Nickel Oxide Nanoparticles Trigger Caspase- and Mitochondria-Dependent Apoptosis in the Yeast Saccharomyces cerevisiae

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Supporting Information

ABSTRACT: The expansion of the industrial use of nickel oxide (NiO) nanoparticles (NPs) raises concerns about their potential adverse effects. Our work aimed to investigate the mechanisms of toxicity induced by NiO NPs, using the yeast Saccharomyces cerevisiae as a cell model. Yeast cells exposed to NiO NPs exhibited typical hallmarks of regulated cell death (RCD) by apoptosis [loss of cell proliferation capacity (cell viability), exposure of phosphatidylserine at the outer cytoplasmic membrane leaflet, nuclear chromatin condensation, and DNA damage] in a process that required de novo protein synthesis. The execution of yeast cell death induced by NiO NPs is Yca1p metacaspase-dependent. NiO NPs also induced a decrease in the mitochondrial membrane potential and an increase in the frequency of respiratory-deficient mutants, which supports the involvement of mitochondria in the cell death process. Cells deficient in the apoptosis-inducing factor (aif1Δ) displayed higher tolerance to NiO NPs, which reinforces the involvement of mitochondria in RCD by apoptosis. In summary, this study shows that NiO NPs induce caspase- and mitochondria-dependent apoptosis in yeast. Our results warn about the possible harmful effects associated with the use of NiO NPs.

INTRODUCTION

Because of their unique physicochemical properties and reactivity, metal oxide nanoparticles (NPs) have been manufactured in greater amounts in the past decade. Nickel oxide (NiO) NPs are used for multiple applications, such as catalysts, memory cells, electrode materials in multilayer ceramic capacitors, diesel-fuel additives, and pigments for ceramics and glasses.1 As the nanotechnology industry expands, exposure to NiO NPs emerges as a significant occupational hazard. In addition, a deeper understanding of the potential impact of NiO NPs on the environment and human health is required.

Adverse effects as a consequence of exposure to NiO NPs, such as cytotoxicity, inflammation, and genotoxicity in human pulmonary cells, have been reported.2−3 The International Agency for Research on Cancer classified nickel compounds as group 1 (carcinogenic to humans) and metallic nickel as group 2B (possibly carcinogenic to humans).4

Cell death can be classified into not programmed “accidental cell death” (ACD) and “regulated cell death” (RCD). The first subtype of cell death (ACD) occurs as a consequence of the exposure of cells to severe physical, chemical, or mechanical stimuli; cells die in an uncontrollable manner as a result of their immediate loss of structural integrity. RCD involves a genetically encoded machinery and can result from multiple signaling pathways, including apoptosis. The designation “programmed cell death” (PCD) is used to refer to RCD that occurs in specific physiological scenarios.7,8 Many human diseases, such as several neurological disorders (for example, Huntington’s, Parkinson’s, and Alzheimer’s disease) and cancer, are associated with the dysregulation of cell death pathways.9 In this context, it has been shown that exposure to NiO NPs induces apoptosis in different cell lines, such as in...
human bronchial epithelial (BEAS-2B),\textsuperscript{10} lung epithelial (NCIH460),\textsuperscript{11} liver (HepG2),\textsuperscript{12,13} and neuronal (SH-SY5Y) cells.\textsuperscript{14}

The yeast Saccharomyces cerevisiae is a relevant eukaryotic cell model in different fields, such as biochemistry and molecular and cellular biology. In the past decade, this unicellular microorganism has been used in the study of RCD.\textsuperscript{15,16} This yeast displays some similarities (orthologues) with mammalian (including human) cells, which makes it an effective cell model for human diseases.\textsuperscript{17} In a previous work, it was found that exposure of \textit{S. cerevisiae} cells to NiO NPs induced the loss of metabolic activity and cell viability in a dose-dependent manner.\textsuperscript{18} In a subsequent work, it was observed that the loss of cell viability induced by NiO could be attributed to the intracellular generation of reactive oxygen species (ROS) and glutathione depletion.\textsuperscript{19} It has been described that oxidative stress is a common denominator of RCD initiation.\textsuperscript{20} The work presented here aims to characterize the cell death observed in \textit{S. cerevisiae} upon exposure to NiO NPs. For this purpose, yeast cell death induced by NiO NPs was assessed by monitoring the loss of plasma membrane integrity and cell viability. Subsequently, the presence of typical morphological and biochemical features of apoptotic cell death was examined in yeast cells exposed to a cytotoxic concentration of NiO NPs. In addition, the dependence of known apoptotic regulators/executors (Yca1p and Aif1p) in RCD induced by NiO NPs was evaluated.

