) were processed with by Ready Prep 2D Cleanup (BioRad) and applied to pH 3–10 or pH 4–7 IPG strips (BioRad) and incubated overnight at room temperature. Isoelectric focusing and second dimension SDS-PAGE were run under standard conditions.

### **Phosphoprotein Enrichment**

Phosphoprotein-enriched mitochondrial fractions were isolated on phosphoprotein enrichment columns (QIAGEN).

### **Immunocapture**

Complex II and FpSDH were immunocaptured from mitochondria isolated from either cells or mouse liver. For details see Supplemental Experimental Procedures.

### **In Vitro Fgr Phosphorylation**

Recombinant Fgr kinase (Abnova) was used to phosphorylate complex II immunocaptured from cells expressing WT and Y604F FpSDH (Salvi et al., 2007).

### **Immunological Analysis**

Antibodies used in this study are listed in Supplemental Experimental Procedures.

### **H<sub>2</sub>O<sub>2</sub> Production**

H<sub>2</sub>O<sub>2</sub> production was measured in cultured cells grown in the absence or presence of 5 mM NAC for 7 days (Moreno-Loshuertos et al., 2006).

## Isolation and Stimulation of Naive CD4<sup>+</sup> and CD8<sup>+</sup> T Cells

CD4<sup>+</sup> and CD8<sup>+</sup> T cells were purified from splenic cell suspensions obtained from 8- to 10-week-old Fgr<sup>+/-</sup> and Fgr<sup>-/-</sup> male mice. To obtain differentiated T lymphoblasts, naive T cells were cultured with concanavalin A (Sigma) and human recombinant IL-2 (50 U/ml, Glaxo).

## Flow Cytometry ROS Production Determination

ROS production in naive or activated T lymphocytes and in cells before and after overexpression of detoxifying enzymes was assessed by flow cytometry of 2,7-DCFH<sub>2</sub>-DA or MitoSOX staining (Kami ski et al., 2012).

## Determination of Apoptosis in Cultured Cells

Apoptosis was monitored by flow cytometry detection of annexin V and propidium iodide (PI) staining in 10<sup>6</sup> cells resuspended in 10 mM HEPES/NaOH (pH 7.4), 140 mM NaCl, 2.5 mM CaCl<sub>2</sub>.

## Mouse Strains

Animal studies were approved by the local ethics committee. Analyses were performed in 8- to 11-week-old male mice.

## In Silico Analysis

See Supplemental Experimental Procedures.

## Statistical Analysis

Comparisons between groups were made by one-way ANOVA. Pairwise comparisons were made by Fisher's PLSD post hoc test. Differences were considered statistically significant at  $p < 0.05$ . Data were analyzed with StatView (Adept Scientific, UK).

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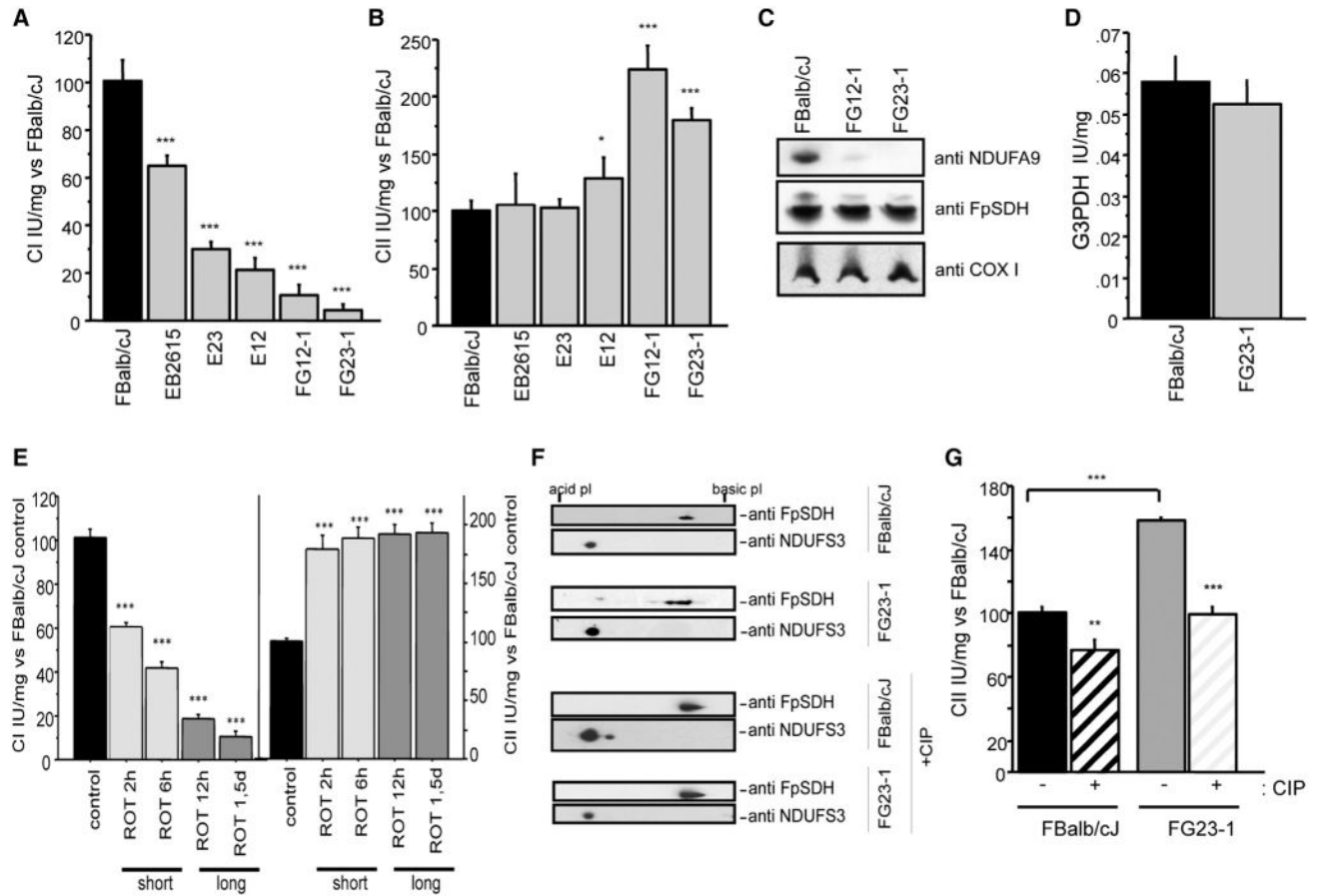
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**Figure 1. Phosphorylation of FpSDH Increases CII Activity in CI-Deficient Cells**

