

RESEARCH PAPER

The antidote effect of quinone oxidoreductase 2 inhibitor against paraquat-induced toxicity *in vitro* and *in vivo*

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BACKGROUND AND PURPOSE

The mechanisms of paraquat (PQ)-induced toxicity are poorly understood and PQ poisoning is often fatal due to a lack of effective antidotes. In this study we report the effects of N-[2-(2-methoxy-6H-dipyrido{2,3-a:3,2-e}pyrrolizin-11-yl)ethyl]-2-furamide (NMDPEF), a melatonin-related inhibitor of quinone oxidoreductase2 (QR2) on the toxicity of PQ *in vitro* & *in vivo*.

EXPERIMENTAL APPROACH

Prevention of PQ-induced toxicity was tested in different cells, including primary pneumocytes and astroglial U373 cells. Cell death and reactive oxygen species (ROS) were analysed by flow cytometry and fluorescent probes. QR2 silencing was achieved by lentiviral shRNAs. PQ (30 mg·kg⁻¹) and NMDPEF were administered i.p. to Wistar rats and animals were monitored for 28 days. PQ toxicity in the substantia nigra (SN) was tested by a localized microinfusion and electrocorticography. QR2 activity was measured by fluorimetry of N-benzylidihydronicotinamide oxidation.

KEY RESULTS

NMDPEF potently antagonized non-apoptotic PQ-induced cell death, ROS generation and inhibited cellular QR2 activity. In contrast, the cytoprotective effect of melatonin and apocynin was limited and transient compared with NMDPEF. Silencing of QR2 attenuated PQ-induced cell death and reduced the efficacy of NMDPEF. Significantly, NMDPEF (4.5 mg·kg⁻¹) potently antagonized PQ-induced systemic toxicity and animal mortality. Microinfusion of NMDPEF into SN prevented severe behavioural and electrocortical effects of PQ which correlated with inhibition of malondialdehyde accumulation in cells and tissues.

CONCLUSIONS AND IMPLICATIONS

NMDPEF protected against PQ-induced toxicity *in vitro* and *in vivo*, suggesting a key role for QR2 in the regulation of oxidative stress.

LINKED ARTICLE

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Abbreviations

Ab, antibody; APF, 3'-(p-aminophenyl)fluorescein; BNAH, N-benzylidihydronicotinamide; DPI, diphenylene iodonium; FACS, flow cytometry; LDH, lactate dehydrogenase; MDA, malondialdehyde; MitoSOX, 3,8-phenanthridinediamine,

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Introduction

Paraquat (1,1'-dimethyl-4,4'-bipyridinium; PQ) is a herbicide used worldwide and the acute exposure to PQ, after accidental or suicidal ingestion, causes thousands of deaths each year. However, clinical management of PQ poisoning is often unsuccessful due to a lack of effective antidotes (Dinis-Oliveira *et al.*, 2009a). PQ poisoning leads to fatal injury of lungs, kidneys, liver, brain and other organs. PQ is toxic to all cell types, but the active accumulation of PQ via diamine transporters in alveolar epithelial cells (pneumocytes type I and II) has been proposed to be responsible for the high sensitivity of lung tissue (Dinis-Oliveira *et al.*, 2008).

Chronic exposure to PQ and other pesticides has been implicated in the pathogenesis of neurodegenerative disorders such as Parkinson's disease (Hatcher *et al.*, 2008) a common neurodegenerative disorder in the elderly, characterized by the loss of dopaminergic neurons in substantia nigra pars compacta (SN). In support of a causative role of PQ in Parkinsonian pathology, many animal models have been developed that show certain features of the disease, like nigrostriatal degeneration or dyskinesia, following acute or chronic exposures to the herbicide or to a structurally related toxin, MPTP (Duty and Jenner, 2011; Nistico *et al.*, 2011). PQ has been shown to cause a selective death of dopaminergic neurons, a process mediated by oxidative stress and a marked activation of microglial cells (Peng *et al.*, 2005; 2009). These events correlate with potent behavioural, electrocortical (ECoG) and neurodegenerative effects following PQ injection into several areas of the rat brain (Mollace *et al.*, 2003). These effects can be prevented by administration of a non-peptidyl superoxide dismutase (SOD) mimetic (Mollace *et al.*, 2003), suggesting that oxidative stress is involved in the neurotoxic response to PQ.

The mechanism whereby PQ induces oxidative stress and elicits cell damage involves a cyclic reduction/oxidation of the PQ di-cation with consumption of NAD(P)H and production of superoxide anion (Bus and Gibson, 1984; Drechsel and Patel, 2008). This process is likely to be catalysed by diaphorases, i.e. enzymes with oxidoreductase activity, localized in different cellular compartments (Drechsel and Patel, 2008; Drechsel and Patel, 2009). Several NAD(P)H-dependent diaphorases have been identified as PQ-reducing enzymes. These include, cytoplasmic NADPH oxidases of the Nox family (Miller *et al.*, 2007; Cristovao *et al.*, 2009), a diphenylene iodonium (DPI)-sensitive thioredoxin reductase, highly expressed in alveolar epithelium (Gray *et al.*, 2007), mitochondrial oxidoreductases present in the electron transport Complex I (Cocheme and Murphy, 2008) and Complex III (Castello *et al.*, 2007; Drechsel and Patel, 2009) or a mitochondrial NADH-dependent reductase VDAC1 (Shimada *et al.*, 2009). Finally, it cannot be excluded that oxidoreductases that preferentially use an alternative electron donor to NAD(P)H, are also involved in PQ-induced cytotoxicity.

N-ribose dihydronicotinamide (NRH): quinone oxidoreductase 2 (NQO2), also known as quinone reductase 2 (QR2), is a NADPH-independent flavoenzyme typically catalysing two-electron reductions of quinones and oxidation of N-alkyl and N-ribosyl nicotinamide (NADH) derivatives (Mailliet *et al.*, 2005); however, one and four electron reactions have also been proposed to be catalysed by this enzyme. QR2 is a homolog of quinone oxidoreductase 1 (QR1), a more extensively studied enzyme of phase II drug metabolism (Bianchet *et al.*, 2008). QR2 and QR1 have partially overlapping substrate specificity, but distinct co-substrates and tissue expression profiles. For this reason both enzymes may play different biological roles.

Quinone and quinone-derivative conversion to hydroquinones is usually detoxifying, but in some cases leads to the bioactivation of cytotoxic compounds, such as menadione and mitomycin (Celli *et al.*, 2006). A remarkable feature of human QR2 is its ability to act as a nitroreductase with high specificity for selected substrates. This property is essential for the bioactivation of the anti-tumour, DNA cross-linking agent CB1954 (Knox *et al.*, 2003). The majority of QR2 substrates are xenobiotics and only a few endogenous substrates are known. For example, a possible role of this enzyme has been suggested in dopamine quinone metabolism and Parkinson's disease (Fu *et al.*, 2008). Interestingly, genetic studies have identified an association of a promoter region polymorphism in the NQO2 gene locus characterized by a 29-bp deletion, with idiopathic and pesticide-induced Parkinson's disease (Harada *et al.*, 2001). In light of more recent findings, this deletion disrupts the sp3 repressor binding site on the NQO2 promoter and favours an increased expression of QR2 (Wang *et al.*, 2004; Yu *et al.*, 2009).

Another important feature of QR2 is its ability to bind several ligands with inhibitory properties, for example natural flavonoids, such as resveratrol and quercetin, anti-tumour drugs such as imatinib, and the neurohormone melatonin. Importantly, QR2 has been identified as the melatonin MT₃ receptor located in the cytoplasm (Mailliet *et al.*, 2005) which may mediate some of the antioxidant activities of melatonin, such as its potent protective effect against PQ systemic toxicity in rats (Melchiorri *et al.*, 1995).

Given the role of QR2 in nitroreduction, bioactivation of cytotoxic xenobiotics and in melatonin signalling, we examined a potential involvement of QR2 in PQ-induced toxicity. We found that a melatonin-related and specific QR2 inhibitor, N-[2-(2-methoxy-6H-dipyrido{2,3-a:3,2-e}pyrrolizin-11-yl)ethyl]-2-furamide (NMDPEF), prevented the cytotoxic effects of PQ *in vitro* in several cell lines and in primary pneumocytes. Knock-down of QR2 partially rescued the cells from PQ-induced toxicity and reduced the protective effect of the QR2 inhibitor. Importantly, NMDPEF showed a potent antidote effect *in vivo* in animal toxicity studies. Our data suggest a novel pharmacological approach to the treatment of PQ poisoning and demonstrate a role for QR2 in the regulation of oxidative stress.