\section*{Experimental Procedures}

\textbf{Preparation of Nickel Oxide Nanoparticle Suspensions.} Commercial NiO NPs (catalog no. 637130, with a particle size of <50 nm and a purity of 99.8%, trace metal basis) were purchased from Sigma-Aldrich (St. Louis, MO).

NiO NP stock suspensions (at 0.5 g/L) were prepared in deionized water, sterilized under an ultraviolet lamp for 30 min, and stored at 4 °C (for up to one month). Before being tested, stock suspensions of NiO NPs were vigorously vortexed and sonicated for 1 h in an ultrasonic bath (80–160 W, Bandelin Sonorex RK 100).

Physicochemical characterizations of NiO NPs, namely, particle size and morphology, agglomeration, surface charge, and dissolution in 10 mM 2-(N-morpholino)ethanesulfonic acid (MES) buffer (Sigma-Aldrich), were previously carried out.\textsuperscript{15}

\textbf{Strains, Media, Growth Conditions, and Treatments.} Experiments were carried out using \textit{S. cerevisiae} BY4741 (MATa; his3Δ1; leu2Δ0; met15Δ0; ura3Δ0) as a wild-type strain (WT); in addition, the respective knockout mutants in the \textit{AIF1} or \textit{YCA1} genes were also used. Strains were obtained from the European Saccharomyces cerevisiae Archive for Functional Analysis (EUROSCARF).

\textit{S. cerevisiae} BY4741 was maintained at 4 °C on YPD agar containing 5 g/L yeast extract (Difco-BD), 5 g/L peptone (Difco-BD), 10 g/L glucose (Merck), and 15 g/L agar (VWR Chemicals). The knockout strains were maintained on YPD agar with 0.2 g/L Geneticin (G418 disulfate salt, Sigma-Aldrich). The assessment of respiratory-deficient (RD) mutants (see below) was carried out using YPGly agar medium. YPGly agar is similar to YPD agar, with glucose being replaced with 2% (v/v) glycerol (Merck). In the canavanine assay (see below), the synthetic dextrose (SD) minimal agar medium,\textsuperscript{21} containing 6.7 g/L yeast nitrogen base without amino acids (Difco-BD), 20 g/L glucose, 15 g/L agar, 0.02 g/L uracil (Sigma-Aldrich), 0.02 g/L L-methionine (Sigma-Aldrich), 0.1 g/L L-leucine (Sigma-Aldrich), and 0.02 g/L L-histidine HCl (Sigma-Aldrich) without or with 60 mg/L canavanine (Sigma-Aldrich) (SDU
\textsubscript{can}), was used.

All strains were grown in YPD broth to exponential phase by incubating the cells overnight, to an OD\textsubscript{600} of ~1.0, with continuous shaking (150 rpm) at 30 °C.

After growth, cells were harvested by centrifugation (2500g, 5 min), washed twice with water, resuspended at a density of 1 × 10\textsuperscript{7} cells/mL in 10 mM MES buffer (pH 6.0) with 20 g/L glucose, and incubated without (control) or with 100 mg/L NiO NPs at 30 °C and 150 rpm for ≤48 h.

The effect of 25 mg/L cycloheximide (cyh, Sigma-Aldrich) on the toxicity induced by NiO NPs was assessed by pre-incubation with this compound for 30 min before exposure to NiO NPs. Control experiments have shown that, at the concentration used, cycloheximide was not toxic to yeast cells.

\textbf{Survival Assay.} The effect of NiO NPs on yeast cells was assessed by a cell viability assay (clonogenic test), as previously described.\textsuperscript{18} Briefly, after exposure to NiO NPs, yeast cells were serially diluted in deionized water, plated on YPD agar, and incubated for 72–96 h at 30 °C. Cell viability was calculated considering the number of colony-forming units (CFUs) per milliliter at time zero as a reference (100%).

\textbf{Evaluation of Plasma Membrane Integrity.} Plasma membrane permeability was assessed using the nonpermeant propidium iodide (PI) probe. Yeast cells (1 × 10\textsuperscript{7} cells/mL) were incubated with 4.5 μM PI (Sigma) for 10 min at room temperature in the dark.