(A) Fibroblast lines with mutations in ND6 show decreased CI activity in proportion to the mutation load. Activity is expressed as the percentage of activity in FBalb/cJ cells (WT, no mutation; n = 12): EB2615 (70%; n = 6); E23 (34%; n = 6); E12 (20%; n = 9); FG12-1 (5%; n = 9); and FG23-1 (2%; n = 9).

(B) CII activity (CoQ reduction) in cell lines with high loads of the iC13887 ND6 mutation, expressed as the percentage of activity in FBalb/cJ cells (n = 18): EB2615 (n = 9); E23 (9 n = 9); E12 (n = 12); FG12-1 (n = 9); and FG23-1 (n = 20).

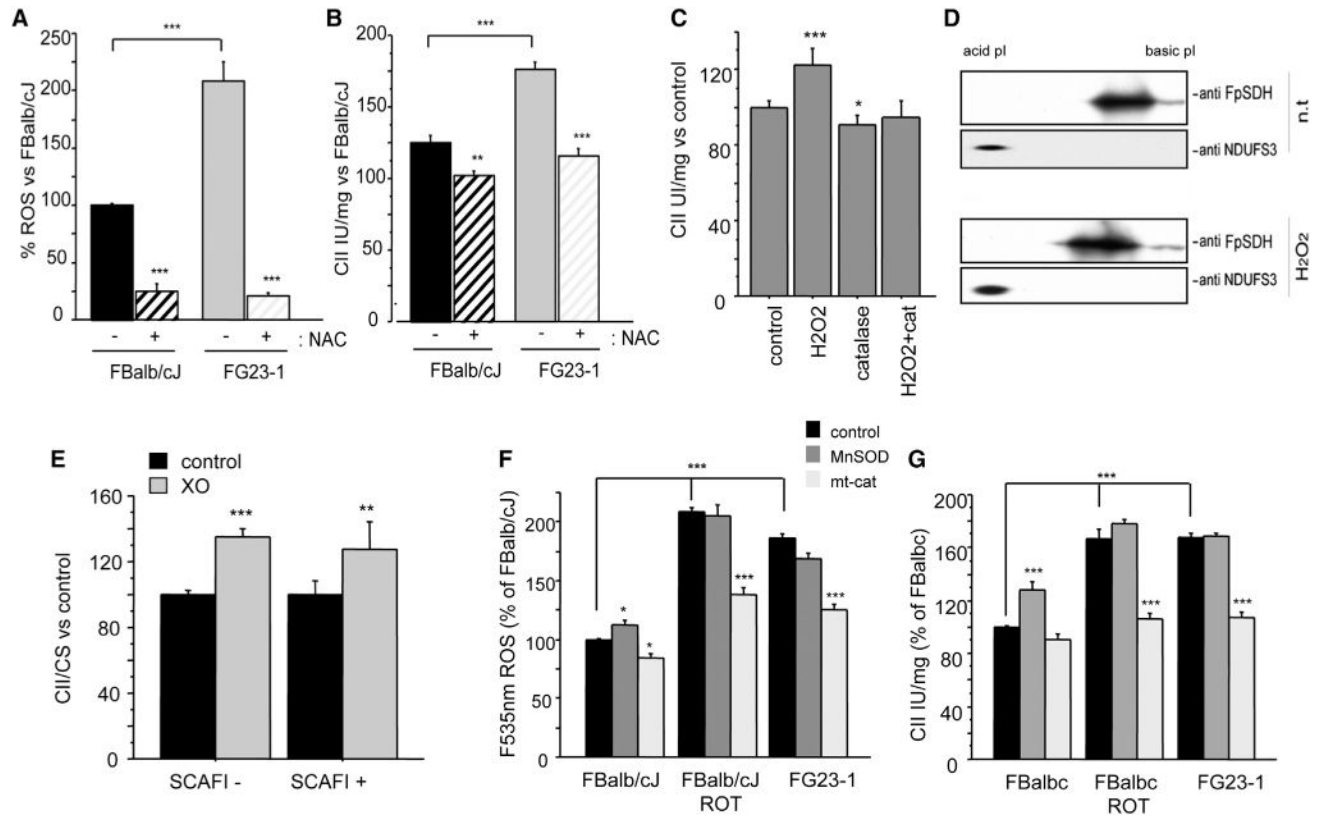
(C) Western blot after BNGE, detecting subunits of CI (NDUFA9), CII (FpSDH), and CIV (COX I).

