

RESEARCH PAPER

Inhibition of Na⁺-H⁺ exchange as a mechanism of rapid cardioprotection by resveratrol

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BACKGROUND AND PURPOSE

Resveratrol is a polyphenol abundantly found in grape skin and red wine. In the present study, we investigated whether resveratrol exerts protective effects against cardiac ischaemia/reperfusion and also explored its mechanisms.

EXPERIMENTAL APPROACH

Infarct size and functional recovery in rat isolated perfused hearts subjected to no-flow global ischaemia followed by reperfusion were measured. Cultured neonatal rat cardiomyocytes were exposed to H₂O₂ (100 μmol·L⁻¹) to induce cell injury. Intracellular ion concentrations were measured using specific dyes. Western blotting was used to quantify protein expression levels.

KEY RESULTS

In rat isolated perfused hearts, treatment with resveratrol (20 and 100 μmol·L⁻¹) 15 min before ischaemia considerably improved left ventricular functional recovery and infarct size. In cultured neonatal rat cardiomyocytes, resveratrol significantly attenuated the increase in reactive oxygen species (ROS) and loss of mitochondrial inner membrane potential. Resveratrol also suppressed the increase in intracellular concentrations of Na⁺ ([Na⁺]_i) and Ca²⁺ ([Ca²⁺]_i) after H₂O₂ application; however, it did not suppress the ouabain-induced [Ca²⁺]_i increase. By measuring changes in intracellular pH recovery after acidification, we also confirmed that acid-induced activation of the Na⁺-H⁺ exchanger (NHE) was prevented by pretreatment with resveratrol. Furthermore, resveratrol inhibited the H₂O₂-induced translocation of PKC-α from the cytosol to the cell membrane; this translocation is believed to activate NHE.

CONCLUSION AND IMPLICATIONS

Resveratrol exerts cardioprotection by reducing ROS and preserving mitochondrial function. The PKC-α-dependent inhibition of NHE and subsequent attenuation of [Ca²⁺]_i overload may be a cardioprotective mechanism.

Abbreviations

ΔΨ_m, mitochondrial inner membrane potential; GAPDH, glyceraldehyde 3-phosphate dehydrogenase; LV, left ventricular; LVDP, left ventricular developed pressure; mitoK_{ATP}, mitochondrial ATP-sensitive K⁺ channels; mPTP, mitochondrial permeability transition pore; NCX, Na⁺-Ca²⁺ exchanger; NHE, Na⁺-H⁺ exchanger; PI, propidium iodide; PMA, phorbol 12-myristate 13-acetate; ROS, reactive oxygen species; TMRE, tetramethylrhodamine ethyl ester

Introduction

Resveratrol, a polyphenol abundantly found in red wine and grapes, has been reported to exert beneficial effects such as

anti-inflammation, inhibition of tumour cell growth, extension of lifespan and cardioprotection against ischaemia/reperfusion injury. Several mechanisms of these effects have been proposed, such as antioxidative action (Ray *et al.*, 1999;

Danz *et al.*, 2009), up-regulation of NO synthesis (Hattori *et al.*, 2002; Das *et al.*, 2005) and activation of the sirtuin-1 (Sirt1) pathway (Chen *et al.*, 2009). However, the precise mechanisms of resveratrol-induced cardioprotection have not yet been established.

The sarcolemmal $\text{Na}^+\text{-H}^+$ exchanger (NHE) plays a role in regulation of intracellular pH (pH_i) by H^+ extrusion driven by the transmembrane Na^+ gradient in cardiomyocytes. NHE activity is increased during ischaemia in response to intracellular acidosis (Avkiran and Marber, 2002). Increases in the intracellular concentration of Na^+ ($[\text{Na}^+]_i$) induced by increased $\text{Na}^+\text{-H}^+$ exchange facilitate Ca^{2+} influx through the $\text{Na}^+\text{-Ca}^{2+}$ exchanger (NCX) working in reverse mode and exacerbate cell injury (transporter nomenclature follows Alexander *et al.*, 2011). Hence, the inhibition of NHE is expected to exert cardioprotection by indirectly limiting increases in intracellular concentration of Ca^{2+} ($[\text{Ca}^{2+}]_i$). It has also been reported that several NHE inhibitors protected cardiomyocytes subjected to oxidative stress and ischaemia/reperfusion with consistent improvement in functional recovery (Scholz *et al.*, 1995; Knight *et al.*, 2001; Teshima *et al.*, 2003; Garcia-rena *et al.*, 2011). Furthermore, a clinical study revealed that the inhibition of NHE during reperfusion therapy preserved cardiac function in patients with acute myocardial infarction (Rupprecht *et al.*, 2000). In addition, the sub-analysis of a large clinical trial showed that the use of cariporide, a NHE inhibitor, in patients who underwent high-risk coronary artery bypass graft surgery reduced the incidence of myocardial infarction and mortality (Boyce *et al.*, 2003). Recently, resveratrol has been reported to decrease the expression of the NHE isoform 1 (NHE-1; SLC9A1) by repressing its promoter in rat myoblasts (Jhumka *et al.*, 2009). This result indicates a possible link between resveratrol and the inhibition of NHE. In the present study, we investigated whether the inhibition of NHE is a possible mechanism of the cardioprotective effects of resveratrol.

Methods

All animal care and experimental procedures were performed in accordance with the *Guide for the Care and Use of Laboratory Animals* published by the US National Institutes of Health (NIH Publication No. 85-23, revised 1996) and the Oita University animal care guidelines.

Isolated perfused heart experiments

Isolated perfused heart experiments and infarct size measurements were performed using the Langendorff apparatus. Sprague-Dawley rats (250–300 g) were anaesthetized with an intraperitoneal injection of a mixture of ketamine (60 $\text{mg}\cdot\text{kg}^{-1}$) and xylazine (10 $\text{mg}\cdot\text{kg}^{-1}$). After the depth of anaesthesia was confirmed by stable heart rate and a lack of flexor responses to a paw pinch, rat hearts were isolated and perfused in retrograde mode at 37.0°C under a constant pressure of 75 mmHg. After stabilization for 20 min, the hearts were subjected to no-flow global ischaemia for 20 min, followed by reperfusion for 30 min. In the resveratrol-treated group, the hearts were perfused with 20 $\mu\text{mol}\cdot\text{L}^{-1}$ resveratrol in Krebs-Henseleit buffer (composition (in $\text{mmol}\cdot\text{L}^{-1}$) 118

