NF-κB pathway controls mitochondrial dynamics

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The Optic atrophy 1 protein (OPA1) is a key element in the dynamics and morphology of mitochondria. We demonstrated that the absence of IκB kinase-α, which is a key element of the nonclassical NF-κB pathway, has an impact on the mitochondrial network morphology and OPA1 expression. In contrast, the absence of NF-κB essential modulator (NEMO) or IκB kinase-β, both of which are essential for the canonical NF-κB pathway, has no impact on mitochondrial dynamics. Whereas Parkin has been reported to positively regulate the expression of OPA1 through NEMO, herein we found that PARK2 overexpression did not modify the expression of OPA1. PARK2 expression reduced the levels of Bax, and it prevented stress-induced cell death only in Bak-deficient mouse embryonic fibroblast cells. Collectively, our results point out a role of the nonclassical NF-κB pathway in the regulation of mitochondrial dynamics and OPA1 expression.

Results and discussion

Overexpression of Park2 is not associated with higher levels of OPA1. On the basis of the observation that NEMO-deficient MEF cells display a mitochondrial network morphology without fragmentation (Figure 1a) and exhibit the same levels of OPA1 isoforms in comparison with wild-type (WT) MEF cells (Figure 1b), we analyzed the mitochondrial network in Parkin-deficient MEF cells (Figure 1c). Confocal microscopy revealed a decrease in the size of mitochondria, although maintaining similar numbers per cell when compared with WT MEF cells (Figures 1c–e). Previous reports from fibroblasts of patients with PARK2 mutations showed similar degrees of branching under basal culturing conditions.1,2 In the presence of protonophore carbonyl cyanide 3-chlorophenylhydrazone (CCCP), mitochondrial network is fragmented in Parkin-deficient cells (Figure 1d). Analysis of OPA1 expression in Parkin-deficient cells revealed a decrease of ~24% in the amount of OPA1 compared with WT cells (Figure 1f), in agreement with Müller–Rischart et al.13

To further examine the role of Parkin in OPA1 regulation, we cotransfected WT, Parkin-, and NEMO-deficient MEF cells with an expression plasmid for PARK2 and a GFP plasmid to monitor the percentage of transfected cells. We achieved a cell transfection efficiency of ~40%, as quantified by flow embryonic fibroblast (MEF) cells display a normal mitochondrial network morphology, we decided to re-examine the role of Parkin in regulating OPA1 expression through the NF-κB signaling pathway.

Mitochondria perform multiple functions that are critical to the maintenance of cellular homeostasis. Mitochondrial dysfunctions have been linked to the development of degenerative diseases and aging. Damaged mitochondria are removed by mitophagy, a process partially regulated by the PARK2-encoded E3 ubiquitin ligase (Parkin) in a PTEN-induced putative protein kinase 1 (PINK1)-dependent manner.1–4 During mitophagy, the phosphorylation of mitofusin (Mfn) 2 by PINK1 has been suggested to induce the recruitment of Parkin to the mitochondria in cardiomyocytes.5 However, previous groups have shown that that Mfn 1 and 2 are dispensable for Parkin-dependent mitophagy in fibroblasts, whereas the Parkin-dependent degradation of these proteins may impair fusion of damaged mitochondria with the healthy network.6–8 PINK1 and Parkin thus act as a quality control machinery on the outer mitochondrial membrane (OMM) to preserve mitochondrial integrity through the ubiquitination of OMM proteins.9,10 Moreover, through its E3 ubiquitin ligase activity,11,12 Parkin was reported to bind to the linear ubiquitin chain assembly complex (LUBAC) and to increase the ubiquitination of NF-κB essential modulator (NEMO),13 a component of the classical NF-κB signaling pathway.14 Müller–Rischart et al. also proposed that Parkin positively regulates the expression of the mitochondrial guanosine triphosphatase Optic atrophy 1 protein (OPA1) through linear ubiquitination of NEMO.13 OPA1 is a regulator of mitochondrial inner membrane fusion and cristae remodeling.15–17 A defect in OPA1 expression is associated with mitochondrial network fragmentation and enhanced sensitivity of the cells to undergo apoptosis by promoting cytochrome c release from the mitochondria.18–20 Because NEMO-deficient mouse