The plasma membrane potential was monitored by staining the cells with bis-1,3-dibutylbarbituric acid trimethine oxonol [DiBAC\textsubscript{4}(3)], known as Oxo (Oxo) (Molecular Probes, Invitrogen). Yeast cells (1 × 10\textsuperscript{7} cells/mL) were incubated with 1.94 μM Oxo (final concentration) for 10 min at 30 °C in the dark. In the double-staining protocol (Oxo/PI), cells were first stained with Oxo and, subsequently, with PI, as described above.

As a positive control (cells with a depolarized and permeabilized membrane), yeast cells were heated at 65 °C for 1 h. The percentage of PI or Oxo negative cells was determined with a fluorescence microscope (Leica DMLB), equipped with an HBO-100 mercury lamp and filter set 13 (excitation filter (band-pass filter, BP) BP 450–490, dichromatic mirror S10, and suppression filter (long pass filter, LP) LP S15) from Leica. Doubly stained (Oxo/PI) yeast cells were observed using filter set 13, GFP (excitation filter BP 450–490, dichromatic mirror 500, and suppression filter BP S50–550), or N2.1 (excitation filter BP S15–560, dichromatic mirror S80, and suppression filter LP S90) from Leica. In each experiment, two samples of at least 200 cells (total of ≥400 cells) were counted in randomly selected microscope fields.

Images were acquired with a Leica DC 300 F camera and processed using Leica IM 50-Image Manager software.

\textbf{Annexin V Staining.} Annexin V [fluorescein isothiocyanate (FITC), FITC-conjugated], a protein that specifically binds to phosphatidylserine (PS) residues, was used to detect the externalization of PS.\textsuperscript{23} After treatment, yeast cells were collected, washed twice with 40 mM phosphate-buffered saline (PBS) (pH 6.8), and resuspended at a density of 1 × 10\textsuperscript{7} cells/mL in sorbitol buffer [1.2 M sorbitol, 0.5 mM MgCl\textsubscript{2}, and 35 mM K\textsubscript{2}HPO\textsubscript{4} (pH 6.8)]. Cell walls were digested by incubation with 90 units/mL lyticase (Sigma-Aldrich) in sorbitol buffer for 45 min at 28 °C. Then, cells suspensions (1 mL) were centrifuged (250g, 5 min), resuspended in 60 μL of incubation buffer (containing the Annexin-V-FLUOS labeling solution and PI) (Roche), and incubated for 10 min at 25 °C. Cells were observed in a fluorescence microscope using filter set 13. In each experiment, two samples containing at least 100 cells (total of ≥200 cells) were counted in randomly selected microscope fields. Images were acquired and processed as described above.

\textbf{Assessment of the Mitochondrial Membrane Potential.} The evaluation of the mitochondrial membrane potential (ΔΨ\textsubscript{m}) was carried out using the probe 3,3'-dihexyloxacarbocyanine iodide [DiOC\textsubscript{6}(3)] (Sigma-Aldrich).\textsuperscript{24,25} Yeast cells (1 × 10\textsuperscript{7} cells/mL) were collected and incubated with 30 nM DiOC\textsubscript{6}(3) for 20 min at 30 °C in the dark. Then, the cells were resuspended (200 μL per well), in quintuplicate in a 96-well flat microplate (Orange Scientific). The fluorescence intensity [in relative fluorescence units (RFUs)] was measured in a PerkinElmer (Victor3) microplate reader at a fluorescence excitation wavelength of 485/14 nm and an emission wavelength of 535/25 nm. ΔΨ\textsubscript{m} was expressed as the ratio between
Figure 1. Cell viability and plasma membrane integrity in yeast cells of S. cerevisiae BY4741 exposed to NiO NPs. Cells were suspended in MES buffer in the absence (control) or presence of 100 mg/L NiO NPs. (A) Cell viability evaluated by colony-forming unit (CFU) counting. (B) Membrane permeability assessed by propidium iodide (PI) exclusion. (C) Membrane polarization monitored by Oxonol (Oxo) exclusion. (D) Photomicrographs of cells not exposed (control) or exposed to 100 mg/L NiO for 24 h and then doubly stained with PI and Oxo. Cells were visualized by fluorescence, using filter set I3 (green and orange-red fluorescence), GFP (green fluorescence), or N2.1 (orange-red fluorescence), or by phase contrast microscopy. The data represent the means (±SD) of at least three independent experiments.