NaCl , 4.7 KCl , 2.5 CaCl_2 , 1.2 MgSO_4 , 1.2 KH_2PO_4 , 25 NaHCO_3 , 10 glucose, 10 HEPES) 15 min before ischaemia. Left ventricular (LV) pressure was monitored using a pressure transducer to record the peak positive and negative first derivatives of LV pressure ($\text{dP}/\text{dt}_{\text{max}}$ and $\text{dP}/\text{dt}_{\text{min}}$ respectively). LV developed pressure (LVDP) was defined as the difference between LV systolic and diastolic pressures. LV pressure, coronary perfusion pressure and electrocardiograms were continuously recorded using a polygraph recorder (Nihon Kohden, Tokyo, Japan). Data were analysed using Lab Chart software (ADInstruments, Colorado Springs, CO, USA). At the end of reperfusion, the hearts were rapidly removed from the Langendorff apparatus and sliced across the long axis of their left ventricles into 2-mm-thick transverse sections. These slices were incubated in 1% triphenyltetrazolium chloride in a phosphate buffer (0.1 $\text{mol}\cdot\text{L}^{-1}$; pH 7.4) at 37°C for 20 min. Infarct areas were enhanced by storage in 10% formaldehyde solution for 24 h and determined by planimetry of each slice. The volumes were calculated using image analysis software (Image J) by multiplying each area by the slice thickness and summing them for each heart.

Primary culture of neonatal rat cardiomyocytes

Cardiac ventricular myocytes were prepared from Sprague-Dawley rats aged 1–3 days and cultured as described elsewhere (Teshima *et al.*, 2003). In brief, rat pups were killed by decapitation and the cardiac ventricles were removed and digested with trypsin (2 $\text{mg}\cdot\text{mL}^{-1}$) at 37°C . The cardiomyocytes were then isolated and cultured in Dulbecco's modified Eagle's minimal essential medium (DMEM) supplemented with 5% fetal bovine serum, penicillin (100 $\text{U}\cdot\text{mL}^{-1}$), streptomycin (100 $\text{mg}\cdot\text{mL}^{-1}$), vitamin B_{12} (2 $\mu\text{g}\cdot\text{mL}^{-1}$) and bromodeoxyuridine (0.1 $\text{mmol}\cdot\text{L}^{-1}$). These cells were placed in serum-free DMEM for 24 h before the experiments. In each isolation, we harvested cardiomyocytes from 11 neonatal rats, which were divided into appropriate groups. An experiment from one isolation was counted as $n = 1$.

Assay for cell death

For quantification of cellular viability, the cardiomyocytes were stimulated with hydrogen peroxide (H_2O_2) (100 $\mu\text{mol}\cdot\text{L}^{-1}$) for 2 h with or without 20 $\mu\text{mol}\cdot\text{L}^{-1}$ resveratrol (Sigma, St Louis, MO, USA), which was administered 1 h before the application of H_2O_2 . The cardiomyocytes were stained with 50 $\mu\text{g}\cdot\text{mL}^{-1}$ propidium iodide (PI) (Roche, Basel, Switzerland), and images were captured by confocal microscopy (543 nm He/Ne laser line) using a $\times 20$ objective lens (Carl Zeiss, Oberkochen, Germany). After scanning, the cells were permeabilized with saponin (50 $\mu\text{mol}\cdot\text{L}^{-1}$) (Sigma) to calculate the percentage of PI-positive cells in each group.

Assay for mitochondrial inner membrane potential

To assay mitochondrial inner membrane potential ($\Delta\Psi_m$), FACS analysis was performed. The cardiomyocytes were stimulated by exposure to H_2O_2 (100 $\mu\text{mol}\cdot\text{L}^{-1}$) for 30 min with or without incubation with 20 $\mu\text{mol}\cdot\text{L}^{-1}$ resveratrol (except for the experiments using various concentrations of resveratrol), which was added 1 h before H_2O_2 (except for the

experiments in various pre-incubating durations with resveratrol). Other chemicals were added 30 min before resveratrol. Tetramethylrhodamine ethyl ester (TMRE; 100 nmol·L⁻¹) (Invitrogen, Carlsbad, CA, USA) was loaded into the cells at 37°C for 30 min. The cells were harvested by trypsinization and analysed using the FACSCalibur system (Becton Dickinson, Bedford, MA, USA). Fluorescence intensity was monitored at 585 nm (FL-2 channel).

Assay for reactive oxygen species

We also investigated the changes in levels of reactive oxygen species (ROS) by FACS analysis. Resveratrol (20 µmol·L⁻¹) was added 1 h before H₂O₂ application. The cardiomyocytes were loaded with 2 µmol·L⁻¹ CM-H₂DCFDA (Invitrogen) in the absence or presence of 100 µmol·L⁻¹ H₂O₂ and incubated at 37°C for 20 min. The cells were harvested by trypsinization and fluorescence intensity was monitored at 530 nm (FL-1 channel). Data were analysed using the WinMDI analysis software.

Assays for [Na⁺]_i and [Ca²⁺]_i

CoroNa Green (1 µmol·L⁻¹) and Fluo-4 (1 µmol·L⁻¹) (Invitrogen) were used to examine [Na⁺]_i and [Ca²⁺]_i levels respectively. The cardiomyocytes were loaded with dyes at 37°C for 30 min. The scanning began immediately after the application of 100 µmol·L⁻¹ H₂O₂, and fluorescence intensity was sequentially monitored every 5 min by confocal microscopy (488 nm argon laser line; ×40 objective lens) (Carl Zeiss).

Measurement of pH_i

pH_i was measured by monitoring the fluorescence of a pH indicator, BCECF-AM (Invitrogen). Cardiomyocytes were plated on glass-bottomed dishes and loaded with BCECF-AM (2 µmol·L⁻¹) for 15 min at room temperature in Krebs-HEPES solution (in mmol L⁻¹: 135 NaCl, 5.9 KCl, 1.5 CaCl₂, 1.2 MgCl₂, 11.6 HEPES, 11.5 D-glucose). The cells were dually excited at wavelengths of 488 and 458 nm with a fixed emission at 540 nm, and images were sequentially acquired at 0.5 s intervals using a confocal microscope system (LSM 710, Carl Zeiss). At the end of the experiments, pH_i was calculated from the ratio of the two fluorescence intensities (488 nm/458 nm) using the nigericin high-K⁺ protocol (Thomas *et al.*, 1979). In this protocol, the cells were superfused with a high-K⁺ solution (in mmol L⁻¹: 145 KCl, 5 NaCl, 0.01 EDTA, 5 HEPES) at varying pH levels ranging from 5.86 to 8.5 in the presence of 5 mg·L⁻¹ nigericin. We confirmed that the ratios of fluorescence intensity linearly correlated with the pH_i levels in a preliminary experiment.