Methods

Cell culture

The human astrocytic cell line U373 (kind gift of M. Pollicita e S. Aquaro, Tor Vergata University, Rome) and human embryonic kidney cells HEK293 were cultured in the complete medium: low-glucose (1%) Dulbecco's modified Eagle's medium (DMEM) supplemented with 10% heat inactivated foetal calf serum (FCS; Lonza, Basel, Switzerland), 2 mM glutamine, 1 mM sodium pyruvate, 100 units·mL⁻¹ penicillin, 100 µg·mL⁻¹ streptomycin (Pen/Strep). Mouse mammary epithelial cells EpH4 were cultured as previously described (Janda *et al.*, 2002). A549 lung carcinoma cells were cultured in F12-K medium, supplemented with 10% FCS and Pen/Strep. Only few specified experiments were carried in high-glucose DMEM (4.5% glucose) complete medium.

Measurement of QR2 activity

Fluorometric assay of QR2 activity was performed as previously described (Mailliet *et al.*, 2005) with a few modifications. Cells were washed once in PBS, collected in PBS with a rubber scraper and centrifuged (400×g, 5 min, 4°C). The resulting cell pellet was resuspended in GPS reaction buffer (50 mM Tris/HCl, pH 7.5, 1 mM n-octyl-β-D-glucopyranoside) and homogenized by a tight glass Dounce homogenizer (Weaton) with 18 strokes on ice. A amount of 0.5 mL of GPS buffer was used for 80% confluent 10 cm dishes. Nuclei and debris were removed by centrifugation (1000×g, 5 min, 4°C). Aliquots of the supernatant were stored at -80°C until required. Protein concentration was determined by BCA assay (Pierce, Rockford, IL, USA). 1–2.5 µg of cell lysate was used for one 170 µL reaction. Reactions were carried out at room temperature in 96 well plates. N-benzylidihydronicotinamide (BNAH, Santa Cruz Biotech., Santa Cruz, CA, USA) oxidation was measured at 1 min intervals using a Victor 1420 fluorometer (Perkin Elmer, Waltham, MA, USA) with excitation at 355 nm and emission at 460 nm. Baseline measurements were taken before the addition of the enzyme source and the reaction followed for 15–20 min. Data were manipulated using Microsoft Excel and then analysed by GraphPad Prism 3.0 software for Michaelis-Menten kinetics (GraphPad, San Diego, CA, USA). The velocity of the reaction was determined by curve fitting and corresponded to the velocity 2 min. after the addition of the enzyme. BNAH fluorescence coefficient per well (96 well plate) was calculated experimentally as 3.5 × 10⁸ M⁻¹.

Isolation of pneumocytes. Primary rat alveolar epithelial cells (mainly type II pneumocytes) were isolated from lung tissue obtained from Wistar rats using techniques described previously (Dusinska *et al.*, 1998) with modifications indicated in the Supporting Information.

Flow cytometry (FACS) analysis of cell viability and apoptosis

U373 astroglial cells were plated at 1 × 10⁴ cells cm⁻² on 24 well plates 2 days before treatment. Growth medium was removed and 500 µL of fresh regular medium containing test compounds or control vehicle was added to each well. At desired

time points the cells were washed with PBS (pH 7.4) once and the cell supernatants were collected into FACS tubes.

EpH4 cells were seeded 2.6 × 10⁴ cells cm⁻² and treated at different time points before analysis, which was done at day 5, after plating. Cells were trypsinized for 3–4 min, (trypsin 0.05%) neutralized in pre-warmed regular medium and collected in FACS tubes.

Primary rat pneumocytes were seed after isolation at a density of 2 × 10⁵ cells cm⁻² on 96 well plates and treatments were started 24 h later. Twenty-four and 72 h after treatment the adherent cells were detached by trypsinization (0.1%, 8 min, 37°C) and pooled with non-adherent cells as described earlier. The cells were then pelleted by centrifugation for 5 min at 1000 × g, and resuspended in 300 µL PBS containing 1% FBS. Trypan blue solution in PBS (0.016%; 100 µL) was added immediately and incubated for 5–10 min before FACS analysis (FACSCanto II, BD Biosciences, Erenbodem, Belgium). The data were collected in 630 nm (PerCP-Cy5) channel.

LDH cell toxicity assay

LDH released into the medium was measured with a colorimetric kit (CytoTox96, Promega, Madison, WI, USA) as described in the Supporting Information.

QR2 knock-down with lentivirus short harpin RNA (shRNA)

Lentivirus particles coding for shRNA against QR2 were produced in 293FT cells (Invitrogen, Carlsbad, CA, USA), transfected with specific pLKO plasmids from Mission shRNA library (Sigma-Aldrich, St Louis, MO, USA) targeting human NQO2 (TRCN0000036709-13) and with lentiviral packaging vectors (kind gift of G. Morrone, UMG, Catanzaro, Italy). G418-resistant 293FT cells (3 × 10⁶) were co-transfected with 2.5 µg pLKO, 1 µg VSVG and 5 µg Delta 8.9 plasmids and Lipofectamine 2000 (Invitrogen). The lentivirus supernatant was collected 42 h post-transfection and was used directly to infect U373 cells. U373 cells (70% confluent) plated on 6 cm dishes were overlaid with 2.5 mL supernatant containing 4 µg·mL⁻¹ polybrene (Sigma-Aldrich). The infection was carried out for 14 h. Infected U373 cells were tested in cytotoxicity assays within 10 days post-infection. EpH4 cells were infected with shRNA lentiviruses prepared as described above, but coding for mouse NQO2 (TRCN0000041873-7) and selected for 8 days with puromycin (2 µg·mL⁻¹). QR2 down-regulation was confirmed by Western blots (see below). Aliquots of cells were frozen and analysed a few days after thawing, in cytotoxicity assays. For cytotoxicity assays, EpH4 cells were seeded on 24 well plates and cultured in 4.5% glucose DMEM, 10% FCS and cytotoxicity measured as described previously.

FACS detection of reactive oxygen species (ROS) in live cells

3,8-phenanthridinediamine, 5-(6'-triphenylphosphonium hexyl)-5,6 dihydro-6-phenyl (MitoSOX; Mito-hydroethidium) and 3'-(p-aminophenyl)fluorescein (APF), (Molecular Probes, Eugene, OR, USA) were used to monitor mitochondrial and cytoplasmic ROS levels, respectively, according to the procedure described in the Supporting Information.

Western blot analysis

For Western blot analysis of QR2, U373 cells (80% confluent) and EpH4 cells (90–100% confluent) were lysed in RIPA buffer (50 mM Tris-HCl, 150 mM NaCl, Igepal 1%) and processed as described by Janda *et al.*, 2011). After addition of Laemmli sample buffer, the lysates were incubated 20 min at 90°C. Protein lysates (50–100 µg) were loaded on 8% or 10% SDS-PAGE gel. For Bis-Tris gradient NuPage gels (Invitrogen) lysates were processed according to the manufacturer's indications. The gels were blotted on PVDF membrane and blocked with 5% milk in Tris buffered saline supplemented with 0.05% Tween-20 (TTBS) for 3 h. QR2 and QR1 were detected by overnight incubation with goat anti-QR2 (Sc-18574, Santa Cruz Biotech.) diluted 1:100 in 5% BSA and then with secondary horseradish peroxidase-conjugated anti-goat antibody (Ab) (Santa Cruz Biotech., Sc-2020) 1:5000 in 5% milk.

Immun-Star WesternC Chemiluminescent kit was used for the chemiluminescent (ECL) signal detection. Mouse anti- α -tubulin (Santa Cruz Biotech.) was used to detect tubulin on the same PVDF membrane by standard procedures, but the ECL signal was recorded by ChemiDoc XRS (Biorad, Hercules, CA, USA).

Studies of systemic toxicity in Wistar rats

All animal care and experimental procedures used in the PQ-induced toxicity studies were approved by the Italian Ministry of Health according to the law DL 116/92. Wistar rats (290–310 g; Harlan, Italy) were used. Animals were housed under controlled temperature (20 + 2°C), humidity (60%) and a 12 h light/12 h dark cycle (light on at 0800 h); food and water were available *ad libitum*. Rats were divided in five groups (six per group): G1 – Control, G2 – PQ only, G3 – NMDPEF only and G4, G5 – PQ plus NMDPEF 3 and 4.5 mg·kg⁻¹ respectively. A 50 mg·mL⁻¹ NMDPEF solution in DMSO was diluted in pre-warmed 150 mM NaCl solution (1 or 1.5:100) and vigorously mixed. PQ was diluted to 60 mg·mL⁻¹ in 150 mM NaCl.

Animals (groups G2 and G4) and were injected i.p. with PQ (30 mg·kg⁻¹). NMDPEF solution (0.5 mg·kg⁻¹ [G4 group] or 0.75 mg·kg⁻¹ [G5 group]) was injected i.p. at 2, 4, 24, 28, 48 and 52 h after PQ treatment. Control animals (G1) were injected with the same amount of vehicle (NaCl solution plus DMSO) at the same times. G3 animals were treated with vehicle (0.9% NaCl) instead of PQ solution, followed by NMDPEF i.p. injections at the same times as the G5 animals. The animals were monitored daily for signs of toxicity for 28 days and longer. A group of surviving treated animals was killed 3.5 months after PQ treatment to assess any late PQ toxicity, such as lung fibrosis.