Abbreviations: CCCP, carbonyl cyanide 3-chlorophenylhydrazone; IKKα, IκB kinase-α; IKKβ, IκB kinase-β; LUBAC, linear ubiquitin chain assembly complex; Mfn, mitofusin; OMM, outer mitochondrial membrane; MEF, mouse embryonic fibroblasts; ΔΨm, mitochondrial membrane potential; NEMO, NF-κB essential modulator; OPA1, Optic atrophy 1; PI, propidium iodide; PINK1, PTEN-induced putative protein kinase 1; STS, Staurosporine

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cytometry at 24 h (Figure 2a). Parkin expression was assessed by western blotting, and actin was used as a control for loading. Analysis of OPA1 expression in PARK2-transfected cells did not reveal an increase in the amount of OPA1 in WT and Parkin−/− (Figures 2b and c). In NEMO-deficient MEFs such as WT or Parkin-deficient MEFs, the overexpression of Parkin following transfection at a dose of 1 μg of plasmid rather resulted in a decrease, although not significant, in the amount of OPA1 (Figure 2d). Because Müller–Rischart et al.13 reported a significant upregulation in protein levels of OPA1, both under basal conditions and in the presence of CCCP, we treated WT- and NEMO-deficient MEF cells with CCCP (Figure 3). CCCP induced the degradation of the long isoform of OPA1 (Figures 3a and b), as previously described.15,16,18–20 In the presence of Parkin, no difference in the levels of OPA1 was observed either at 3 or 24 h post-
This result is consistent with a previous report from Lutz et al. showing that the absence of parkin does not alter the proteolytic processing of OPA1 in SH-SY5Y cells. It has been proposed that mitochondrial fission related to Drp1 induces mitochondrial network fragmentation in Parkin-deficient cells. Furthermore, in both cell lines, CCCP induced mitochondrial network fragmentation (Figures 3c and d) and dissipation of mitochondrial membrane potential (ΔΨm loss) (Figures 3e and f). Moreover, in the presence of Parkin, the mitochondrial membrane potential of CCCP-treated MEF is even reduced, suggesting a removal of damaged mitochondria (Figures 3c and f), which is consistent with previous reports on the functional role of Parkin on mitochondria.

IKKa-deficient MEF cells display lower levels of OPA1 and a fragmented mitochondrial network. The NF-κB/Rel family is composed of several members, including RelA (p65), RelB, c-Rel, NF-κB1 p50, and NF-κB2 p52. NF-κB proteins are regulated through their interaction with specific inhibitors, the IκB proteins (IκBα, IκBβ, or IκBε), which keep them in an inactive form in the cytoplasm of cells. The classical NF-κB pathway involves a huge complex containing NEMO and IκB kinase-β (IKKβ), but not IκB kinase-α (IKKα). IKKα and IKKβ phosphorylate IκB proteins, which trigger their...

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**Figure 2** OPA1 protein expression in Parkin-transfected MEF cells. (a) Efficiency of cell transfection. Cells were cotransfected with Parkin and GFP (1 μg) or mock. Green fluorescent intensity was determined by flow cytometry after 24 h (b–d). Profile of OPA1 expression in WT, Parkin−/−, and NEMO−/− MEFs (left panels), and quantification of the amount of OPA1 isoforms (right panels). The values indicate the relative levels of the long isoforms with respect to the baseline (value 1), in the absence (none) or after 24 h of transfection with either a GFP or PARK2 vector at the dose of 0.5 and 1 μg. HSP60 was used as a loading control to normalize protein content. Immunoblots were also probed with specific Parkin antibodies. Actin was used as a loading control. Four experiments were performed.
proteasome-dependent degradation. An alternative nonclassical pathway depends on IKKα and NF-κB-inducing kinase. Our results showed that, in contrast to NEMO-deficient cells (Figure 1), the amount of long and short isoforms of OPA1 was drastically decreased in IKKα- and IKKαβ-deficient MEFs but not in IKKβ-deficient MEFs (Figures 4a and b). In addition, IKKα- and IKKαβ-deficient MEFs displayed a fragmented mitochondrial network (Figures 4c and d), in the absence of OMM permeabilization, as judged by cytochrome c release (Figures 4c and d). Furthermore, we demonstrated in both IKKα- and IKKαβ-deficient MEFs that mitochondrial network is recovered (Figures 5a and b) and OPA1 expression (Figures 5c and d) after the overexpression of IKKα but not after IKKβ or NEMO expressions.