(D) G3PDH activity in FG23-1 and FBalb/cJ (n = 6).

(E) Activities of CI (left) and CII (right) in a time course rotenone treatment of FBalb/cJ (200 nM) as a percentage of activity in untreated cells (n = 6). Short: 2 and 6 hr treatment; long: 12 hr and 1.5 day treatment.

(F) FpSDH immunoblot after 2D IEF/SDS-PAGE (IEF strip pH 3–10) of control and calf intestine phosphatase (CIP)-treated mitochondrial protein (100 µg) from FBalb/cJ and FG23-1 cells. The acidic shift of FpSDH spots in FG23-1 cells is blocked by CIP. The CI subunit NDUFS3 was used to align and compare the blots.

(G) CII activity in CIP-treated mitochondria from FBalb/cJ and FG23-1 cells (n = 6). Data are presented as the percentage of activity in untreated FBalb/cJ. \*\*p < 0.001; \*\*\*p < 0.0001. Enzyme activities were determined spectrophotometrically. All data are presented as mean ± SD. See also Figure S1.



### Figure 2. FpSDH Phosphorylation and CII Activity Are ROS Dependent

(A) ROS (H<sub>2</sub>O<sub>2</sub>) levels measured in nontreated (n.t.) FG23-1 and FBalb/cJ cells and in cells treated with NAC (5 mM, 7 days) (control, n = 35; NAC, n = 12).

(B) Effect of NAC treatment on CII activity in FBalb/cJ and in FG23-1 cells (untreated, n = 15; NAC, n = 4). Data in (A) and (B) are presented as the percentage of nontreated FBalb/cJ cells.

(C) CII activity in isolated mouse liver mitochondria treated for 10 min with H<sub>2</sub>O<sub>2</sub> (50 μM), catalase (25 U/ml), or both (n = 4).

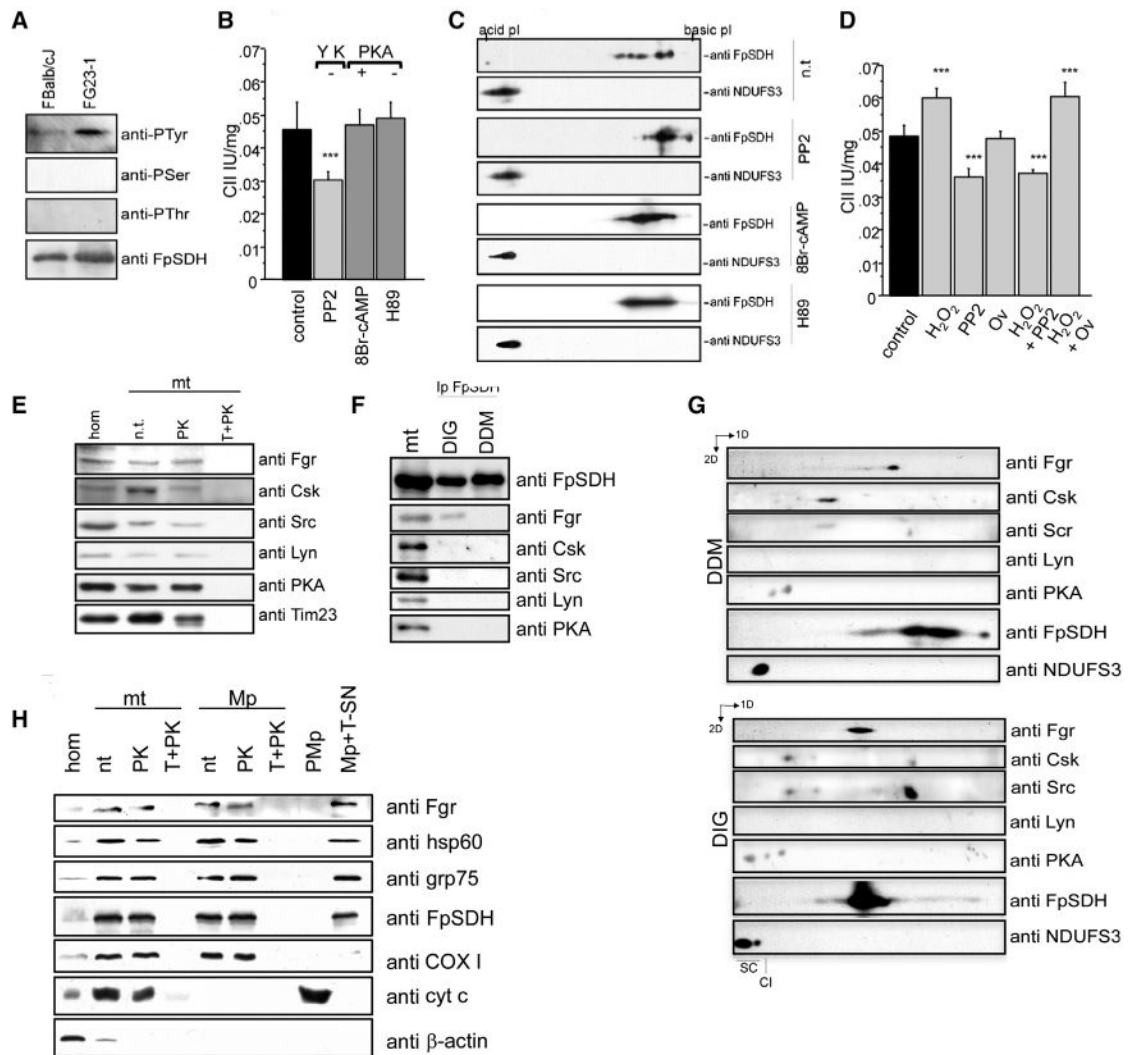
(D) 2D IEF/SDS-PAGE (IEF strips pH 4–7) Western analysis of FpSDH from H<sub>2</sub>O<sub>2</sub>-treated mouse liver mitochondria.