Microinfusion into the SN

PQ-induced brain toxicity studies were approved by the Italian Ministry of Health according to the law DL 116/92. Animals were housed as described earlier. Rats were anaesthetized with chloral hydrate (400 mg·kg⁻¹ i.p.) and positioned in a stereotaxic frame (D. Kopf Instruments, Tujunga, CA, USA). A stainless steel guide cannulae (25 gauge) was then implanted unilaterally into the SN under stereotaxic guidance and secured to the skull with a dental acrylic. In the

same session, four electrodes were chronically inserted into each frontoparietal cortex to permit recording of ECoG activity (Bagetta *et al.*, 1994). Rats were allowed 2 days of recovery before experiments were carried out. No alteration of posture or motor activity changes have been found over this period. For testing, the animals were placed into individual transparent Perspex cages and allowed to acclimatize to the new environment for 15 min.

PQ was solubilised in PBS. Drugs were solubilised in DMSO and then diluted in PBS. Microinfusion of drugs or vehicle (5% DMSO diluted in PBS) was carried out by means of a 10 µL Hamilton syringe connected by a Teflon tube to an injector cannula. Test compounds were microinjected in a total volume of 1 µL. The injector cannula was left in place for a further 2 min after completing the injection. Drugs or vehicle were infused for 30 min before PQ injection. Each animal was treated only once and monitored continuously for 4 h after treatment. Postural, locomotor changes and convulsive behaviour were recorded by two independent observers who were unaware of the treatment the animals had received.

Malondialdehyde (MDA) determination in brain tissue and isolated cells

MDA standard was prepared by hydrolysis of 1,1,3,3-tetraethoxypropane. Endogenous MDA was determined indirectly by quantifying the content of 2,4-dinitrophenylhydrazone, which is the product of the derivatization reaction of MDA with 2,4-dinitrophenylhydrazine (DNPH).

In order to obtain brain tissue, the animals were killed 24 h after treatment and the SN area excised. 0.5 g of tissue per sample was homogenized in phosphate buffer (0.01 M, pH 7.0). Cultured cells (8 × 10⁶) were treated as required with PQ plus NMDPEF or vehicle (DMSO) and 18 h later cells were resuspended in phosphate buffer by scraping. The biological material was sonicated and then centrifuged at 4500 × g for 5 min. The supernatant was collected and used for the derivatization reaction, which was carried out for 30 min at room temperature. The solid-phase extraction procedure was applied to isolate 2,4-dinitrophenylhydrazone from biological samples.

Quantitative analysis of derivatization reaction products was performed on Agilent 1200 using a 20 µL loop injection valve (Agilent, Santa Clara, CA, USA). The chromatographic system was attached to a diode-array detector. A methanol solution of hydrazone and acid solution of DNPH were injected into the column and identified by their relative retention times determining the maximum wavelength of absorption of the hydrazone at 310 nm. A mixture of H₂O acidified by CF₃COOH (pH = 2.20) and acetonitrile, in ratio of 45:55 (v/v) was used as mobile phase. It was delivered at a flow rate of 1.0 mL·min⁻¹ through a Nucleosil 100-5 RP/18 (25 × 4.6 cm, 4 µm) reverse phase column (Agilent), with a relative guard column (4.5 × 0.46 cm).

Data and statistical analysis

Data are expressed as means ± SEM for the number *n* of independent experiments or as means ± SD for the number *n* of independent samples, from a representative experiment

out of at least 3 independent experiments that yielded comparable results, as indicated for each figure. Numerical data from all the experiments were analysed with ANOVA, followed by Student–Newman–Keuls test. IC_{50} was calculated by XLfit software package.

Materials

NMDPEF, a kind gift from Philippe Delagrangre, Servier, France [100 or 20 mM stock in DMSO], melatonin (Sigma-Aldrich, 500 mM stock in DMSO), apocynin (Sigma-Aldrich, 200 mM stock in EtOH), PQ (Sigma-Aldrich, 100 mM in PBS), rotenone (Sigma-Aldrich, 10 mM in PBS) were stored in aliquots at -80°C , and dilutions were prepared shortly before treatments.

Results

The QR2 inhibitor NMDPEF prevents non-apoptotic PQ-induced cell death in astroglial and epithelial cells

In order to examine a potential involvement of QR2 in PQ-induced toxicity we tested a specific and potent QR2 inhibitor NMDPEF (Mailliet *et al.*, 2005) (Figure 1A), in different cell systems, including lung, brain, kidney and mammary gland-derived cell lines. NMDPEF exerted a marked cytoprotective effect against PQ-induced cell death in human U373 astroglial cells, in embryonic kidney HEK293 cells (Supporting Information Figure S2A and C) and moderate or poor cytoprotective effect in other cell types. These observations may be a reflection of different expression levels of QR2 in different cell lines (Supporting Information Figure S2E).

We chose U373 astroglial cells, which responded very well to NMDPEF, as a model system to evaluate the QR2 inhibitor *in vitro*. We first determined the IC_{50} of the cytoprotective effect of NMDPEF in U373 cells in the presence of lethal doses of PQ (100 μM). This concentration of PQ induced a non-apoptotic cell death in 60–86% U373 cells 72 h after treatment (Supporting Information Figure S1). NMDPEF prevented cell death in a dose-dependent fashion, with a maximum effect at 20–50 μM (Figure 1B) and an IC_{50} = 6.5 μM . We also analysed the protective effect of NMDPEF over time at different PQ concentrations. NMDPEF delayed cell death induced by 100 μM PQ (Figure 1C and D) and also against millimolar concentrations of PQ (Supporting Information Figure S2A). Furthermore, when tested against lower concentrations of PQ (1 and 10 μM) NMDPEF fully prevented cell death over a period of 7 days (Figure 1D).

NMDPEF is structurally related to melatonin and has been reported to inhibit QR2 by binding to the same site on the enzyme as melatonin (Mailliet *et al.*, 2005) and has been shown to inhibit PQ toxicity *in vivo* (Melchiorri *et al.*, 1995). Therefore, we tested if this neurohormone was able to counteract the cytotoxicity of PQ in astroglial cells. Melatonin effectively blocked morphological changes induced by PQ in astroglial cells at 24–48 h, but showed only a modest and transient protection against PQ-induced cell mortality (100 μM) at later time points (Figure 1E). This effect of melatonin demonstrates a 10,000 fold weaker inhibition of QR2

activity, compared with that of NMDPEF (Mailliet *et al.*, 2005). Apocynin, a NADH oxidase inhibitor was also shown to have a significant protective effect against PQ-induced toxicity *in vitro* and *in vivo* (Cristovao *et al.*, 2009). However, apocynin only delayed the onset of PQ-induced cell death for several hours and had only a marginal effect at 72 h (Figure 1F). These observations correlated with our findings that, in U373 cells melatonin and apocynin had no inhibitory effect and melatonin only a weak inhibitory effect on QR2 activity (Figure 2A and B).

PQ is toxic to all organs and tissues, but the major cause of death is respiratory failure mainly as a consequence of damage to alveolar epithelial cell (pneumocytes type I and II) (Dinis-Oliveira *et al.*, 2008). For this reason we evaluated the protective effect of NMDPEF on PQ-induced cell death in pneumocytes. We decided to perform experiments on freshly isolated primary pneumocytes to avoid variability in the response due to genetic alterations present in existing cell lines (Supporting Information Figure S2). It is interesting to note that the QR2 gene is localized on a distal region of chromosome 6 (6p25), which is frequently deleted in many lung tumours (Merlo *et al.*, 1994). Rat alveolar epithelial cells were several times more sensitive to PQ than astroglial cells and 20 μM PQ was sufficient to kill 80% of cells within 72 h (Figure 1G). NMDPEF significantly counteracted cell death induced by high and low doses of PQ (Figure 1G and H).

The effect of NMDPEF and PQ on QR2 activity in vitro

NMDPEF had a significant effect on QR2 activity in U373 cell lysates (IC_{50} between 0.1 and 1 μM). As expected, the inhibitory effect of melatonin (100 μM) was weak, but significant, whereas apocynin did not inhibit QR2 activity (Figure 2A and B). In order to determine whether PQ has any effect on QR2 activity we performed two types of assays. In the first assay we determined QR2 activity from U373 cell lysates in the presence of PQ with or without NMDPEF added to the *in vitro* reaction (Figure 2C). In the second assay, we tested QR2 activity in protein lysates prepared from cells treated for 24 h with PQ with or without the QR2 inhibitor (Figure 2D). PQ added exogenously to the QR2 activity assay did not increase BNAH consumption in the absence of menadione (Figure 2C) and had no significant effect on QR2 activity in the presence of substrate (Figure 2C). Interestingly, we could detect a statistically significant increase in QR2 activity in cells treated for 24 h with PQ (100 μM) (Figure 2D), which could be inhibited by NMDPEF (20 μM). Both assays showed that NMDPEF inhibited QR2 activity, however, the concentration required to detect a significant inhibition of QR2 in whole cells (20 μM) (Figure 2D) was much higher than the concentration required for potent inhibition of QR2 *in vitro* (0.1–1 μM) (Figure 2C).