RESULTS

Impact of NiO NPs on Cell Viability and Plasma Membrane Integrity. Cell viability was evaluated by quantification of the reduction of the number of CFUs after different periods of contact time with yeast NiO NPs. Exposure to NiO NPs induced an abrupt loss of cell cultivability. Cell viability dropped to ~60% in the first 8 h (Figure 1A); later exposure to NiO NPs induced a slower reduction in cell viability, ~33% of the cells being viable at 48 h (Figure 1A).

To define a lethal scenario in yeast cells exposed to NiO NPs, a kinetic assessment of cell membrane integrity was carried out through the monitoring of PI uptake. It is assumed that viable cells exclude PI (PI negative cells) (Figure 1D, control). As one can see in Figure 1B, a small proportion of the cell population (2%) remained permeable to PI after contact with NiO NPs for 8 h.

The quantification of RD mutants induced by NiO NPs was assessed by the complete lack of growth of these mutants on nonfermentable medium (YPGly).27,28

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The uptake of PI requires significant damage to the cell membrane. This means that the permeability to PI represents irreparable damage of the plasma membrane (permanent loss of barrier function)\(^{31}\) and cell death.\(^{7,32}\) On the other hand, it has also been reported that the uptake of the lipophilic anionic probe DiBAC\(_4\) (3), known as Oxonol (Oxo), occurs in less damaged cells\(^{33}\) with a reduced membrane potential (depolarized cell membrane).\(^{22,34}\) Nonstressed cells displayed a polarized membrane that remained unstained (Oxo negative cells) (Figure 1D, control). To obtain further information about the impact of NiO on the yeast plasma membrane, after exposure to NPs, cells were stained with Oxo. The exposure of yeast cells to NiO NPs for \(\leq 8\) h did not have any impact on the plasma membrane potential, as cells remained Oxo negative (Figure 1C). After this time, the cells became permeable to Oxo. Thus, after exposure to NPs for 24 and 48 h, \(\sim 20\) and \(\sim 50\)% of the cell population were Oxo positive, respectively (Figure 1C). After the same period of exposure to NPs, \(\sim 8\) and 15% of the yeast cells were PI positive, respectively (Figure 1B). Comparative analysis of the cells stained with Oxo and PI revealed that there was a fraction of cells which membrane presented a reduced potential, becoming Oxo positive, although they maintained their impermeability to PI (Figure 1B,C). These results indicate that under NiO stress, de-energization precedes permeabilization of the cell membrane. This possibility was confirmed by fluorescence microscopic observation of the yeast cells exposed to NPs and doubly stained with Oxo and PI, where it was possible to detect PI negative/Oxo positive cells (Figure 1D). However, all cells that were PI positive presented a depolarized membrane (Oxo positive) (Figure 1D). Altogether, the results show that the loss of cultivability preceded the loss of membrane potential and plasma membrane integrity (Figure S1).

After the exposure of yeast cells to NiO NPs for 24 h, \(\sim 8\)% of yeast cells have passed the “point of no return,”\(^{35}\) as they exhibited irreversible plasma membrane permeabilization (PI positive cells) and thus can be classified as dead.\(^{7}\) At this time (24 h), although only \(\sim 55\)% of the cell population displayed the ability to undergo proliferation (Figure 1A and Figure S1), \(\sim 80\)% of the cell population had intact membranes (PI negative/Oxo negative) and, thus, can be classified as viable, which suggests RCD in yeast cells exposed to NiO NPs. These results prompted us to further characterize the cell death phenotype NiO NPs induced through the assessment of different morphological and biochemical features.

**NiO NPs Induce Cell Death with Typical Apoptotic Hallmarks.** In a previous work, yeast cells exposed to NiO NPs presented an intracellular overload of reactive oxygen species (ROS),\(^{19}\) which can mediate yeast cell death by apoptosis. To confirm this possibility, yeast cells exposed to NiO NPs were doubly stained with Annexin V-FITC and PI to detect the externalization of phosphatidylserine (PS). The exposure of PS at the outer leaflet of the plasma membrane (loss of membrane asymmetry) is considered an early sign of apoptosis and one of the most reliable apoptosis markers.\(^{36}\) Cells that were Annexin V positive and PI negative (Figure 2A) are considered apoptotic.\(^{15}\) As one can see in Figure 2B, after exposure to NiO NPs for 6 h, the percentage of yeast cells that were Annexin V positive and PI negative was significantly different from the control; after this time, an increase in the number of apoptotic cells was observed.