Assays for NHE activity

NHE activity was measured by monitoring the recovery rate of pH_i from rapid acidification using the NH₄Cl pre-pulse technique as described elsewhere (Sabri *et al.*, 1998). The cells were superfused with Krebs-HEPES solution for 5 min, followed by NH₄Cl solution (25 mmol·L⁻¹) for 6 min; thereafter, the solution was replaced with Na⁺-free Krebs-HEPES solution. After 10 min of perfusion, the perfusate was switched back to Na⁺-containing Krebs-HEPES solution and the cells were superfused for 5 min. The solution in the dishes was continuously changed during the experiment by perfusing

fresh solution while aspirating the old solution at the same rate (5 mL·min⁻¹) using a Harvard syringe system (Harvard Apparatus, Holliston, MA, USA). The change in pH_i was sequentially analysed. During superfusion with Na⁺-free Krebs-HEPES solution, pH_i was stable at an acidic level and did not recover until the perfusate was switched back to Krebs-HEPES solution. The rate of pH_i recovery was operationally defined as NHE activity. We quantified this rate by measuring the slope of a straight line fitted to the initial 1 min from the onset of recovery. Resveratrol was added 1 h before the experiment. H₂O₂ and other reagents were added from the beginning of perfusion with NH₄Cl and continued until the end of the experiment.

Western immunoblot analysis

The cells were incubated with resveratrol for 1 h. H₂O₂ and phorbol 12-myristate 13-acetate (PMA) were loaded 10 min before cell harvesting. Membranous and cytosolic proteins were extracted using the Subcellular Proteome Extraction Kit (Merck, Darmstadt, Germany) according to the manufacturer's instructions. Western immunoblot analyses were performed as previously described (Thuc *et al.*, 2010). The primary antibody for PKC-α was purchased from Cell Signaling Technology (Danvers, MA, USA). Glyceraldehyde 3-phosphate dehydrogenase (GAPDH) was used as an internal control. Equal amounts of protein were fractionated on 8.5% SDS-polyacrylamide gels and transferred electrophoretically onto a polyvinylidene difluoride membrane. The membranes were blocked with 5% non-fat milk in Tris-buffered saline containing 0.05% Tween 20 for 1 h at room temperature, and incubated with primary antibodies at a 1:1000 dilution for 1 h at room temperature. After washing, the membranes were incubated with a 1:4000 dilution of horseradish peroxidase-conjugated secondary antibodies for 1 h at room temperature. The blots were visualized by enhanced chemiluminescence and exposure to Hyper-film. Band signals were quantified densitometrically using Image J software. All the band signals were normalized by the GAPDH protein levels and the values were expressed as percentage of the mean value of the control.

Statistical analysis

All data are expressed as mean ± SE. Multiple comparisons among groups were performed by one-way ANOVA with Bonferroni *post hoc* test. Sequential changes were analysed by two-way ANOVA. A value of *P* < 0.05 was considered statistically significant.

Results

LV functional recovery and infarct size

At baseline, all haemodynamic parameters, such as LVDP, coronary perfusion pressure and heart rate, were the same between each group. The hearts subjected to global ischaemia exhibited considerable decrease in LVDP during reperfusion. Treatment with 20 or 100 µmol·L⁻¹ resveratrol 15 min before global ischaemia showed a higher recovery of LVDP; however, treatment with 10 µmol·L⁻¹ resveratrol did not improve LVDP recovery (Figure 1A). The infarct size was significantly

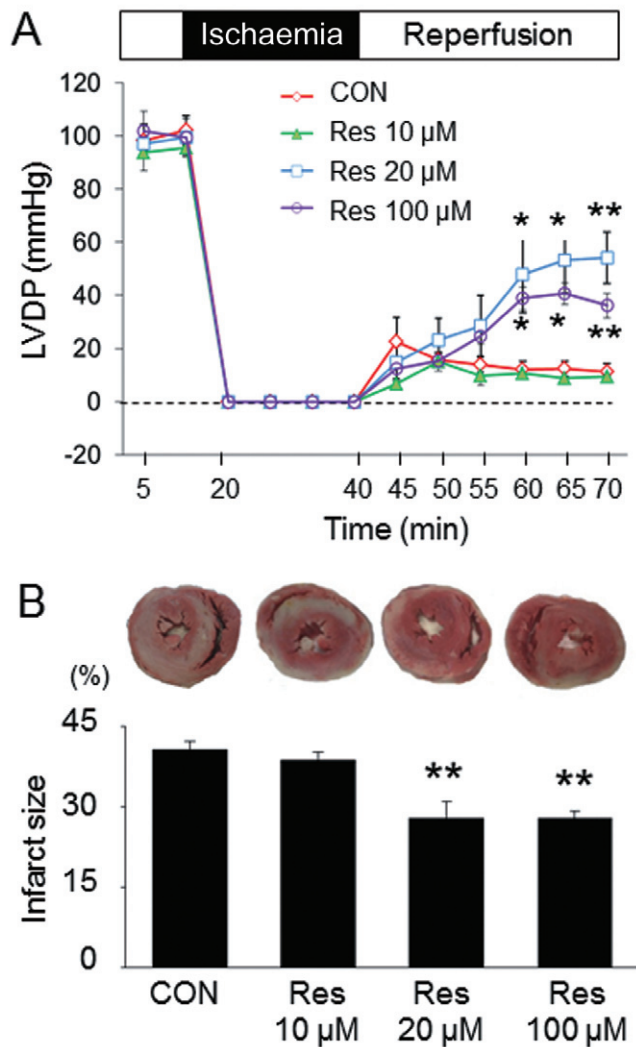


Figure 1

Effects of resveratrol on LV functional recovery and infarct size. (A) Treatment with 20 or 100 $\mu\text{mol}\cdot\text{L}^{-1}$ resveratrol (Res) 15 min before ischaemia improved the recovery of LVDP in rat isolated perfused hearts subjected to 20 min of global ischaemia, followed by 30 min of reperfusion. (B) Treatment with 20 or 100 $\mu\text{mol}\cdot\text{L}^{-1}$ resveratrol reduced the infarct size compared with the control group. Treatment with 10 $\mu\text{mol}\cdot\text{L}^{-1}$ resveratrol neither improved LV functional recovery nor reduced the infarct size in rat hearts. Data are means $\pm$ SE ($n = 5$ for each group). * $P < 0.05$, ** $P < 0.01$ significantly different from control (CON).

decreased in the 20 and 100 $\mu\text{mol}\cdot\text{L}^{-1}$ resveratrol-treated groups compared with that in the control group. Treatment with 10 $\mu\text{mol}\cdot\text{L}^{-1}$ resveratrol did not reduce infarct size as compared with control (Figure 1B).