QR2 inhibition prevents ROS generation in response to PQ and other oxidative agents

Next, we evaluated the ability of NMDPEF to prevent ROS generation in response to PQ, hydrogen peroxide and rotenone. ROS generation was determined by an indirect method based on the mean fluorescence intensity (MFI) measurement by FACS of mitochondrial superoxide-reactive MitoSOX

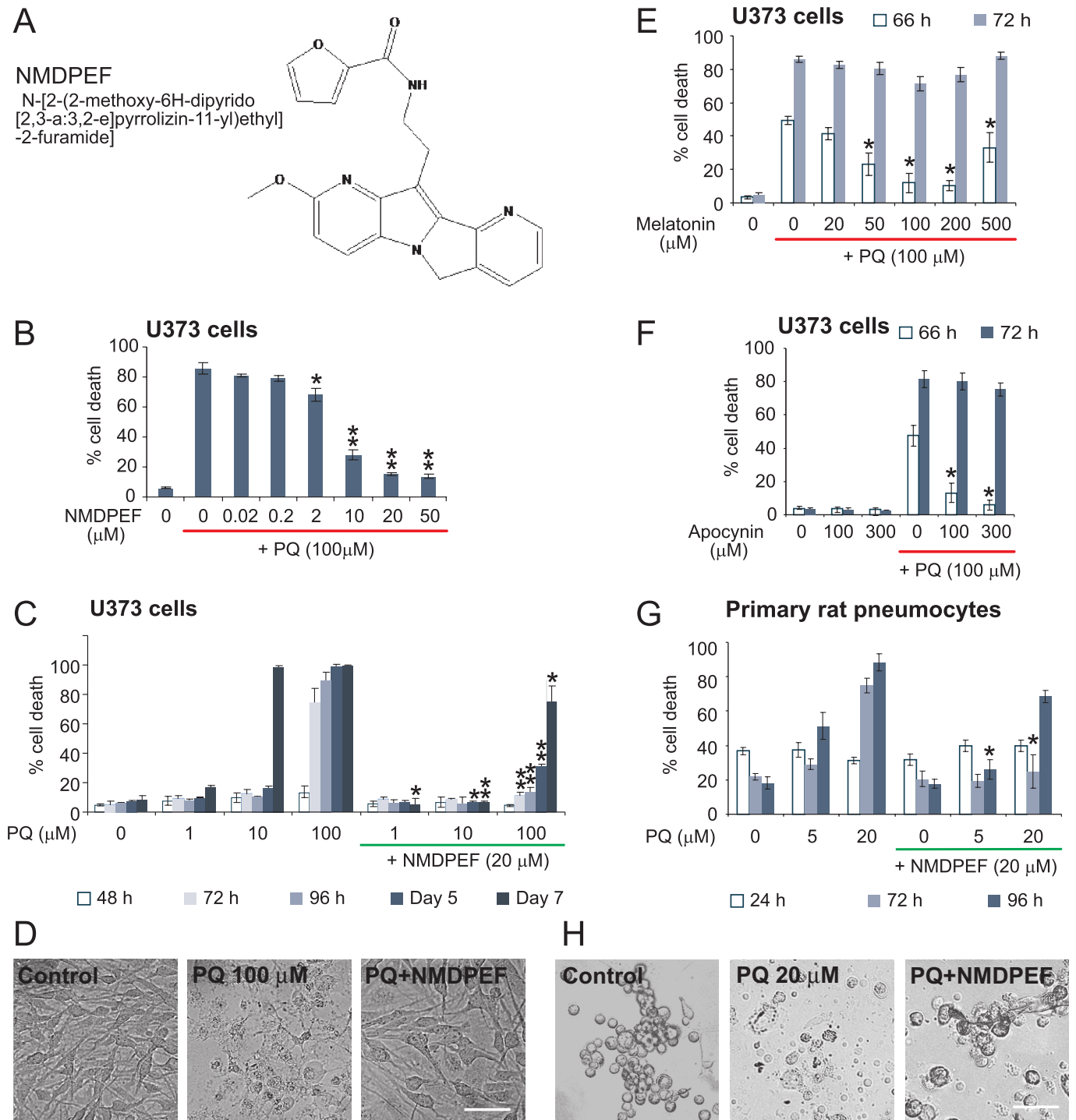


Figure 1

Protective effect of the QR2 inhibitor against PQ-induced cell death *in vitro*. (A) Structure of QR2 inhibitor NMDPEF. (B) Dose-dependent effect of NMDPEF against PQ-induced cell death in human astroglial cells U373. Cells were treated with 100 μM PQ and different concentrations of NMDPEF. Cell death was assessed by Trypan blue and FACS analysis 72 h post-treatment. (C) Long-term protective effect of NMDPEF (20 μM) against different concentrations of PQ was analysed in U373 cells for 7 days. (D) Phase-contrast micrographs of U373 cells treated with 100 μM PQ +/- 20 μM NMDPEF taken 72 h post-treatment. Bar 100 μm . (E,F) Weak and transient protective effect of melatonin (E) and a NADPH inhibitor apocynin (F) against PQ-induced cell death in U373 cells. The experiments were performed as in B but the cells were analysed at 66 and 72 h post-treatment. (G) Protective effect of NMDPEF against PQ-induced cell death in primary rat alveolar epithelial cells. Cell death was assessed by Trypan blue and FACS analysis. (H) Phase-contrast micrographs of primary pneumocytes treated with 20 μM PQ with or without 20 μM NMDPEF for 72 h. Bar 50 μm . The graphs in (B), (E) and (G) show the mean \pm SEM of 3 independent experiments performed in triplicate ($n = 9$). The graphs C and G show the mean \pm SD, $n = 3$, of a representative experiment out of at least three independent experiments performed in triplicate that yielded similar results. * $P < 0.05$, ** $P < 0.01$; significant effect of NMDPEF, $n = 9$.

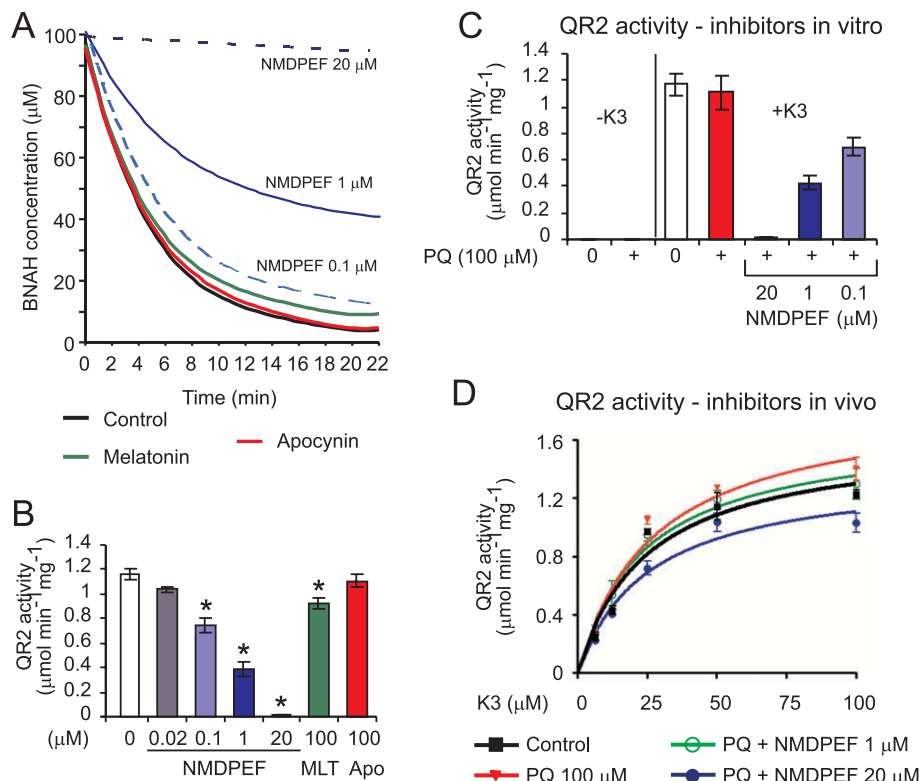


Figure 2

Effect of NMDPEF and PQ on QR2 activity is dependent on the assay. (A) QR2 reaction progress in the presence of substrate K3 (menadione, 100 μM), co-substrate BNAH (100 μM), U373 cell lysate (2.5 μg) and indicated conc. of NMDPEF or melatonin (100 μM) or apocynin (100 μM). The decrease in co-substrate (BNAH) fluorescence was measured by fluorimetry in triplicate. Best-fitted curves to all experimental points are shown. (B) QR2 activity in the presence of NMDPEF, melatonin (MLT) and apocynin (APO) at indicated concentrations. The reactions were carried out as described in (A) and QR2 activity was calculated. The graph shows the mean ± SEM of three independent experiments ($n = 9$). * $P < 0.01$; significantly different from control. (C) No effect of PQ on QR2 activity *in vitro*. U373 cell lysate (1 μg) was incubated with 100 μM PQ and 20, 1 and 0.1 μM NMDPEF in the presence or absence of K3. The graph shows the mean ± SEM of three independent experiments ($n = 9$). (D) Effects of PQ and NMDPEF on QR2 activity in whole cells. Cells were treated for 24 h with PQ (100 μM) with or without NMDPEF (1, 20 μM), lysed and 1 μg of the lysates (in quadruplicate) was assayed in the presence of different concentrations of substrate as indicated above. Michaelis-Menten curves were fitted by non-linear regression. The graph shows a representative experiment out of three independent experiments performed in quadruplicate, means ± SEM. Note, QR2 activity is enhanced in cells exposed to PQ for 24 h and blocked by cytoprotective doses of NMDPEF.