(E) CII activity in mitochondria isolated from SCAFI- (left) and SCAFI+ (right) fibroblasts treated with the ROS generator xanthine oxidase (XO) (n = 4). Data are presented as the percentage of activity in nontreated fibroblasts.

(F) H<sub>2</sub>O<sub>2</sub> production in FG23-1 and rotenone-treated FBalb/cJ fibroblasts is prevented by expression of mitochondrially targeted catalase (mt-cat, n = 5).

(G) Elevated CII activity in FG23-1 and rotenone-treated FBalb/cJ cells is prevented by expression of mt-cat but not MnSOD (n = 3). \*p < 0.01; \*\*p < 0.001; \*\*\*p < 0.0001.

All data are presented as mean ± SD. See also Figure S2.



**Figure 3. The Src-Type Kinase Fgr Phosphorylates CII**

(A) Immunoblot phosphoprotein analysis in FpSDH immunoprecipitates from FBalb/cJ and FG23-1 cells.

(B) Complex II activity in mouse liver mitochondria incubated with the Tyr kinase inhibitor PP2, the PKA agonist 8Br-cAMP, or the PKA antagonist H89 (n = 5).

(C) 2D IEF/SDS-PAGE (3–10 IEF strips) Western analysis of mouse mitochondria treated with PP2, 8Br-cAMP, or H89. PP2 induces a basic shift in FpSDH compared with nontreated mitochondria (n.t.). Modulation of PKA activity had no effect on FpSDH phosphorylation. NDUFS3 (CI) was detected to align and compare blots.

(D) H<sub>2</sub>O<sub>2</sub>-induced activation of complex II is blocked by the Tyr kinase inhibitor PP2, but not by the Tyr phosphatase inhibitor orthovanadate (Ov) (n = 6).

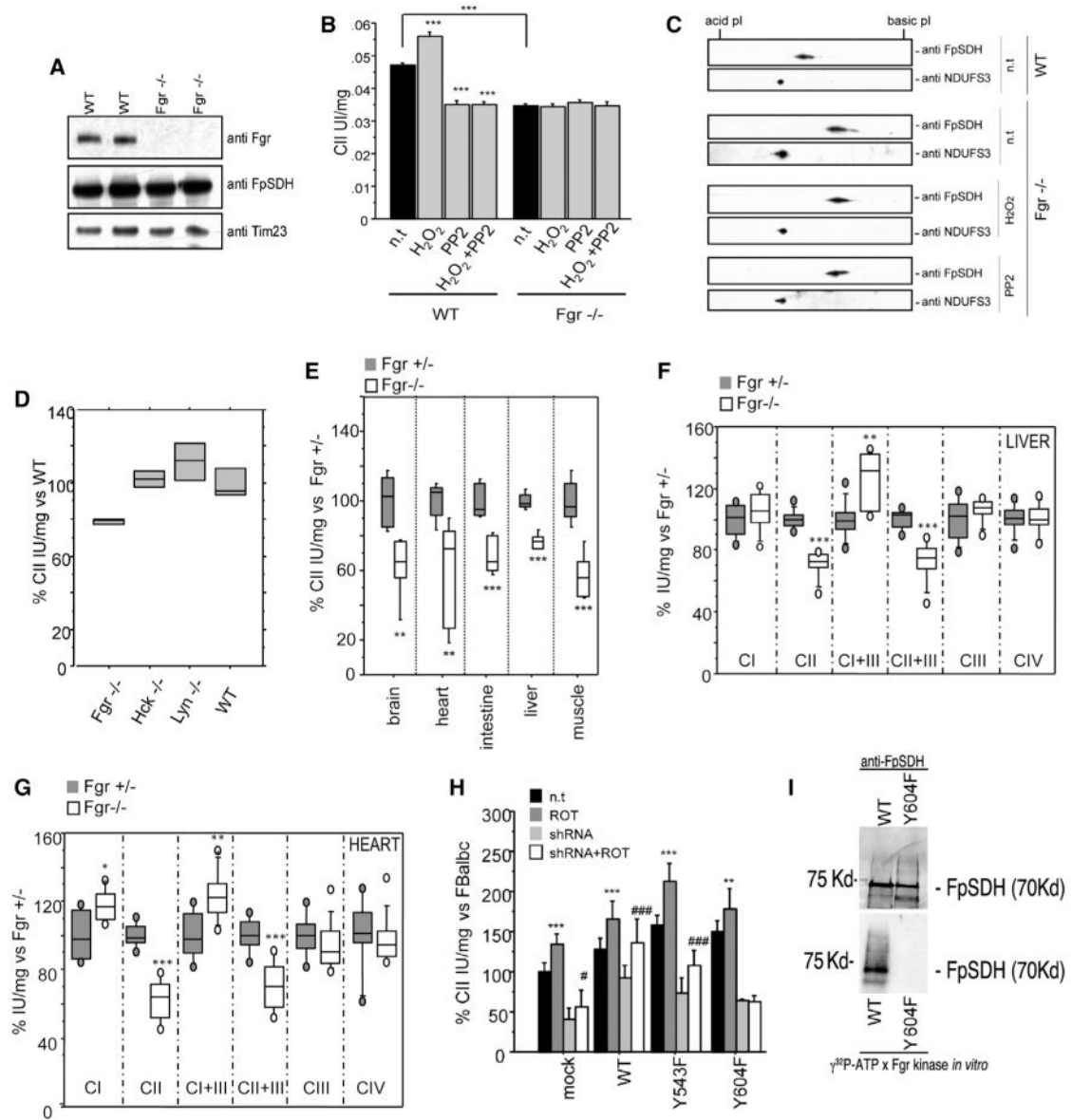
(E) Intact mouse mitochondria (mt) protect a portion of Fgr, Csk, Src, Lyn, and PKA kinases from digestion by proteinase K: hom, cell homogenate; n.t., non-treated; PK, proteinase K; T+PK, proteinase K treatment of mitochondria solubilized with Triton X-100. Tim23 was used as a marker of intact mitochondria.