To gain further insight into the role of Parkin in the regulation of OPA1 expression through the classical and nonclassical NF-κB pathways, we analyzed the impact of PARK2 overexpression in MEFs deficient for either IKKα, IKKβ, or both.14 Our western blot results showed that, in none of the MEFs tested, transfection with the PARK2 vector, at a dose of 0.5 or 1 μg, did significantly increase the amount of OPA1 normalized to mitochondrial HSP60 levels (Figures 6a–c). Furthermore, we did not observe major changes in OPA1 mRNA expression by RT-PCR after transfection with the PARK2 vector compared

Figure 3  Park2 overexpression in stressed MEF cells. (a) Profile of OPA1 expression in WT and (b) NEMO−/− MEFs. Cells were transfected with Park2 at a dose of 1 μg. OPA1 expression was assessed at 3 and 24 h in the absence or presence of CCCP. Hsp60 and actin were used as loading controls to normalize protein content. (c) WT and (d) NEMO−/− MEF cells were stained with specific antibodies directed against Hsp60 (in red). Mitochondrial network was analyzed after overexpression of Park2 in the absence or presence of CCCP after 24 h. (e) Mitochondrial membrane potential of WT MEF cells assessed by flow cytometry in the absence or presence of CCCP. (f) Histograms of ΔΨm from WT and NEMO−/− MEF cells (mean fluorescence intensity) of transfected Park2 cells in the absence or presence of CCCP. Mean ± SD of three independent experiments; *P<0.05.
with the GFP vector alone in all the MEFs tested (Figure 6d). Therefore, by using different cell lines, we excluded an impact of cell culture conditions on the absence of Parkin effect on OPA1 expression. Thus, overexpression of Parkin appeared to have a minor role in regulating OPA1 expression in MEF cells. This is in agreement with a previous report, in which the overexpression of Parkin in Hela cells was not associated with an increase in the amount of OPA1.

Overexpression of Park2 reduces Bax expression and cell death. To delineate the role of Parkin in apoptosis, we analyzed the effect of its overproduction in the different MEF lines on the expression of two proapoptotic molecules with a crucial role in the control of mitochondria permeabilization, Bax, and Bak. Our results revealed that, at the dose of 1 μg of PARK2 vector, the amount of Bax normalized to actin was lower in WT, Parkin-deficient, and IKKαβ-deficient MEFs than
in nontransfected cells or in cells transfected with a GFP vector (Figures 7a and c). We did not observe differences in the amount of Bak in any of the cell lines following overexpression of Parkin.

Finally, we assessed whether the ectopic expression of Parkin protected WT or PARK2-deficient MEFs from cell death induced by staurosporine (STS) (Figure 8). Ectopic expression of Parkin in MEF cells had no protective effect against STS-mediated cell death in WT (Figures 8a and b) or in Parkin-deficient MEFs (Figure 8c). Because in MEF cells Bax and Bak are functionally redundant in regulating apoptosis, the reduction in the amount of Bax is probably not sufficient to provide protection in these conditions. These results are consistent with an early report showing that Parkin had a minor effect on a large panel of apoptotic stimuli including STS. The protective effect of Parkin observed in neuronal cells by Johnson et al. in similar conditions probably reflects the fact that Bax is essential to prevent death in neurons, whereas Bak has no role. Therefore, we assessed PARK2 overexpression in Bak-deficient MEF cells (Figures 8d and e). In this