probe. Although MitoSOX does not allow quantitative measurements of superoxide, it is a method of choice for comparative evaluations of oxidative stress in mitochondria (Mukhopadhyay *et al.*, 2007). In U373 cells PQ (100 μM) induced a threefold increase in MitoSOX fluorescence at 24 h and a lower, but significant increase at earlier time points (Figure 3A). Co-treatment with NMDPEF strongly attenuated mitochondrial ROS induction by PQ at each tested time point (Figure 3A and B). Rotenone (10 μM) caused a comparable increase in mitochondrial superoxide levels, but the decrease in MitoSOX-MFI caused by NMDPEF was significantly less pronounced (Figure 3A and B). In line with these findings, NMDPEF delayed or prevented cell death induced by rotenone, but was more effective in preventing PQ-dependent cytotoxicity (Supporting Information Figure S3). The antioxidant activity of NMDPEF was further confirmed by the use of another fluorescent probe, APF, which is sensitive to hydroxyl radicals and NO, but not to hydrogen peroxide levels in the cytoplasm (Setsukinai *et al.*, 2003). PQ induced a maximal shift in cytoplasmic ROS levels around 11 h post-treatment. A

similar effect was observed when H₂O₂ (1 mM) was applied. The increase in APF-MFI was attenuated by NMDPEF, but its antioxidant effect was significantly weaker against H₂O₂-induced oxidative stress (Figure 3C and D).

One consequence of an oxidative insult to whole cells is lipid peroxidation, which we measured by determining the MDA content in cells treated with PQ and NMDPEF. NMDPEF efficiently inhibited the increase in MDA induced by PQ (Figure 3E–H).

These data suggest that NMDPEF is able to attenuate the oxidative stress induced by various stimuli, but with a higher specificity against PQ-induced ROS.

Silencing of QR2 attenuates PQ-induced cell death in astroglial and epithelial cells

Next we determined if QR2 is a key player in PQ-induced toxicity and if NMDPEF exerts its protective activity by binding to QR2. To answer this question we expressed shRNA against QR2, initiated by lentiviral particles, to down-regulate QR2 expression. First we evaluated the ability of four different

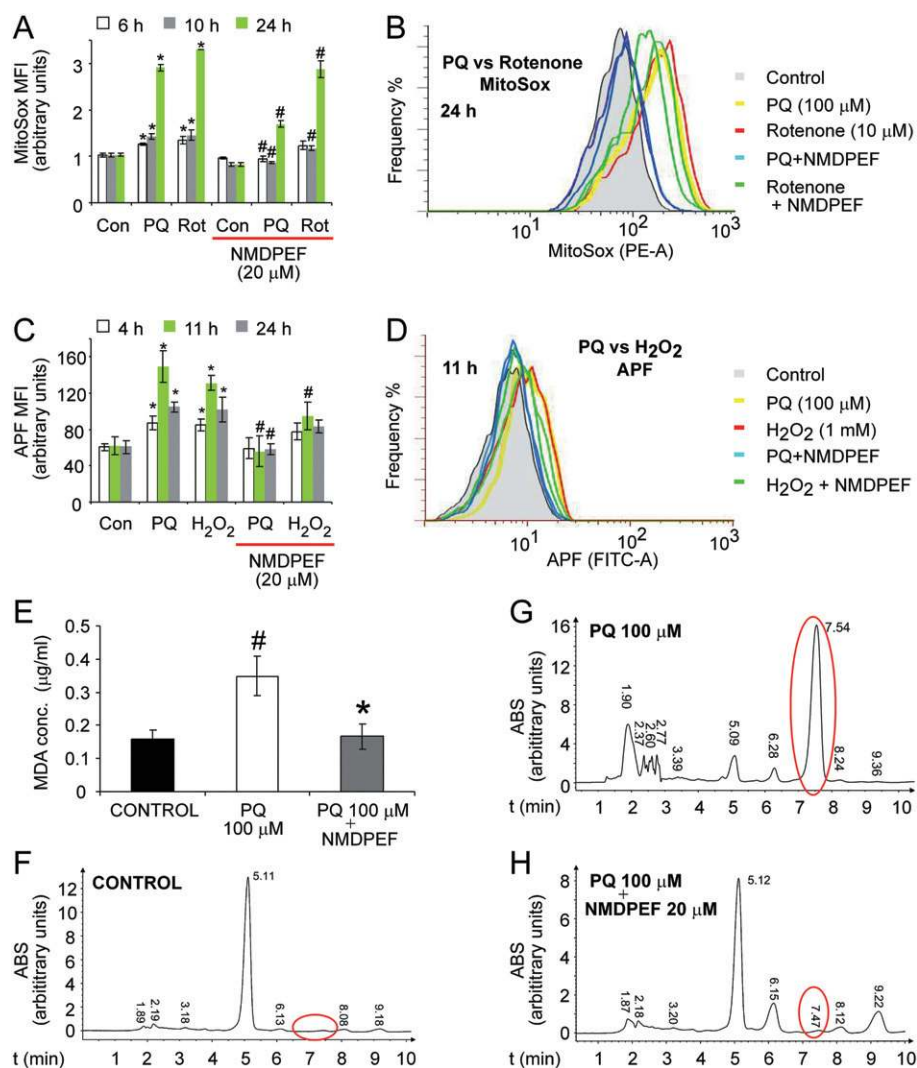


Figure 3

QR2 inhibitor NMDPEF prevents ROS production induced by PQ, hydrogen peroxide and rotenone, but the protective effect is greater against PQ. The oxidative stress was measured by an MFI of MitoSOX using FACS. (A) U373 cells were treated with PQ (100 µM) or rotenone (10 µM) +/– NMDPEF (20 µM). Six, 10, 24 h later the cells were loaded with MitoSOX, trypsinized and analysed by FACS. (B) Representative fluorescence histograms of the experiment shown in A (two histograms for each 24 h treatment). (C) U373 cells were treated with PQ (100 µM) and/or H₂O₂ (1 mM) and APF probe was used to quantify ROS levels. (D) Representative FITC channel fluorescence histograms of the experiment shown in C (two histograms for each 11 h treatment). Data represent the mean ± SEM, *n* = 6, of three independent experiments performed in duplicate. **P* < 0.05, ***P* < 0.01, significant differences between untreated versus PQ-, rotenone- or H₂O₂-treated cells. #*P* < 0.05, significant effect of NMDPEF. (E–H) NMDPEF blocks lipid peroxidation in cells exposed to PQ. (E) Cells were treated for 20 h with 100 µM PQ plus 20 µM NMDPEF or with PQ plus vehicle (DMSO) and compared with control cells (DMSO treated). MDA was measured by HPLC. Mean ± SEM, *n* = 3, of three independent experiments. (F) Representative HPLC profiles of MDA-derivatization products extracted from control cells, (G) cells treated with PQ (100 µM) for 20 h and (H) cells co-treated with indicated concentrations of PQ and NMDPEF for 20 h. ABS, 280 nm absorbance; red circles, MDA derivatization product – 2,4-dinitrophenylhydrazine; #*P* < 0.05, significant difference between untreated and PQ-treated cells. **P* < 0.05, significant effect of NMDPEF.

lentiviral constructs to silence QR2 expression. Two shRNAs were able to decrease strongly QR2 levels in U373 cells (Figure 4A). U373 cells expressing different shRNAs were then tested in an LDH-based cytotoxicity assay 72 h after exposure to PQ. QR2 down-regulation strongly decreased LDH release, compared with the cells with normal QR2 levels (Figure 4B). Although QR2 silencing significantly augmented basal cell mortality in astroglial cells, it strongly attenuated the increase (~20%) in cell death in response to 100 µM PQ, as

determined by FACS analysis (Figure 4C, right panel). However, shRNA QR2 cells did not show a significant resistance to 1 mM PQ (Figure 4C, left panel). In addition, QR2 silencing led to a markedly reduced response to NMDPEF in U373 cells treated with 1 mM and 100 µM PQ (Figure 4C). The higher resistance to PQ and to NMDPEF correlated with lower expression levels of QR2 in cells expressing one of the shRNAs against QR2 – h10 (Figure 4D). We confirmed these observations in a non-transformed mammary epithelial cell