Cell death

Cell death was induced by incubation with H_2O_2 for 2 h. Figure 2A shows the representative images and quantitative results of PI staining in the cardiomyocytes. Incubation with

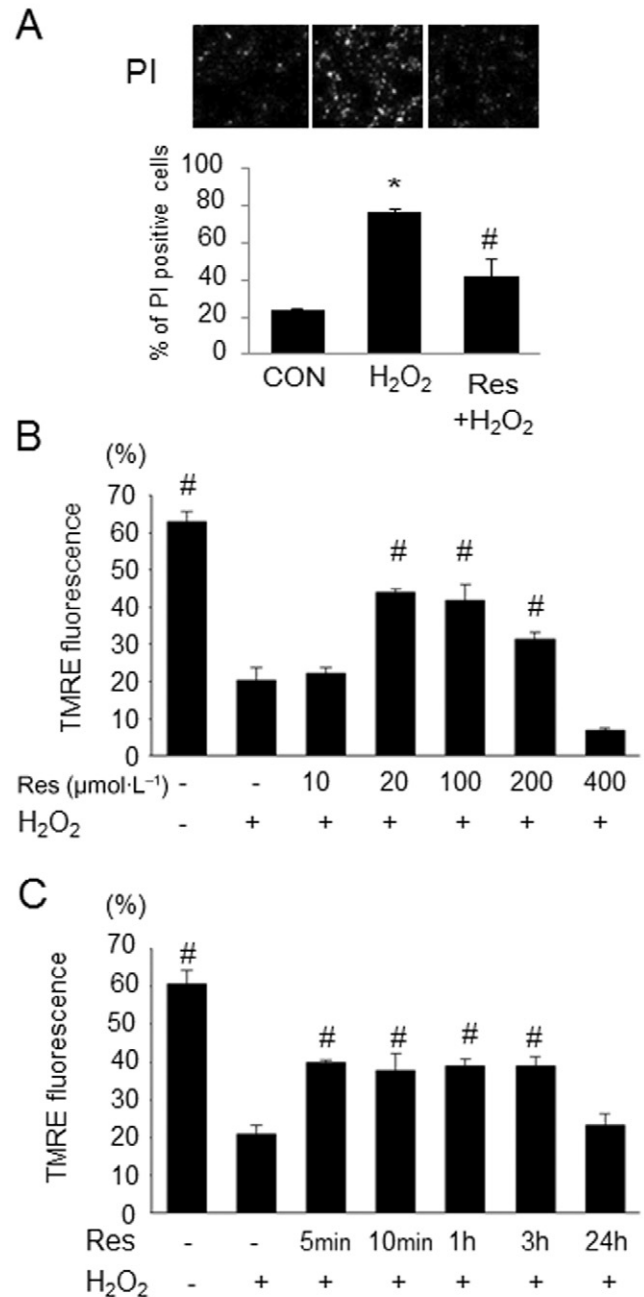


Figure 2

Effects of resveratrol on cell death and the dissipation of $\Delta\Psi_m$ induced by H_2O_2 . (A) Representative images of PI staining of cardiomyocytes stimulated by H_2O_2 (100 $\mu\text{mol}\cdot\text{L}^{-1}$) in the presence or absence of resveratrol (Res; 20 $\mu\text{mol}\cdot\text{L}^{-1}$) and quantitative percentages of PI-positive cells. Resveratrol suppressed the increase of PI positivity induced by H_2O_2 . (B) Summarized data of FACS analysis with various concentrations of resveratrol. Resveratrol preserved $\Delta\Psi_m$ in a dose-dependent manner. (C) Quantitative results of FACS analysis with various durations of pre-incubation with resveratrol. Data are means $\pm$ SE ($n = 7$ for each group). * $P < 0.05$ significantly different from control (CON); # $P < 0.05$ significantly different from H_2O_2 .

H₂O₂ significantly increased the number of PI-positive cells, which was suppressed by pretreatment with resveratrol (20 $\mu\text{mol}\cdot\text{L}^{-1}$).

Mitochondrial inner membrane potential

Dissipation of $\Delta\Psi_m$ is a critical event in the process of cell death (Kroemer *et al.*, 1998). We examined whether the cardioprotective effect of resveratrol is associated with the preservation of $\Delta\Psi_m$ by assessing the change in TMRE fluorescence by FACS analysis. Incubation with H₂O₂ decreased TMRE fluorescence, indicating the depolarization of $\Delta\Psi_m$. The percentage of cells exhibiting higher TMRE fluorescence was evaluated, as described elsewhere (Thuc *et al.*, 2010). As shown in Figure 2B, $\Delta\Psi_m$ is preserved at resveratrol concentrations of 20–200 $\mu\text{mol}\cdot\text{L}^{-1}$, but not at concentrations as low as 10 $\mu\text{mol}\cdot\text{L}^{-1}$ or as high as 400 $\mu\text{mol}\cdot\text{L}^{-1}$. The protective effect of pre-incubation with resveratrol (20 $\mu\text{mol}\cdot\text{L}^{-1}$) occurred as early as 5 min and continued up to 3 h. Cardioprotection was not observed with pre-incubation for 24 h (Figure 2C).

Levels of ROS

Oxidative stress is one of the key determinants of myocardial ischaemia/reperfusion injury (Zweier and Talukder, 2006). We also examined the role of ROS in the cardioprotective effect of resveratrol. Figure 3 shows the representative histograms and quantitative results from FACS analysis of ROS levels in cardiomyocytes. Treatment with H₂O₂ shifted the peak of the histogram to the right, indicating an increase in ROS level (Figure 3A). The summarized data showed a significant increase in DCF fluorescence in H₂O₂-treated cells, which was suppressed by pretreatment with resveratrol (Figure 3B).

[Na⁺]_i and [Ca²⁺]_i

[Ca²⁺]_i overload following the accumulation of [Na⁺]_i leads to the dissipation of $\Delta\Psi_m$. Therefore, we examined the effects of resveratrol on H₂O₂-induced [Na⁺]_i and [Ca²⁺]_i increases in cardiomyocytes. Figure 4A shows the representative images and sequential changes of CoroNa Green fluorescence in six randomly selected cells incubated with or without resveratrol. [Na⁺]_i gradually increased during the first 30 min after H₂O₂ application and thereafter the rate of increase was augmented to reach its peak at approximately 45 min. Incubation with resveratrol diminished the increase in [Na⁺]_i. Figure 4B shows the representative images and quantitative results of Fluo-4 fluorescence in six randomly selected cardiomyocytes. [Ca²⁺]_i was rapidly and considerably increased at approximately 30 min after H₂O₂ application and reached a plateau at approximately 40 min. Incubation with resveratrol almost completely inhibited the increase in [Ca²⁺]_i due to H₂O₂ application. The fluorescence of CoroNa Green and Fluo-4 were steeply decreased 50 min after the incubation with H₂O₂ because of the rupture of the cardiomyocytes. To examine a possible effect of resveratrol on NCX, we investigated the changes in [Ca²⁺]_i by ouabain, a Na⁺/K⁺ ATPase inhibitor. [Ca²⁺]_i continuously increased during the 60 min scanning after ouabain application. Incubation with resveratrol did not affect the increase in [Ca²⁺]_i due to ouabain