(F) Anti-FpSDH specifically coimmunoprecipitates Fgr kinase from mitochondria solubilized with digitonin (DIG), which preserves intact supercomplexes, but not dodecyl maltoside (DDM), which resolves individual respiratory complexes. mt, fully solubilized mitochondria.

(G) 2D BNGE/SDS-PAGE of DDM- and DIG-solubilized mitochondria, showing comigration of Fgr and FpSDH in DIG-treated mitochondria. Note comigration of Csk and Src in both preparations.

(H) Mitochondrial Fgr is located in the matrix. Fractionation experiment showing protection of Fgr from PK digestion in intact mitochondria (mt) and mitoplasts (Mp) but not in either fraction solubilized with Triton X-100 (T+PK). Fgr was present in the supernatant fraction of Triton X-100-solubilized mitoplasts (Mp+T-SN). P-Mp, post-mitoplast fraction. Markers used: hsp60 and Grp75, mitochondrial matrix; FpSDH, association with inner membrane; COXI, inner membrane; cyt c, intermembrane space;  $\beta$ -actin, cytoplasm.

All data are presented as mean  $\pm$  SD.



#### Figure 4. CII Is Regulated by Fgr Phosphorylation at Tyr604

(A) SDS-PAGE western analysis of Fgr tyrosine kinase in liver mitochondrial protein preparations from control or  $Fgr^{-/-}$  mice.

(B) CII activity in isolated liver mitochondria from WT or  $Fgr^{-/-}$  mice in the presence of  $H_2O_2$ , PP2, or both. Data are presented as mean  $\pm$  SD.

(C) 2D IEF/SDS-PAGE (IEF strips pH 4–7) Western analysis of FpSDH from WT liver mitochondria and from nontreated and  $H_2O_2$ - or PP2-treated  $Fgr^{-/-}$  liver mitochondria. NDUFS3 (CI) is used to align and compare blots.

(D) Complex II activity in liver mitochondria from different Tyr kinase null mice.

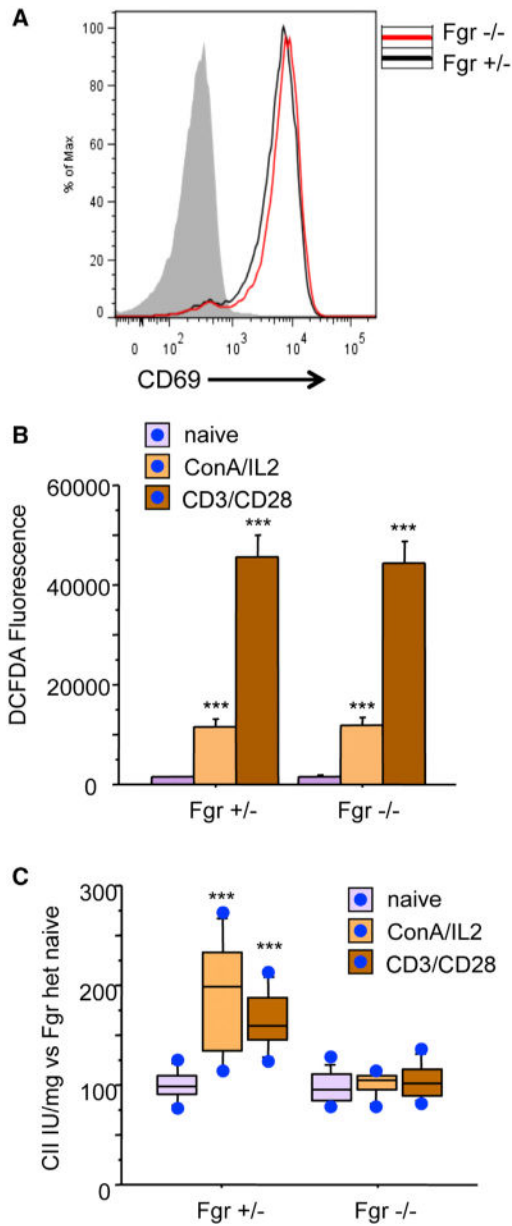
(E) Complex II activity in tissue homogenates from  $Fgr^{+/-}$  and  $Fgr^{-/-}$  null mice ( $n = 4$ ).