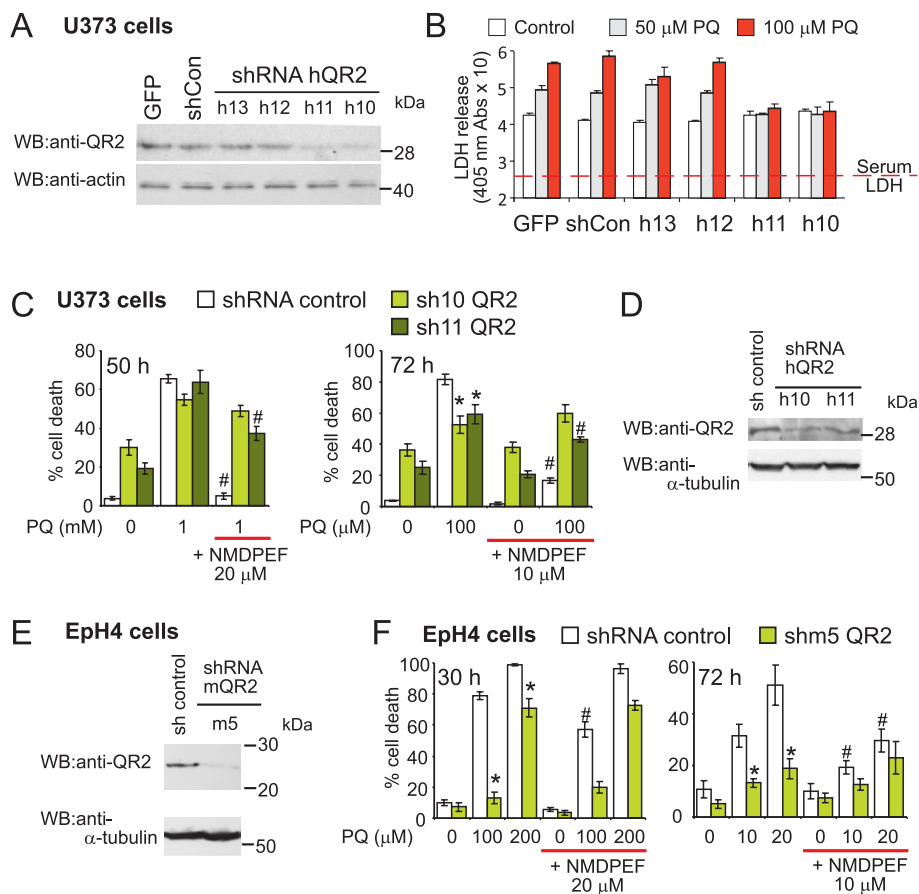


Figure 4

QR2 silencing attenuates cell death induced by PQ. (A) U373 cells were infected with four different lentiviruses encoding shRNA against human QR2 (shQR2) or control lentivirus [Lenti GFP and shRNA control (shCon)] and efficacy of QR2 silencing was assessed by Western blot 5 days after infection. Cell lysate (40 μ g) was run on 8% SDS-PAGE gel. (B) In parallel, the cells analysed in A, were seeded in 24 well plates and incubated in the presence of PQ. Seventy-two hours post-treatment the PQ-induced toxicity was assessed by colorimetric assay for LDH content. This experiment was performed three times and yielded comparable results. Means \pm SD, $n = 4$, of a representative experiment. The values start from the mean serum LDH background absorbance. (C) Human U373 cells infected as in A, with anti-hQR2 shRNA h10 and h11, lentiviruses effective in blocking the expression of QR2 (see Western blot in A), were incubated for 50 h with 1 mM PQ (left panel) or 72 h with 100 μ M PQ (right panel). The PQ treatment was started 8 days post-infection (Day 0). Cell mortality was assessed by Trypan blue and FACS. Means \pm SD, $n = 3$, of a representative experiment out of three experiments that yielded comparable results. (D) Western blot analysis of QR2 expression in U373 cells used in (C). Cells were lysed on day 0 of the experiment showed in C and analysed as in (A). (E) Western blot analysis of QR2 silencing in mouse epithelial cells Eph4. Cells were infected with lentiviruses producing shRNA against mouse QR2 (mQR2) and selected with puromycin for 8 days. Cell lysate (50 μ g) was run on a gradient (4–12%) NuPage gel. (F) High resistance to a high and low dose of PQ in Eph4 shRNA m5 (shm5) expressing cells compared with pLKO-control shRNA cells. Eighty per cent confluent Eph4 cells, were treated with 100 μ M PQ and analysed 30 h later (left panel) or with 10 μ M PQ and analysed 72 h later (right panel) as described for C. Means \pm SD, $n = 3$, of a representative experiment out of three independent experiments, performed in triplicate that yielded comparable results. * $P < 0.05$, statistically significant differences for shRNA QR2 cells versus the 'no target' shRNA pLKO control cells. # $P < 0.05$, significantly different from NMDPEF-untreated cells.

line Eph4 (Supporting Information Figure S2B). Eph4 cells required puromycin selection to achieve a satisfactory down-regulation of QR2 (Figure 4E), but in contrast to astroglial cells, Eph4 QR2 shRNA expressing cells showed lower basal cell mortality compared with control cells. This observation correlated with the marked resistance to PQ-induced cell death in response to high (Figure 4F, left panel) as well as low dose PQ (Figure 4F, right panel), as compared with control (shRNA virus infected and puromycin-resistant cells). Significantly, QR2 knock-down led to a complete loss of the protective effect mediated by NMDPEF in epithelial cells.

NMDPEF prevents systemic toxicity of PQ

As NMDPEF was a potent inhibitor of PQ effects in cultured cells, we decided to test the effects of the compound *in vivo*. In the systemic toxicity studies, animals were injected i.p. with PQ (30 mg·kg⁻¹, $n = 6$) or vehicle. Two and 4 h later animals were injected i.p. with NMDPEF. The injections of NMDPEF were subsequently repeated on days 1 and 2 after the PQ treatment. PQ given i.p. caused visible toxicity in animals (piloerection, tremors, extended limbs, limited motor activity) which was observed the day after treatment. Animal deaths usually occurred between 4 and 14 days after injection

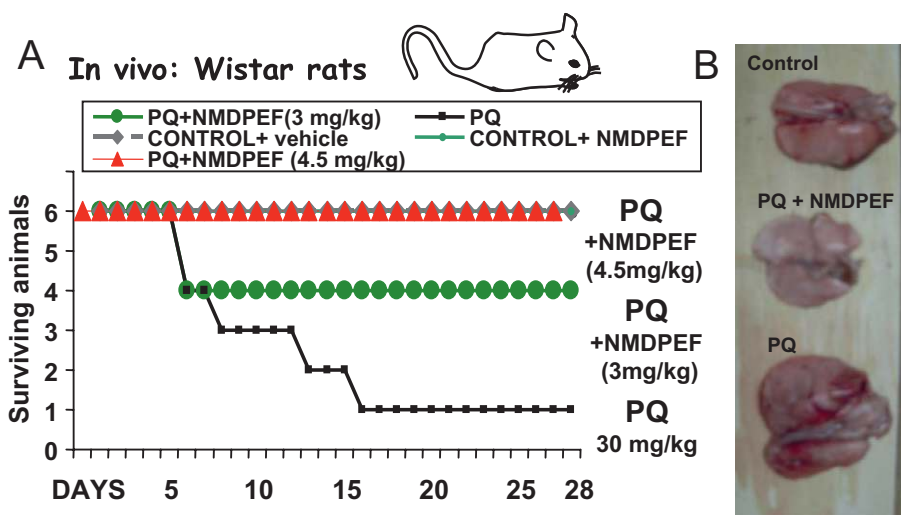


Figure 5

QR2 inhibitor prevents PQ-induced systemic toxicity and mortality in Wistar rats. (A) The animals (six per group) were injected i.p. with PQ ($30 \text{ mg}\cdot\text{kg}^{-1}$) followed by i.p. injections of NMDPEF ($0.5 \text{ mg}\cdot\text{kg}^{-1}$ or $0.75 \text{ mg}\cdot\text{kg}^{-1}$ each) 2 and 4 h later (Day 0). The drug was also administered twice on the Day 1 and twice on the Day 2 post-PQ treatment, $0.5 \text{ mg}\cdot\text{kg}^{-1}$ or $0.75 \text{ mg}\cdot\text{kg}^{-1}$ each time, to the final dose 3 and $4.5 \text{ mg}\cdot\text{kg}^{-1}$ respectively. The animals were monitored daily for signs of toxicity for 28 days post-treatment. (B) The animals surviving after the PQ treatment and controls were killed 3.5 months later and lungs were analysed for macroscopic signs of fibrosis. Note, enlarged lungs and extended areas of fibrosis in animals that survived from PQ treatment and normal size and normal appearance of lungs isolated from animals treated with $4.5 \text{ mg}\cdot\text{kg}^{-1}$ of NMDPEF and PQ.