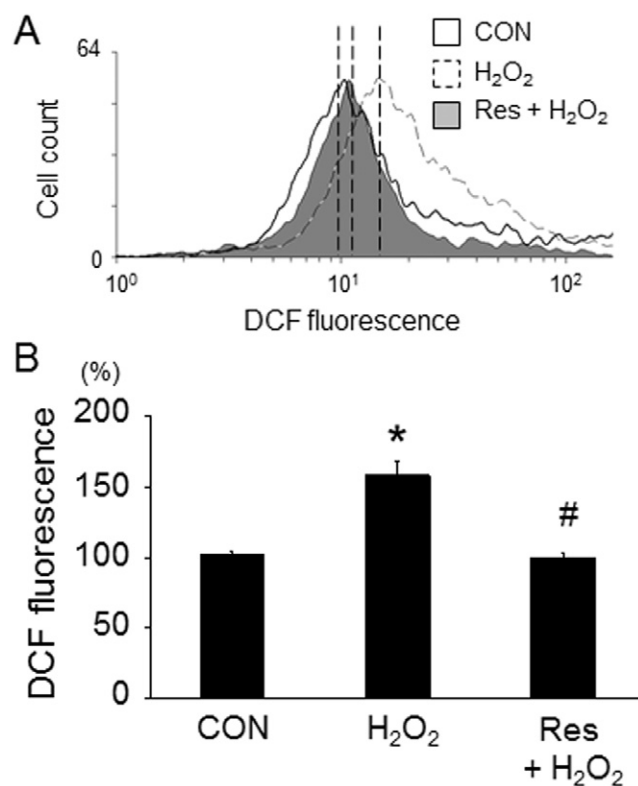


Figure 3

Effects of resveratrol on ROS level. (A) Representative FACS histograms obtained from cardiomyocytes loaded with CM-H₂DCFDA (2 $\mu\text{mol}\cdot\text{L}^{-1}$) in the presence or absence of H₂O₂ (100 $\mu\text{mol}\cdot\text{L}^{-1}$). The area under black line represents control, the grey area represents resveratrol-treated cells and the area under dotted line represents the cells stimulated by H₂O₂. Incubation with H₂O₂ for 20 min shifted the histogram to the right, indicating an increase in ROS level. Treatment with resveratrol (Res; 20 $\mu\text{mol}\cdot\text{L}^{-1}$) for 1 h before H₂O₂ application almost totally suppressed the increase in ROS level induced by H₂O₂. (B) Quantitative results analysed the intensity of fluorescence in the peak of the histograms. Data are means $\pm$ SE ($n = 6$ for each group). * $P < 0.05$ significantly different from control, # $P < 0.05$ significantly different from H₂O₂.

application (Figure 4C). Taken together, these results indicate that resveratrol may affect NHE, but not NCX, to suppress [Ca²⁺]_i overload.

NHE activity

Next, we examined the effect of resveratrol on NHE activity. Figure 5A shows the representative alteration in the pH_i of cardiomyocytes. Perfusion with NH₄Cl solution increased pH_i, which rapidly decreased after the perfusate was switched to a Na⁺-free solution. During 10 min of Na⁺-free perfusion, pH_i was stable at approximately 6.5. The recovery of pH_i began after the cells were superfused with Na⁺-containing Krebs solution. A bold line indicates the slope of the initial 1 min of pH_i recovery in each experiment (Figure 5B). Figure 5C shows the summarized data of the pH_i recovery rates from five independent experiments. Exposure to H₂O₂ for 10 min increased NHE activity, which was attenuated by