(F and G) Individual and combined mitochondrial complex activities in isolated mitochondria from liver (F) and heart (G) ( $n = 4$ ). In (D)–(G), lines extending from the boxes indicate the variability outside the upper and lower quartiles.

(H) CII activity in FBalb/cJ cells silenced for endogenous FpSDH and re-expressing WT and mutant FpSDH variants (n = 4). ROT, rotenone (200 nM); shRNA, silencing of endogenous FpSDH. Data are presented as the percentage of activity in nontreated, mock-infected FBalb/cJ (mean  $\pm$  SD). Statistical significance versus nontreated: \*\*p < 0.001; \*\*\*p < 0.0001. Statistical significance versus shRNA: #, p < 0.01; ###, p < 0.0001.

(I) Fgr in vitro phosphorylation (bottom) and FpSDH immunodetection (top) of immunocaptured complex II from FBalb/cJ cells expressing WT FpSDH (left line) or the Y604F mutant (right line). See also Figure S3.



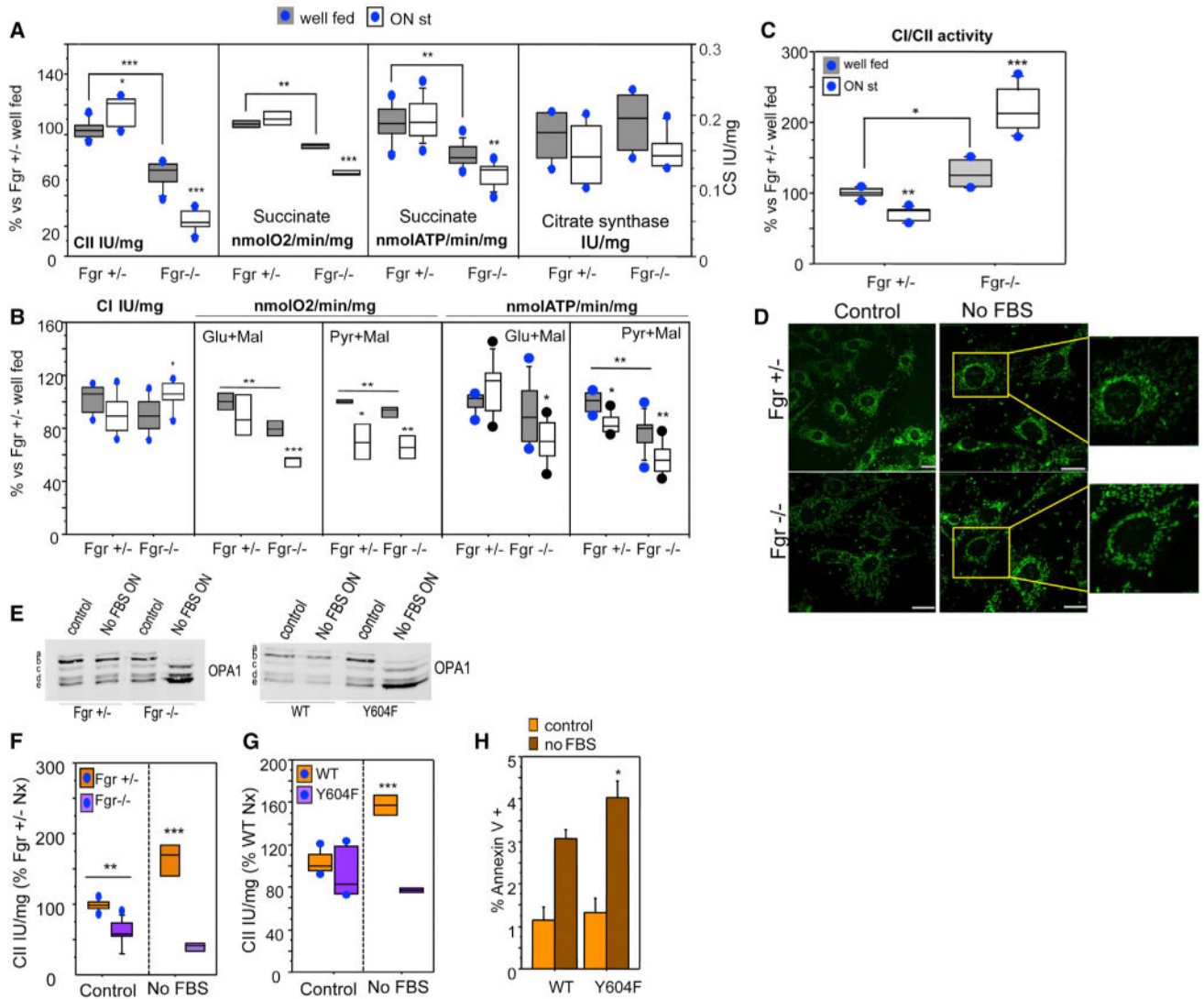


**Figure 5. T Cell Activation and CII Activity Response**

(A) Flow cytometry analysis of T cell activation. Naive T cells from  $Fgr^{+/-}$  and  $Fgr^{-/-}$  mice were activated with anti-CD3/CD28, and activation was assessed by detection of CD69. Gray line: negative control. The chart shows data from one representative experiment of four.