Another feature of apoptosis is the requirement of de novo protein synthesis.\(^{37-41}\) To test this possibility, the effect of cyh, a protein synthesis inhibitor in eukaryotes,\(^{42}\) on RCD induced by NPs was investigated. Yeast cells were incubated for 6 or 24 h with NiO in the presence or absence of 25 mg/L cyh, and CFUs were counted. NiO- and cyh-treated cells displayed a higher percentage of CFUs compared to the NiO-treated cells in the absence of cyh (Figure 2C). This result suggests that the loss of cell viability in response to NPs stress requires the participation of cellular machinery, namely de novo protein synthesis, which is consistent with an apoptotic program.
Apoptotic cells present condensed chromatin and modification of the shape of the nucleus as a consequence of the proteolysis of key nuclear proteins.\textsuperscript{43} To evaluate if NiO NPs induce chromatin condensation and modification of the morphology of the nucleus, nontreated (control) and NiO-treated yeast cells were stained with the DNA-binding dye DAPI. Fluorescence microscopy observations revealed that NiO-treated yeast cells displayed ring- and kidney-shaped condensed chromatin, characteristic of apoptotic cells (Figure 2D). Control cells displayed homogeneous round nuclei (Figure 2D). A yeast population exposed for 8 h to NiO NPs presented a percentage of cells with apoptotic nuclei significantly different from those of the control (cells not exposed to NiO); the percentage of cells with apoptotic nuclei increased with NP exposure duration (Figure 2E).

Altogether, the results presented above indicate that NiO NPs induced an apoptotic phenotype in yeast cells.

**NiO NPs Induce Mitochondrial Membrane Depolarization.** It has been shown that mitochondria play a central role in the execution of RCD by apoptosis in yeast cells.\textsuperscript{40,44–47} Modification of $\Delta W_m$ led to the disruption of mitochondrial function. In the work presented here, $\Delta W_m$ was assessed by quantification of the fluorescence of cells stained with the membrane potential-sensitive probe DiOC$_6$(3). This probe accumulates in the negatively charged mitochondrial matrix, and the extent of its uptake, as measured by the intensity of cellular fluorescence, reflects the $\Delta W_m$.\textsuperscript{25,48} The observation of the control cells by fluorescence microscopy allowed the visualization of the typical mitochondrial networks as a result of the polarization of the inner mitochondrial membrane (Figure 3A). Fluorescence quantification revealed that after exposure to NiO NPs for 1 h, yeast cells presented a progressive reduction in fluorescence ($\leq$6–8 h), compared to that of control cells (Figure 3B). These results indicate that NiO NPs induce the depolarization of the mitochondrial membrane. Microscopic observation of cells exposed to NPs for 8 h confirmed the presence of yeast cells with decreased green fluorescence and even the presence of yeast cells in which DiOC$_6$(3) failed to accumulate (Figure 3A) probably as a consequence of the dissipation of the mitochondrial membrane potential. Yeast cells exposed to NiO NPs for 24 h presented a fluorescence higher than that of cells exposed for 6–8 h (Figure 3B). However, this increase in fluorescence can be explained by the presence of some cells exhibiting a strong green fluorescence, all over the cells, as a result of the accumulation of DiOC$_6$(3) in the cytosol and not due to the repolarization of the mitochondrial membrane. Double staining of these cells with DiOC$_6$(3) and PI confirmed that the strongly fluorescent cells presented a permeabilized plasma membrane (PI positive cells) (Figure S2).

**NiO NPs Induce Yca1p- and Aif1p-Dependent Apoptosis.** To better characterize RCD, the effect of NiO NPs on the cell viability of knockout mutant strains $\textit{yca1}$Δ and $\textit{aif1}$Δ, devoid of protein known to be involved in apoptotic cell death, was evaluated.