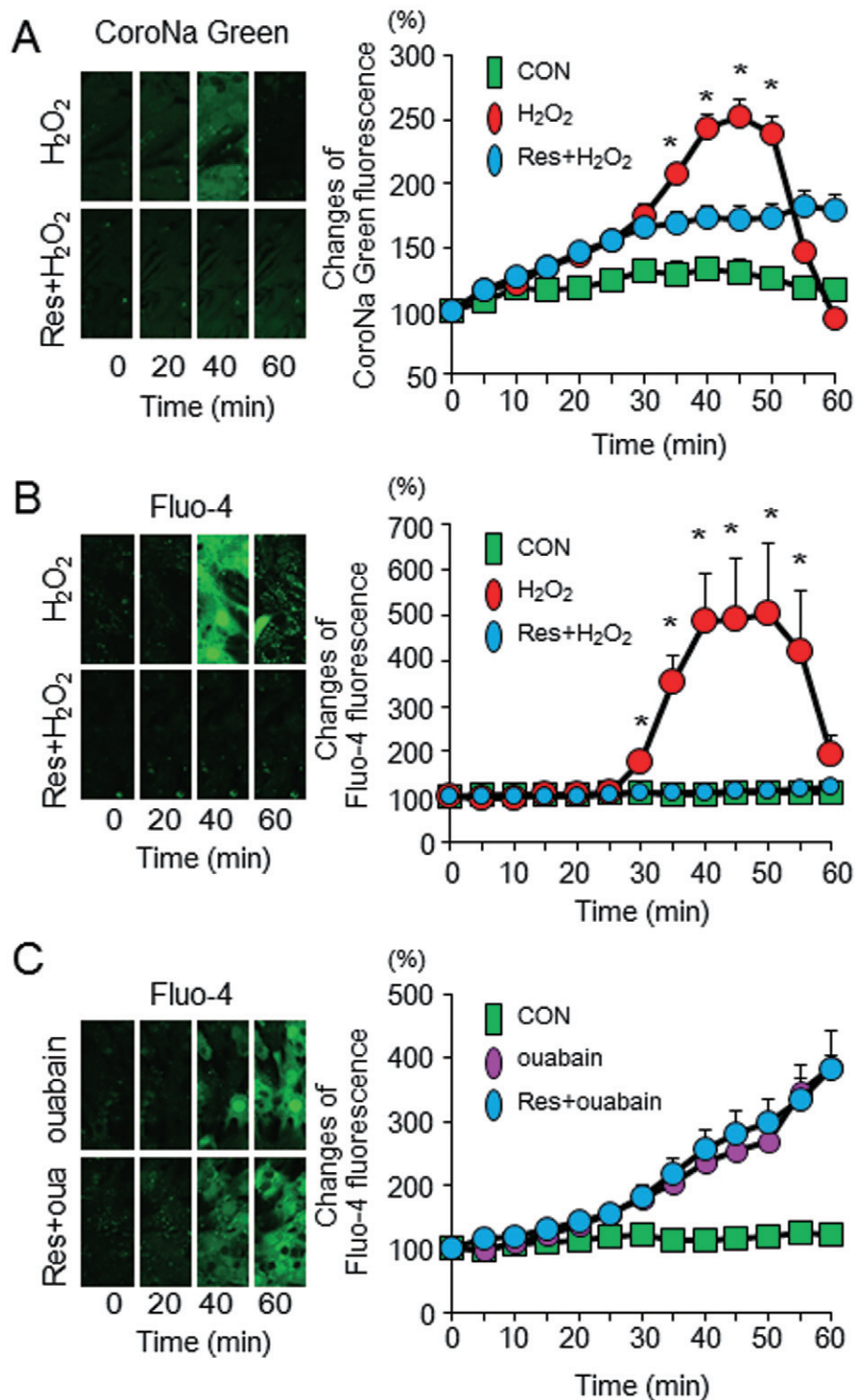


Figure 4

Effects of resveratrol on $[Na^+]_i$ and $[Ca^{2+}]_i$. (A) Representative sequential images of CoroNa Green fluorescence obtained from the cardiomyocytes loaded with H_2O_2 ($100 \mu\text{mol}\cdot\text{L}^{-1}$) in the presence or absence of resveratrol (Res; $20 \mu\text{mol}\cdot\text{L}^{-1}$) and quantitative data from each group. $[Na^+]_i$ started to increase after the H_2O_2 application and peaked at approximately 45 min. This increase was attenuated by pretreatment with resveratrol. (B) Representative sequential images of Fluo-4 fluorescence obtained from the cardiomyocytes loaded with H_2O_2 in the presence or absence of resveratrol and quantitative data from each group. An increase in $[Ca^{2+}]_i$ by the H_2O_2 application was suppressed by pretreatment with resveratrol. (C) Representative sequential images of Fluo-4 fluorescence obtained from the cardiomyocytes loaded with ouabain ($1 \text{ mmol}\cdot\text{L}^{-1}$) in the presence or absence of resveratrol ($20 \mu\text{mol}\cdot\text{L}^{-1}$) and quantitative data from each group. Pretreatment with resveratrol did not affect the increase in $[Ca^{2+}]_i$ by ouabain. Data are means $\pm$ SE ($n = 6$ for each group). * $P < 0.05$ significantly different from Res + H_2O_2 .

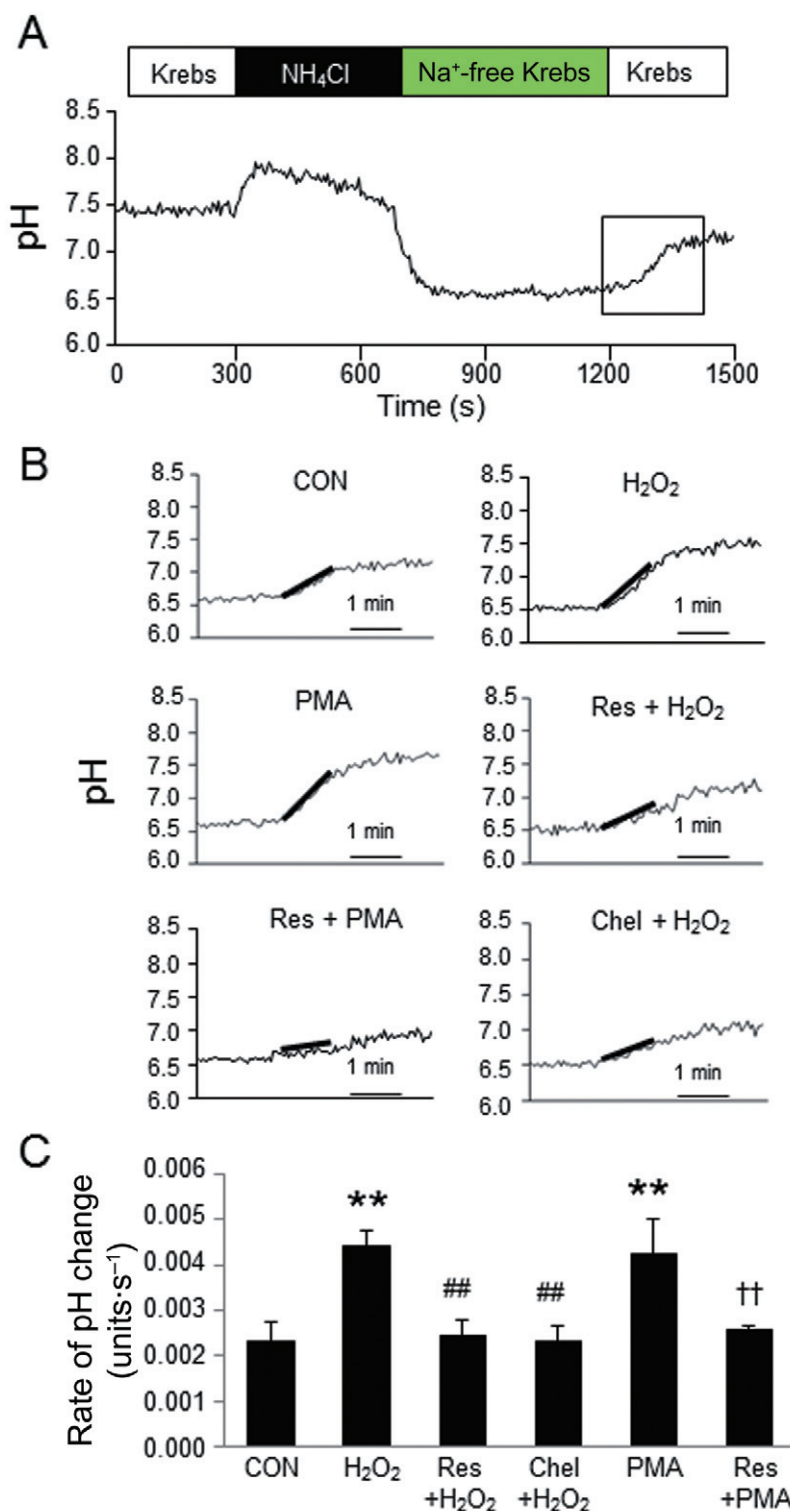


Figure 5

Effect of resveratrol on NHE activity. (A) pH_i was continuously measured using the pH sensitive dye, BCECF. The rate of pH_i recovery during the initial 60 s was defined as NHE activity (outlined by rectangle). (B) Representative pH_i recovery responses from rapid acidification in each group. A line was fitted to pH_i trace during the first minute of recovery. The slopes were measured and compared. (C) Summarized quantitative data from independent experiments in each group. Exposure to H₂O₂ (100 μmol·L⁻¹) increased NHE activity in the cardiomyocytes, which was suppressed by pretreatment with resveratrol (Res; 20 μmol·L⁻¹) or chelerythrine (Chel; 1 μmol·L⁻¹). PMA (5 μmol·L⁻¹) also activated NHE, which was attenuated by pretreatment with resveratrol. Data are mean ± SE (*n* = 5 for each group). ***P* < 0.01 significantly different from control, ##*P* < 0.01 significantly different from H₂O₂, ††*P* < 0.01 significantly different from PMA.