Figure 5  Mitochondrial network fragmentation and OPA1 expression. (a, c) IKKαβ−/− and (b, d) IKKα−/− MEF cells were transfected with plasmids encoding for IKKα, IKKβ, NEMO, or both IKKα+IKKβ at the dose of 1 μg. mock serves as a control. (a and b) After overnight culture, cells were stained with specific antibodies directed against HSP60 (in red) and mitochondrial fragmentation was analyzed (c and d) OPA1 expression. As control Hsp60 was used as a loading control to normalize protein content. Furthermore immunoblots were probed with specific antibodies to IKKα, IKKβ, and NEMO. Actin was used as a control. Mean ± S.D. of three independent experiments; *P < 0.05
context, our results demonstrated a protective effect of Parkin on STS-mediated mitochondrial depolarization (Figure 8d) and cell death (Figure 8e). This effect was not dependent on the dose used, as in a recent report the authors show that at the dose of 20 μg, which is 20-fold higher than the dose used here, Park2 overexpression favors cell death.30

**Conclusions**

In conclusion, our results demonstrate that despite a decrease in the level of OPA1, associated with reductions in mitochondrial network in Parkin-deficient cells, PARK2 overexpression had no impact on the expression of OPA1 in MEF cells. Thus, these findings do not confirm the major role of Parkin in OPA1
regulation reported by Müller–Rischart.\textsuperscript{13} As Bax has a preponderant role over Bak in primary neurons, the protective effect of Parkin may rather be related to the ubiquitination of Bax and the limitation of its mitochondrial translocation to the mitochondria, previously reported to dampen the apoptotic response.\textsuperscript{31} By showing that the absence of IKK\textsubscript{α}, with or without IKK\textsubscript{β}, has an impact on mitochondrial network morphology and OPA1 expression, our results point out a role of the nonclassical NF-κB pathway in the regulation of mitochondrial dynamics and OPA1 expression, rather than to a role of NEMO, which is essential for the canonical NF-κB pathway.

Materials and Methods

Cell culture and reagents. MEF (WT, NEMO\textsuperscript{−/−}, IKK\textsubscript{α}\textsuperscript{−/−}, IKK\textsubscript{β}\textsuperscript{−/−}, and Parkin\textsuperscript{−/−}) cells were cultured in Dulbecco’s modified Eagle’s medium (DMEM; Invitrogen, Carlsbad, NM, USA), supplemented with 10% fetal calf serum, 2 mM L-glutamine, 50 IU penicillin, and 50 μg/ml streptomycin in 5\% CO\textsubscript{2}, at 37 °C. MEF cells were transfected using lipofectamine2000 (Invitrogen) with plasmids encoding for either GFP or PARK2 plasmid. STS was purchased from Sigma (La Chapelle-sur-Endre, France).

Flow cytometry. After 24 h, Parkin-transfected cells were treated overnight with STS. Cells were also cotransfected with GFP, and the percentage of dying cells was assessed using a propidium iodide nuclear probe (PI\textsuperscript{+}, Molecular Probes, Life Technology, Cergy Pontoise, France). Mitochondrial membrane potential was assessed using DioC\textsubscript{6} (Molecular Probes). Cells were analyzed by flow cytometry (FC500 flow cytometer, Beckman Coulter, Villepinte, France).