(B) Flow cytometry analysis of ROS production (2,7-DCFH<sub>2</sub>-DA) in resting T cells and upon CD3/CD28 or ConA activation (n = 3). Data are presented as mean  $\pm$  SD.

(C) CII activity measured in resting and activated T cells (n = 4). Data are presented as the percentage of activity compared to resting T cells. Lines extending from the boxes indicate the variability outside the upper and lower quartiles.



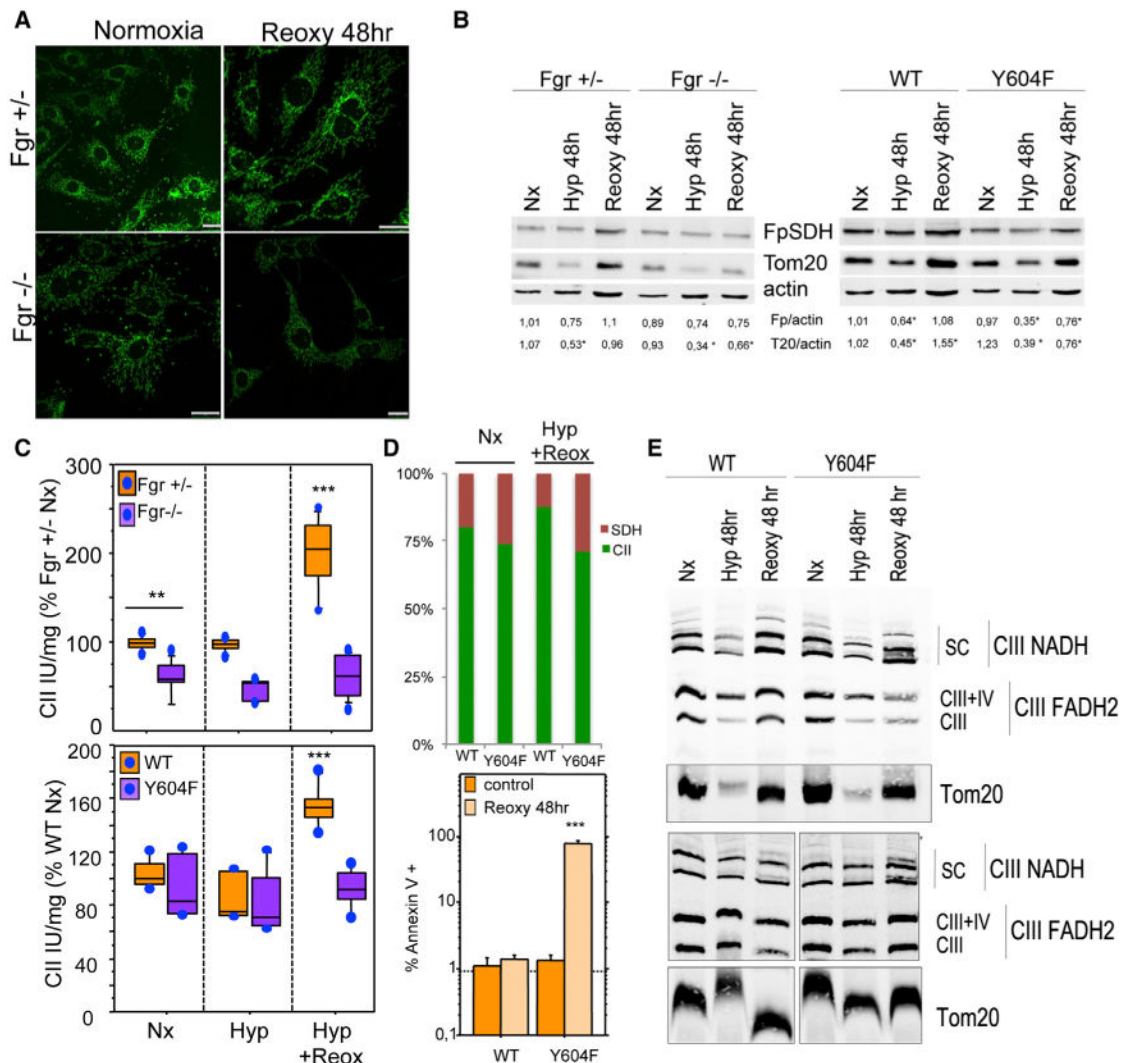
**Figure 6. Mitochondria Lacking Fgr or Expressing Y604F FpSDH Respond Abnormally to Starvation and Serum Deprivation**

(A) Succinate-driven OXPHOS function in mouse liver mitochondria from  $Fgr^{+/-}$  and  $Fgr^{-/-}$  mice fed a normal diet (well fed) or starved overnight (ON st,  $n = 4$ ). CII activity (left), succinate-driven respiration (center left), succinate-driven ATP synthesis (center right), and citrate synthase activity (CS, right).