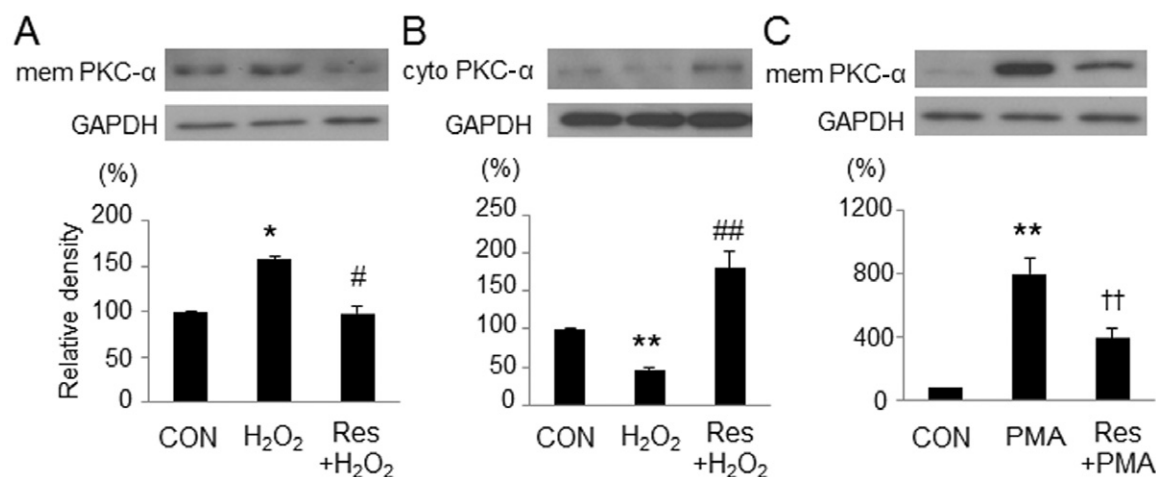


Figure 6

Effects of resveratrol on the translocation of PKC- α . Representative images and quantitative results of immunoblots for PKC- α . All band signals were normalized by GAPDH and compared with control. (A, B) The H₂O₂-induced translocation of PKC- α from the cytosol (cyto) to the sarcolemmal membrane (mem) was reversed by pretreatment with resveratrol (Res). (C) The PMA-induced translocation of PKC- α to the sarcolemmal membrane was attenuated by resveratrol. Data are mean $\pm$ SE ($n = 5$ for each group). * $P < 0.05$, ** $P < 0.01$ significantly different from control; # $P < 0.05$, ## $P < 0.01$ significantly different from H₂O₂; †† $P < 0.01$ significantly different from PMA.

treatment with resveratrol and chelerythrine, a PKC inhibitor. Furthermore, PMA, a PKC activator, increased NHE activity, which was also attenuated by treatment with resveratrol. Taken together, these results showed that resveratrol inhibits NHE activity in a PKC-dependent manner.

Translocation of PKC

PKC isozymes are reported to translocate to different parts of the cell and exert divergent effects in ischaemia/reperfusion injury. During ischaemia, PKC- α is translocated from the cytosol to the cell membrane (Yoshida *et al.*, 1996). Therefore, we examined the effect of resveratrol on the translocation of PKC- α . Incubation with H₂O₂ for 10 min increased the expression of PKC- α in the sarcolemmal membrane and decreased that in the cytosol, which was prevented by pretreatment with resveratrol (Figure 6A,B). The PMA-induced increase in PKC- α expression in the sarcolemmal membrane was also considerably reduced by treatment with resveratrol (Figure 6C). These results indicate that resveratrol inhibited NHE activity by inhibiting the translocation of PKC- α .

Discussion

The main findings of the present study are as follows: (i) resveratrol reduced infarct size and improved LV functional recovery in isolated hearts that were subjected to ischaemia/reperfusion; (ii) resveratrol attenuated cardiomyocyte death by reducing ROS and preserving mitochondrial function; and (iii) resveratrol prevented NHE activation by inhibiting the translocation of PKC- α from the cytosol to the sarcolemmal membrane in cardiomyocytes. This is the first report to demonstrate the PKC-dependent inhibition of NHE as a mechanism of rapid cardioprotection by resveratrol.

On the basis of our results, we have proposed a possible protective pathway for resveratrol in which resveratrol inhibits the translocation of PKC- α from the cytosol to the cell membrane and prevents the subsequent events leading to cell injury (Figure 7). PKC plays a critical role in ischaemia/reperfusion injury. PKC- α is translocated from the cytosol to the cell membrane during ischaemia/reperfusion (Yoshida *et al.*, 1996), and membrane-associated PKC- α has been reported to activate NHE (Maly *et al.*, 2002). In this report, the PKC inhibitor and genetic ablation of PKC- α suppressed the activation of NHE. These results indicate that PKC- α may contribute importantly to the activation of NHE. Furthermore, a study by Slater *et al.*, (2003) showed that resveratrol inhibited the activity of conventional PKC- α in an *in vitro* assay system. In this report, resveratrol inhibited activation of PKC- α by competitive binding to the activator-binding sites of this kinase which prevented PKC- α from binding to plasma membrane and becoming active. These results strongly support our present finding that the inhibitory effect of resveratrol on NHE was PKC- α dependent. Activation of ERK1/2 is another pathway leading to NHE activation by oxidative stress (Snabaitis *et al.*, 2002; Cingolani *et al.*, 2011). Although our data clearly demonstrated a PKC- α -dependent NHE inhibition by resveratrol, the involvement of ERK1/2 is an interesting issue for further study. In a previous study, PKC activated NADPH oxidase (Dekker *et al.*, 2000) which will increase ROS production, suggesting that PKC may work upstream of ROS. However, the present result that activation of NHE by H₂O₂ was inhibited by chelerythrine strongly supports the ROS-PKC-NHE activation sequence as shown in Figure 7. Considering that NHE is activated in response to intracellular acidosis, the link between acidosis and the translocation of PKC- α remains unclear.