The proteases known as caspases are usually activated in the early stages of apoptosis and play a pivotal role in yeast cell death execution.\textsuperscript{49,50} Yeast cells bear the metacaspase Yca1p, which is an ortholog of mammalian caspases.\textsuperscript{51} The disruption of the $\textit{YCA1}$ gene ($\textit{yca1}$Δ strain) renders cells significantly more resistant to the toxic effect of the NPs (Figure 4), which suggests that NiO-induced cell death is caspase-dependent.

The apoptosis-inducing factor (Aif1p) is a flavoprotein with NADH oxidase activity localized in the mitochondrial matrix. The expression of Aif1p is induced by the toxic effect of NiO NPs (Figure 4). To better characterize RCD, the effect of NiO NPs on the cell viability of knockout mutant strains $\textit{yca1}$Δ and $\textit{aif1}$Δ, devoid of protein known to be involved in apoptotic cell death, was evaluated.

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cells increased with NiO NP exposure duration. Consistent with this result, the presence of cells lacking mtDNA (as revealed by DAPI staining) in the population exposed to NPs for 24 h (Figure 2D, NiO-treated cells) was observed. In control cells, the percentage of RD cells remained approximately constant (2%), which corresponded to the values previously observed, was most likely the origin of the damage of nuclear DNA due to intracellular ROS accumulation in yeast cells exposed to NiO. This assay is based on the mutation in the plasma membrane arginine permease (encoded by CAN1), which originates in cells (canR mutants) resistant to l-canavanine (an analogue of l-arginine). It was possible to observe an increase of the number of CanR colonies (among the population of viable cells) in cells exposed to NiO NPs compared to the control. The mutation frequency induced by NiO NPs increased with exposure duration: after exposure to NiO NPs for 6 and 24 h, the frequency of CanR mutations was approximately 7- and 12-fold higher, respectively, compared with that of cells not exposed to NiO NPs (Figure 5B). These results demonstrate the damage of nuclear DNA by NiO NPs. In the control, the frequency of CanR mutations was similar after 6 and 24 h (7 per 10^7 viable cells), which is similar to the rate described in the literature (10–20 per 10^7 viable cells).

**DISCUSSION**

The level of manufacturing and use of metal oxide NPs, such as NiO NPs, has increased greatly in the past decade and, inevitably, raised concerns and uncertainties about the possible adverse effects of NPs on the environment and human health. In a previous work, it was proposed that NiO NPs exert their toxic effect on yeast cells by an indirect mechanism: NiO NPs adsorb to the cell wall and release nickel at the NP–cell wall interface, producing a toxic response. To date, little has been known about the mechanisms underlying the loss of yeast cell viability induced by NiO NPs. To understand deeply the mechanism of the toxicity induced by NiO NPs, the well-known apoptotic model *S. cerevisiae* was used.

It was shown that the exposure of yeast cells to NiO NPs for ≤24 h induced a loss of cell proliferation (cell viability) without disrupting cell membrane integrity (Figure 1). However, extending the exposure of yeast cells to NiO yielded cell death (plasma membrane breakdown, assessed through PI staining) (Figure 1). It was also found that yeast cells exposed to NPs lost their plasma membrane asymmetry (PS exposure to the outer membrane leaflet) (Figure 2). The loss of cell viability induced by NiO required the involvement of the metacaspase Yca1p (Figure 4) and was attenuated by the presence of a protein synthesis inhibitor (cyh) (Figure 2C). All of these features are typical hallmarks of RCD by apoptosis in yeast cells.[32,39,49] Our results are in agreement with those that described the PS externalization[10,14] and the involvement of caspase-3 in apoptosis induced by NiO NPs in mammalian cell lines.[10–12,60] These results also suggest that *S. cerevisiae* and mammalian cell lines, exposed to NiO NPs, share some common apoptotic features.

In a previous work, it was observed that antioxidants [L-ascorbic acid (an efficient scavenger of free radicals)[63,64] and N-tert-butyl-α-phenylnitrene, known as PBN (a free radical spin trapping agent)][65] prevented ROS overload and almost fully abolished the loss of cell viability induced by NiO NPs;[19] these results strongly suggest that oxidative damage plays a central role in yeast cell death. The increased level of ROS, previously observed, was most likely the origin of the damage of the nuclear DNA (observed through the increase in the number of can1R mutants) (Figure 5B). Our results are in agreement with those describing DNA damage in human cell lines exposed to NiO NPs.[7,12,24,66] In addition, in the work presented here, the induction of RD mutants in yeast cells exposed to NPs was shown (Figure 5A), which can also be attributed to the increased ROS levels in these cells. The loss of part or all of the mtDNA was further confirmed through the observation by fluorescence microscopy of yeast cells stained