Real-time PCR analysis. Total RNA was isolated from the different MEF cell lines after 24 h of transfection. cDNA synthesis was performed using the AffinityScript QPCR cDNA Synthesis Kit (Stratagene, Agilent Technologies, Paris, France) with 1 μg of total RNA and random primers. The resulting complementary-DNA product was expanded using the Absolute Blue QPCR SYBR Green ROX Mix (Thermo Scientific, Illkirch, France) with specific primers for murine OPA1 (forward: 5′-ACCACAGGAAGCTCAAAA-3′, reverse: 5′-TCTGATCTCAGAAACAA-3′), sample input was normalized through quantification of the transcript levels of the housekeeping genes Gapdh (forward: 5′-GGCTCAGAAGCTCAAGAA-3′, reverse: 5′-TCTGATCTCAGAAACAA-3′), and Rps18 (forward: 5′-TCTGATCTCAGAAACAA-3′, reverse: 5′-GGCTCAGAAGCTCAAGAA-3′). Detection was performed with the ABI PRISM 7900HT Sequence Detection System (Applied Biosystems, Saint Aubin, France). Threshold cycle (Ct) values were obtained for each gene using the instrument software and auto-Ct function. The relative levels of gene expression were determined using the delta–delta Ct method.

Immunoblotting. Total lysates were extracted using 1% NP40 (150 mM NaCl, 50 mM Tris HCl pH 7.4, 10 mM EDTA) supplemented with a cocktail of antiproteases and antiphosphatases. A total of 20 μg of proteins from each point were resolved by SDS-NuPAGE (4–12% Bis-Tris gels, Novex, Les Ulis, France) and transferred to

Figure 7 Modulation of Bax and Bak expressions in Parkin-transfected MEF cells. Immunoblots from (a) WT, (b) Parkin\textsuperscript{−/−}, and (c) IKK\textsubscript{αβ}\textsuperscript{−/−} as described in Figure 2 and Figure 6 were probed with Bax and Bak antibodies. Values indicate Bax and Bak levels with respect to the baseline (mock, value set as 1) and to actin as a control of loading. Histograms represent mean ± S.D. of three independent experiments; *P < 0.05.
nitrocellulose membranes (Amersham Biosciences, Les Ulis, France). Nonspecific sites were blocked by incubation with 5% milk for 1 h at room temperature, and the membranes were then incubated with mouse monoclonal anti-OP A1 (BD Biosciences Pharmingen, Le Pont de Claix, France; clone 18), anti-Parkin (Millipore, Le Pont de Claix, France), and rabbit anti-Bax (N-20, from Santa-Cruz, Le Pont de Claix, France). Antibodies specific for IKKα, IKKβ, and NEMO were purchased from Cell Signaling (St Quentin Fallavier, France). Equal protein loading was assessed by probing the membranes with anti-actin (Millipore) and anti-HSP60 monoclonal antibodies (Stressgen, St Quentin Fallavier, France). Membranes were treated with horseradish peroxidase-linked goat anti-mouse or anti-rabbit secondary antibodies (Amersham Biosciences). Immunoreactive proteins were detected and quantified by enhanced chemiluminescence (Amersham Biosciences) using a CCD camera (GBOX, SYNGENE, Ozyme, France).

**Fluorescence microscopy.** MEF cells were grown on coverslips and fixed with 4% paraformaldehyde (Sigma-Aldrich, St Quentin Fallavier, France). Cells were then incubated with mouse monoclonal anti-cytochrome c (BD Biosciences, St Quentin Fallavier, France; clone 6H2.B4), or goat anti-HSP60 (Santa Cruz) antibody. Cells were washed and incubated for 2 h with Alexa Fluor secondary anti-mouse, anti-rabbit, or anti-goat antibodies (Molecular Probes). Images were acquired with a Zeiss LSM 510 confocal microscope equipped with an oil-immersion fluorescence objective (Carl Zeiss, Inc., Périgny, France). Cells were also stained with 100 nM Mitotracker green (Invitrogen) for 30 min. Cells were imaged by epifluorescence using a Zeiss Axiovert 200 M inverted microscope. Deconvolution was performed by subtracting a Gaussian blur (σ: 12) to the original image. Deconvolved images were then processed with a certain threshold to obtain a binary image of the mitochondrial network. The binary image was then analyzed using the Analyze Particle function of ImageJ to determine the number and the size of mitochondria in individual cells.

**Conflict of Interest**
The authors declare no conflict of interest.

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