(B) NADH-driven OXPHOS function in mouse liver mitochondria from  $Fgr^{+/-}$  and  $Fgr^{-/-}$  mice fed a normal diet or starved overnight ( $n = 4$ ). CI activity (left), glutamate-driven respiration (center left), pyruvate-driven respiration (center), glutamate-driven ATP synthesis (center right), and pyruvate-driven ATP synthesis (right) are shown.

(C) Rate of use of NADH/FADH reducing equivalents in mouse liver mitochondria from  $Fgr^{+/-}$  and  $Fgr^{-/-}$  mice ( $n = 4$ ). Data in (A)–(C) are presented as the percentage of values obtained in well-fed  $Fgr^{+/-}$  mice. Lines extending from the boxes indicate the variability outside the upper and lower quartiles.

- (D) Immunostaining of mitochondria (Tom20, green) in  $Fgr^{+/-}$  and  $Fgr^{-/-}$  fibroblasts cultured with serum or without serum overnight (No FBS ON).
- (E) Immunoblot showing the migration of OPA1 in lysates from control and serum-deprived  $Fgr^{+/-}$  and  $Fgr^{-/-}$  cells (left) or control and serum-deprived FpSDH-silenced fibroblasts re-expressing WT or Y604F FpSDH (right).
- (F) CII activity in  $Fgr^{+/-}$  and  $Fgr^{-/-}$  fibroblasts grown in 10 mM glucose in the indicated conditions (n = 5). Data are presented as the percentage activity in  $Fgr^{+/-}$  cells grown in normoxia; lines extending from the boxes indicate the variability outside the upper and lower quartiles.
- (G) CII activity in FpSDH-silenced fibroblasts re-expressing WT or Y604F FpSDH, cultured at 5 mM glucose under the indicated oxygenation conditions (n = 5). Data are presented as the percentage of activity in WT-expressing cells grown in normoxia; lines extending from the boxes indicate the variability outside the upper and lower quartiles.
- (H) Apoptotic events in FpSDH-silenced fibroblasts re-expressing WT or Y604F FpSDH and grown with or without serum. Apoptosis was determined by flow cytometry as the percentage of cells positive for annexin V and propidium iodide (n = 3). Data are presented as mean  $\pm$  SD. \*p < 0.01; \*\*p < 0.001; \*\*\*p < 0.0001.  
See also Figures S4 and S5.



### Figure 7. Mitochondria Lacking Fgr or Expressing Y604F FpSDH Respond Abnormally to Hypoxia-Reoxygenation

(A) Immunostaining of mitochondria (Tom20, green) in Fgr<sup>+/-</sup> and Fgr<sup>-/-</sup> fibroblasts cultured in normoxia (21% O<sub>2</sub>) or hypoxia (1% O<sub>2</sub>) for 48 hr followed by 48 hr normoxia (Reoxy 48 hr).

(B) Left: Immunoblot analysis of Fgr<sup>+/-</sup> and Fgr<sup>-/-</sup> cells cultured under normoxia (Nx), hypoxia for 48 hr (Hyp 48 hr), or hypoxia followed by normoxia (Reoxy 48 hr). Right: Immunoblot analysis of FpSDH-silenced fibroblasts re-expressing WT or Y604F FpSDH and cultured under the indicated oxygenation conditions. Numbers beneath blots show FpSDH (Fp):actin and Tom20 (T20):actin ratios (n = 4).

(C) Top: CII activity in Fgr<sup>+/-</sup> and Fgr<sup>-/-</sup> fibroblasts grown in 10 mM glucose under the indicated conditions (n = 5). Bottom: FpSDH-silenced fibroblasts re-expressing WT or Y604F FpSDH, cultured with 5 mM glucose under the indicated oxygenation conditions (n = 5). Data are presented as the percentage activity in Fgr<sup>+/-</sup> cells or WT-FpSDH-re-expressing cells grown in normoxia; lines extending from the boxes indicate the variability outside the upper and lower quartiles.

(D) Top: Relative activities of CII (CoQ reduction) and SDH in WT- and Y604F-re-expressing FpSDH-silenced cells grown in normoxia or through a hypoxia/reoxygenation cycle. For each cell line and condition, 100% = the sum of CII and SDH activities; absolute SDH activity did not differ between cell lines and conditions. Bottom: Apoptotic events in FpSDH-silenced fibroblasts re-expressing WT or Y604F FpSDH and grown in normoxia or through a hypoxia/reoxygenation cycle. Apoptosis was determined by flow cytometry as the percentage of annexin V- and PI-positive cells (n = 3). Data are presented as mean  $\pm$  SD.

(E) BNAGE of FpSDH-silenced fibroblasts re-expressing WT or Y604F FpSDH cultured at 5 mM glucose under the indicated oxygenation conditions; the blot reveals the distribution of CIII (anti-core 1 immunodetection) among the different forms of free complex and supercomplexes. The outer membrane protein Tom20 is used as a mitochondrial protein loading control. Upper and lower panels are taken from two independent experiments. Note that upon reoxygenation the amount of CIII super assembled with CI is abnormally elevated in the Y604F mutant. See also Figure S5.