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*Figure 5. NiO NPs induce mitochondrial and nuclear DNA damage in *S. cerevisiae* BY4741. Cells were suspended in MES buffer in the absence (control) or presence of 100 mg/L NiO NPs. (A) Mitochondrial DNA damage assessed through the formation of respiratory-deficient cell mutants (cells without the capacity to grow on a nonfermentable carbon source, YPGly). (B) Nuclear DNA damage evaluated through the formation of canavanine resistant (CanR) mutants. The data represent the means (±SD) of at least three independent experiments. Mean values are significantly different: *P < 0.05 in comparison with untreated cells (control); unpaired *t* test.*
Figure 6. Sequence of events associated with NiO NP-induced cell death in S. cerevisiae. Proposal based on the results obtained in this work and previous work.18,19

with DAPI; in the cells exposed to NiO for 24 h, it was not possible to visualize mtDNA (Figure 2D, NiO-treated cells).

The involvement of mitochondria in yeast apoptosis upon exposure to different stimuli has been described.20,44,45 Consistent with the possibility of NiO NPs inducing mitochondria-dependent apoptosis, in a previous work, it was shown that the mitochondria is the main contributor for ROS production.19 Here, it is described mitochondrial membrane depolarization (Figure 3), most likely, as a consequence of the permeabilization of the membrane. The dissipation of ΔΨm and the preservation of cell membrane integrity (Figure 1A,B) are usually characteristic of early apoptotic cells.64 The induction of formation of mutants with a petite phenotype, as described above (Figure 5A), together with our previous results, which have shown that cells from a derivative ρ0 mutant (without mitochondrial DNA) presented reduced levels of ROS and increased resistance to NiO NPs,19 reinforces the possibility of an active role of mitochondria in the execution of an apoptotic program in response to exposure to NiO NPs.

Previous studies have described the requirement of the apoptosis inducing factor (Aif1p) in RCD induced by several stimuli.20,45 Yeast cells without the mitochondrial pro-apoptotic factor Aif1p (aif1Δ strain) presented higher resistance to NiO NPs (Figure 4), which suggests the participation of the Aif1p in the execution of RCD by apoptosis. This result also supports the role of mitochondria in apoptosis induced by NiO NPs. It is also described that in response to apoptotic stimuli, Aif1p is released from mitochondria and translocated into the nucleus, where chromatin condensation and DNA fragmentation are carried out and, thus, cellular demise is provoked.52 Modifications of the nucleus morphology (including the increase in the number of cells with ring- and kidney-shaped condensed chromatin) were observed in yeast cells treated with NPs (Figure 2D); these facts are compatible with the possibility of the participation of Aif1 as the executor of NiO-induced RCD.

In summary, it was shown that exposure of yeast cells to NiO NPs induces cell death, which exhibits several typical features of RCD by apoptosis: (i) loss of proliferation capacity dependent on the participation of cellular machinery, namely, de novo protein synthesis and the activation of the metacaspase Yca1p; (ii) externalization of PS at the surface of the cytoplasmic membrane in cells retaining membrane integrity; and (iii) modification of nucleus morphology. Other further insights were also obtained, namely, mitochondrial membrane potential dissipation, induction of petite mutants, and the participation of the Aif1p; these facts support the involvement of mitochondria in the RCD induced by NiO NPs. An overview of the sequence of detrimental effects associated with NiO NPs induced in S. cerevisiae is presented in Figure 6.

In conclusion, the results presented here strongly suggest that NiO NPs induce caspase- and mitochondrial dependent RCD by apoptosis in S. cerevisiae. Using this yeast, as a model organism, new information’s about the events underlying cellular death, more specifically, about the role of mitochondria in the apoptosis induced by NiO NPs, are given. The data presented might contribute to the understanding and prevention of the possible detrimental effects associated with the use of NiO NPs.

■ ASSOCIATED CONTENT

$\$ Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acs.chemrestox.8b00265.

Comparison of the evolution of cell death, plasma membrane integrity, and double staining [DiOC6(3) and propidium iodide] of S. cerevisiae BY4741 exposed to NiO (PDF)

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Notes

The authors declare no competing financial interest.
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