(Figure 5A). The manifestations of PQ-induced toxicity were markedly reduced in all animals treated with NMDPEF. In addition, NMDPEF reduced ($3 \text{ mg}\cdot\text{kg}^{-1}$) or fully prevented ($4.5 \text{ mg}\cdot\text{kg}^{-1}$) animal mortality (Figure 5A), and potentially reduced lung fibrosis in surviving PQ-treated animals (Figure 5B). Moreover, we did not observe any toxic effects after treatment with NMDPEF alone, as measured by animal behaviour or histological analysis of organs (data not shown).

QR2 prevents direct effects of PQ in nigral areas of the brain

In a second *in vivo* experiment PQ was directly microinfused into the SN. We previously reported that such a treatment leads to electrocorticographic (EcoG), epileptogenic discharges and behavioural stimulation in rats that resembles Parkinsonian movements in humans (Mollace *et al.*, 2003). We tested the ability of NMDPEF to prevent these immediate effects of PQ and compared it with a non-peptidyl SOD mimetic, M40401. The microinfusion of $1 \mu\text{L}$ of vehicle into SN failed to modify the animal behaviour and EcoG recordings (Figure 6A) and did not induce any lethality in treated rats (Figure 6B). In contrast, PQ ($25 \mu\text{g}$) produced increased motor activity, jumping and circling of animals, starting from 27 min post-injection. EcoG showed a pattern of desynchronization with high voltage and long lasting (4 h) epileptogenic discharges (Figure 6A). All the animals died $32 \pm 2.5 \text{ h}$ after the microinfusion of PQ into SN. Pre-treatment of rats with NMDPEF or M40401 ($10 \mu\text{g}$ infused into SN; $n = 10$) antagonized motor convulsion, EcoG epileptogenic discharges (Figure 6A) and the mortality of rats receiving PQ (Figure 6B). In particular, NMDPEF ($10 \mu\text{g}$; $n = 10$) injected into SN 30 min before PQ administration, prolonged the latency of symptoms (Figure 6C) and reduced PQ-related

mortality of rats by 80% (Figure 6B). This observation correlated with a strong reduction in MDA levels in the nigrostriatal areas of the brain in animals treated with NMDPEF, compared with PQ-treated animals (Figure 6D). The overall protective effect of NMDPEF was comparable with M40401, in terms of reduction of mortality, latency of symptoms and lipid peroxidation (Figure 6B–D), suggesting that the inhibition of QR2 strongly attenuated the oxidative stress elicited by PQ in this SN toxicity model.

Discussion and conclusions

This report shows that NMDPEF, a novel, melatonin-related tetracyclic compound, previously characterized as a potent inhibitor of QR2 (also known as NQO2) (Mailliet *et al.*, 2005), prevents PQ-induced toxicity *in vitro* and *in vivo* in rat models of systemic and brain toxicity. NMDPEF is several times more potent than melatonin and apocynin, which have been previously shown to antagonize PQ-induced toxicity (Melchiorri *et al.*, 1995; Cristovao *et al.*, 2009). The protective effect of QR2 knock-down in astroglial and epithelial cells confirms the involvement of this enzyme in the complex toxic mechanisms elicited by PQ (Figure 4). In addition, the reduced sensitivity to the QR2 inhibitor, following QR2 silencing in astroglial cells, as well as the lack of response to NMDPEF in epithelial cells, suggested that QR2 was the main target of NMDPEF in the protection against PQ (Figure 4).

To our knowledge this report is the first to suggest a role of QR2 in PQ-induced toxicity, although QR2 has been implicated in the toxic effects of other xenobiotics. In fact, in contrast to QR1, which usually works as a detoxifying enzyme, QR2 was found to mediate the toxicity of exogenous

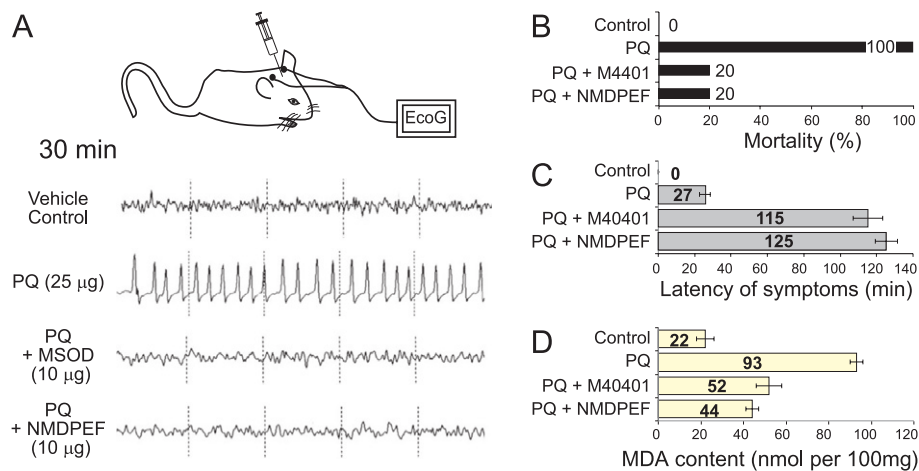


Figure 6

NMDPEF prevents ECoG discharges, lipid peroxidation and death induced by the infusion of PQ into the SN of Wistar rats. (A) Representative traces showing high voltage epileptogenic discharges induced by the infusion of PQ (25 µg) or vehicle with or without pre-treatment (20 min) with NMDPEF or a MnSOD mimetic M40401 (10 µg). Traces are representative of ECoG effect observed 30 min after administration of the herbicide. (B) Mortality following the infusion of PQ (25 µg, $n = 10$) and NMDPEF or M40401 as in A ($n = 10$, each treatment). (C) The effect of NMDPEF and M40401 (10 µg) on the latency (min) of seizures induced by PQ (25 µg). Mean \pm SEM, $n = 10$. (D) MDA accumulation in the brain tissue in rats receiving PQ (25 µg) with or without pre-treatment with NMDPEF or M40401 as in A and B. SN homogenates were subjected to derivatization and analysed by HPLC Data shown are means \pm SEM, $n = 10$. * $P < 0.05$, significantly different from PQ alone.

quinones and other compounds. For example, in the case of menadione, a cytotoxic synthetic analogue of vitamin K, QR2 is a bio-activating enzyme such that cells deficient in QR2 and also QR2 knock-out mice are resistant to menadione toxicity. Furthermore, activation of QR2 with NRH, a non-biogenic co-substrate, augments the toxic effects of menadione (Long *et al.*, 2002; Celli *et al.*, 2006). QR2 also bioactivated mitomycin C leading to the generation of the cytotoxic, nitroreductase anti-tumour drug CB1954. Keratinocytes isolated from skin tumours of QR2 knock-out mice show increased resistance to CB1954 and mitomycin C, whereas addition of NRH or overexpression of QR2 enhances drug toxicity (Celli *et al.*, 2006).

Our data suggest that PQ-induced toxicity is mediated via QR2 activity. Several other oxidoreductases have also been implicated in PQ bioactivation, such as NADPH oxidases of the Nox family (Miller *et al.*, 2007; Cristovao *et al.*, 2009), a DPI-sensitive thioredoxin reductase (Gray *et al.*, 2007), as well as a mitochondrial NADH-dependent reductase (Shimada *et al.*, 2009). However, in spite of a wide specificity of QR2 in terms of substrates and co-substrates, it is unlikely that QR2 directly catalyses the conversion of the PQ di-cation into a PQ radical, thus using PQ as a substrate in a mechanism analogous to the diaphorases listed above. This is a likely explanation because QR2, just as its homologue QR1, is considered to be a two-electron oxidoreductase mediating the transfer of two or four electrons from a co-substrate to a substrate, no matter whether it is a quinone reduction or nitroreduction (Wu *et al.*, 1997; Chen *et al.*, 2000). However, an alternative, yet widely accepted mechanism of PQ reduction involves the single electron transfer on the nitrogen of the pyridine ring (Drechsel and Patel, 2008), which is rather incompatible with the obligatory two-electron transfer mediated by the FAD catalytic centre of QR enzymes (Li *et al.*, 1995). In fact, PQ added to the

QR2 activity fluorometric assay using BNAH as substrate, did not increase BNAH consumption and had no significant effect on BNAH decay *in vitro* in the presence of menadione (Figure 2C). This observation suggested that PQ was not a QR2 substrate and any positive effect on QR2 activity, such as the one observed after 24 h treatment (Figure 2D), was indirect and cell-system dependent. Therefore, even though it has been suggested that QR2 may be able to catalyse one electron reductions (Vella *et al.*, 2005) we would exclude QR2 as a candidate enzyme directly involved in PQ redox cycling. Further experiments should clarify the indirect role of QR2 in PQ redox cycling and PQ-induced toxicity.