An increase in [Ca²⁺]_i precedes irreversible myocardial injury. Much of the increase in [Ca²⁺]_i during ischaemia is due

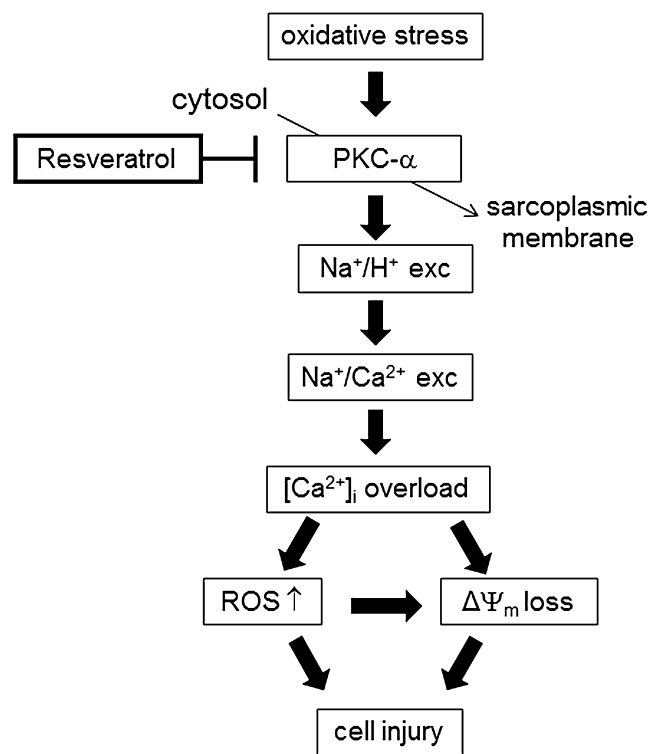


Figure 7

Proposed pathways of cardioprotective effects of resveratrol in ischaemia/reperfusion.

to Ca²⁺ entry because of the reverse mode activity of the NCX. During ischaemia/reperfusion, pH_i becomes acidic, which facilitates the extrusion of H⁺ from the cytosol in exchange for Na⁺ by NHE. Increased [Na⁺]_i can be extruded by NCX in exchange for Ca²⁺, thereby increasing [Ca²⁺]_i. Thus, the inhibition of NCX and NHE should attenuate [Ca²⁺]_i overload and is a possible target for cardioprotection. Previous studies showed that NCX inhibitors such as KB-R7943 and SEA0400 exerted cardioprotective effects against ischaemia/reperfusion injury by attenuating [Ca²⁺]_i overload (Yoshitomi *et al.*, 2005; Motegi *et al.*, 2007). In the present study, resveratrol prevented the increase not only in [Ca²⁺]_i but also in [Na⁺]_i by H₂O₂. Furthermore, resveratrol did not affect the increase in [Ca²⁺]_i by ouabain, which facilitates Ca²⁺ influx through NCX working in reverse mode by inhibiting the Na⁺/K⁺ ATPase pump. These results indicate that resveratrol may affect NHE, but not NCX. We also assayed NHE activity by monitoring pH_i recovery from acid loading and confirmed that resveratrol suppressed H₂O₂-induced NHE activation.

Mitochondria play a key role in the process of cell injury, and loss of the mitochondrial potential, ΔΨ_m, because of the opening of mitochondrial permeability transition pore (mPTP) is a critical step towards cell death. Opening of the mPTP collapses ΔΨ_m and halts oxidative phosphorylation, resulting in ATP depletion and cell death. The association of mitochondrial ATP-sensitive potassium (mitoK_{ATP}) channels with cardioprotection by the inhibition of NHE has been

previously reported (Miura *et al.*, 2001). Aiello and Cingolani (2011) and Mosca and Cingolani (2002), reported that red wine extracts conferred cardioprotection by opening mitoK_{ATP} channels. In contrast, in our preliminary experiments, we did not observe any effect of 5-hydroxydecanoate, a mitoK_{ATP} channel inhibitor, on resveratrol-induced cardioprotection. Considering that PKC is an activator of the mitoK_{ATP} channels (Sato *et al.*, 1998), the cardioprotective effects of resveratrol may be independent of the mitoK_{ATP} channels.

Increase in ROS is a key determinant of ischaemia/reperfusion injury. On reperfusion, re-oxygenation of the cells triggers a burst of ROS; an increase in ROS favours mPTP opening (Crompton *et al.*, 1987). In this study, we showed that resveratrol attenuated a burst of ROS in the cardiomyocytes. Hence, cardioprotection by resveratrol may be mediated by suppressing ROS increase and subsequently preventing mPTP opening. Consistent with the present result, a recent study showed that NHE inhibitors decreased myocardial superoxide production (Garciaarena *et al.*, 2008). However, the mechanisms by which resveratrol suppresses ROS increase remain elusive. Considering that several ROS-generating enzymes are Ca²⁺ dependent (Gordeeva *et al.*, 2003; Brookes, 2004), the prevention of ROS increase may be a secondary effect of the inhibition of [Ca²⁺]_i overload. The preservation of mitochondrial function is also another possible mechanism of ROS reduction. Mitochondria are a major source of ROS production, particularly in a functionally impaired condition, and mitochondrial Ca²⁺ overload can enhance the generation of ROS from mitochondria (Gordeeva *et al.*, 2003; Brookes, 2004). Furthermore, resveratrol may have ROS-scavenging property as suggested in some previous reports (Ray *et al.*, 1999; Danz *et al.*, 2009). In our preliminary experiments, resveratrol exerted cardioprotection against exogenously added H₂O₂ even after the washing out of resveratrol after 1 h pre-incubation. This result excludes the direct effect of resveratrol on H₂O₂. However, it does not exclude a possible scavenging effect of resveratrol after NHE activation.

The plasma concentration of resveratrol in humans after an oral administration of 25 mg resveratrol is reportedly equivalent to 2 μmol·L⁻¹ (Walle *et al.*, 2004). Also, plasma levels after oral administration of 20 mg·kg⁻¹ resveratrol to rats were less than 1 μmol·L⁻¹ (Asensi *et al.*, 2002). These concentrations were lower than those we used in the present study. Higher levels of resveratrol supplementation may be necessary to induce cardioprotection.

In conclusion, we have shown that resveratrol conferred cardioprotection against ischaemia/reperfusion injury and oxidative stress by preserving mitochondrial function and reducing ROS burst. This protection was mediated by the PKC-dependent inhibition of NHE and subsequent attenuation of [Ca²⁺]_i overload.

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Conflicts of interest

None.

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