QR2 may play a more general role in oxidative stress regulation. In fact, NMDPEF attenuated rotenone and hydrogen peroxide-induced ROS levels *in vitro* (Figures 2 and Supporting Information Figure S3). QR2 stabilizes the transcription factor p53 by a MDM2 ubiquitination-independent mechanism (Gong *et al.*, 2007) and p53 is an important cellular mediator of the oxidative stress response (Vousden and Ryan, 2009). Moreover, QR2 knock-down and knockout cells have an increased resistance to benzopyrene-mediated cell death and decreased basal cell mortality, thereby facilitating the survival of cells with DNA damage (Gong *et al.*, 2007) and decreasing sensitivity to chemical carcinogens (Shen *et al.*, 2010). Finally, a function of QR2 in PQ-induced toxicity might be even more indirect and could be related to a potential role in the regulation of cellular metabolism, similar to mitochondrial aconitase. Aconitase has been shown to mediate PQ toxicity not only via release of oxidized iron from its 4Fe-S clusters and thereby favouring Fenton reactions, but also by its active role in the Krebs cycle and respiration. Indeed, knock-down of aconitase caused a decrease in cellular metabolism, which protected against PQ-mediated cell death (Cantu *et al.*, 2011).

QR2 has been proposed to act as the third melatonin receptor MT3 in the cell (Nosjean *et al.*, 2000; Mailliet *et al.*, 2004). The relationship between the many physiological functions of melatonin and this enzyme remains unclear, but the weak inhibitory effect of melatonin on QR2 may explain the anti-oxidant properties of this neurohormone (Vella *et al.*, 2005; Luchetti *et al.*, 2010). Melatonin exhibited significant protection against systemic toxicity of PQ in two rat strains (Melchiorri *et al.*, 1995). We did not directly compare the effects of melatonin and NMDPEF *in vivo*, but considering the high doses of melatonin necessary to attenuate PQ-induced mortality in rats, it is likely that NMDPEF is much more potent *in vivo*, giving full protection at 4.5 mg·kg⁻¹ (Figure 5). In addition, melatonin was only partially cytoprotective in astroglial cells when used at ~100 μM (Figure 1E). These findings are in agreement with a weak inhibitory activity of melatonin on QR2 activity (IC₅₀ = 130 ± 35 μM). In contrast, NMDPEF has been structurally designed to resemble melatonin and bind to the melatonin-binding site on QR2. It is a very powerful QR2 inhibitor (IC₅₀ = 14 nM) (Mailliet *et al.*, 2005) and a potent blocker of PQ-induced toxicity as shown here.

Plant flavonoids such as resveratrol and quercetin also act as QR2 inhibitors (Vella *et al.*, 2005). Interestingly, both compounds have been shown to protect against PQ-induced toxicity and even though these effects have been explained by other mechanisms (Park *et al.*, 2010; Robb and Stuart, 2011), it is likely that some of the antioxidant activity of flavonoids depends on their interaction with QR2.

Equimolar concentrations of QR2 inhibitor compared with PQ (10 μM) were required for full protection against cell death *in vitro* in U373 cells (Figure 1C). Interestingly, NMDPEF was effective at much lower concentrations than PQ *in vivo* in preventing the systemic and nigral toxicity of PQ in the majority of animals, suggesting that NMDPEF had better pharmacokinetic properties than PQ (Figure 5). NMDPEF exhibited no inherent toxicity at 4.5 mg·kg⁻¹ *in vivo* and was effective even if administered after PQ. This profile suggests that NMDPEF could be a candidate for the therapeutic management of PQ poisoning. Indeed, the current methods of clinical intervention in cases of PQ poisoning are often unsuccessful due to the lack of effective treatments (Dinis-Oliveira *et al.*, 2009a). Recently, sodium salicylate (Dinis-Oliveira *et al.*, 2007) as well as lysine acetylsalicylate (Dinis-Oliveira *et al.*, 2009b), commonly available non-steroidal anti-inflammatory drugs, have been proposed as promising antidotes for the treatment of PQ poisoning. The modulation of pro-oxidant and pro-inflammatory pathways has been suggested as essential features of the beneficial effects of both sodium salicylate and lysine acetylsalicylate. However, in animal models, high doses of these drugs (~200 mg·kg⁻¹) are required to protect against PQ (25 mg·kg⁻¹ i.p.) toxicity. Furthermore, in these animals, the compounds themselves demonstrated inherent toxicity at 400 mg·kg⁻¹ kg⁻¹ suggesting a limited therapeutic window for these agents in the treatment of PQ poisoning (Dinis-Oliveira *et al.*, 2009b). Other anti-inflammatory and ROS scavenging drugs are used in current PQ detoxification protocols, including dexamethasone, methylprednisolone, cyclophosphamide and N-acetylcysteine. All these agents seem to offer only a limited benefit for patients with acute PQ poisoning (Dinis-

Oliveira *et al.*, 2009a). Other compounds able to attenuate the toxic effects of PQ *in vitro*, such as apocynin and DPI (Cristovao *et al.*, 2009; Gray *et al.*, 2007), show a protective effect specific for SN dopaminergic degeneration *in vivo* triggered by low dose PQ (Cristovao *et al.*, 2009), yet their efficacy against acute systemic PQ-induced toxicity has not been addressed.

Thus the data presented here suggest a novel pharmacological approach to the treatment of acute PQ poisoning. Future studies should elucidate the molecular mechanisms of the antidote-like effect of NMDPEF and the role of QR2 in the regulation of oxidative stress.

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Conflict of interest

All authors declare no competing financial interests.

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Supporting information

Additional Supporting Information may be found in the online version of this article:

Figure S1 The cell death induced by PQ in U373 and EpH4 cells is not apoptosis. (A) The 48 h PQ treatment failed to induce significant apoptosis in U373 cells. Annexin V staining was performed as above 48 h after the addition of PQ. Mean \pm SEM, $n = 3$. (B) The 18 h PQ treatment (1 and 2 mM) did not induce any significant increase in Annexin V positive

astroglial cells. Cells were trypsinized, stained with PE-labelled Annexin V and 7-AAD to monitor dead cells and analysed by FACS. After gating the populations the percentage of 7-AAD and Annexin V-positive cells was determined. (C) U373 cells, exposed to PQ for 60 h, were treated for 20 min with PE-fmk-zVAD caspase-3 inhibitor (Casp3* Red), recognizing the activated caspase-3. After trypsinization, cells were co-stained with Trypan blue and analysed by FACS. Mean \pm SEM, $n = 3$. (D) PQ failed to induce caspase-3 activation in highly sensitive mammary epithelial cells EpH4. Cells were stained as in C and analysed by FACS. Mean \pm SEM, $n = 3$. (E) Representative dot-plots of double PE-fmk-zVAD (Casp3* Red) and Trypan blue staining from the experiment shown in C.

Figure S2 The protective effect of QR2 inhibitor NMDPEF on PQ-induced cell death correlates with QR2 expression in different cell types. (A) U373 astroglial cells were seeded at standard conditions and treated with PQ (as above) for indicated times and co-treated with 20 μ M NMDPEF. The percentage of dead cells was analysed by Trypan blue and FACS. A representative experiment out of 3 independent experiments performed in triplicate, Mean \pm SD, $n = 3$; * $P < 0.05$, compared with the time-matched PQ alone controls. (B) EpH4 cells were seeded as indicated in methods and treated with PQ as indicated above. The cell death analysis was performed on day 5 after seeding as in A. * $P < 0.05$ and ** $P < 0.01$ compared with the time-matched PQ alone controls. (C) 6×10^4 HEK293 cells were seeded per well (24 well plates), 24 h before the addition of PQ and NMDPEF. The cells were exposed for indicated times to indicated agents. The graph shows the mean of independent experiments performed in triplicate, Mean \pm SD, $n = 6$; * $P < 0.05$ and ** $P < 0.01$ compared with the time-matched PQ alone controls. (D) A549 lung adenocarcinoma cells were seeded at a density 4×10^4 per well (24 well plate). The cells were exposed for indicated times to PQ and NMDPEF added next day after seeding and analysed as above. The graph shows the mean of independent experiments performed in triplicate, Mean \pm SD, $n = 9$. (E) The expression levels of QR2 in different cell lines compared with rat liver and lung tissue. Around 80 μ g of cell lysate were run on 10% SDS-PAGE and QR2 expression was analysed by Western blot. GAPDH expression is shown as loading control. Note the highest expression of QR2 in HEK293 cells and U373 cells that respond very well to NMDPEF and a very low expression in A549 cells, which are not protected by NMDPEF.

Figure S3 QR2 inhibitor NMDPEF protects against PQ and rotenone toxicity, but the protective effect is much stronger for PQ. The cells were treated with PQ (10 μ M) or rotenone (10 μ M) and the percent of dead cells was determined by Trypan blue staining and FACS analysis at time points indicated. Mean \pm SEM of 3 independent experiments performed at least in duplicate, $n = 6$, * $P < 0.05$, compared with the time-matched rotenone alone, # $P < 0.01$, compared with the time-matched PQ alone.