A novel non-toxic inhibitor of the activation of NADPH oxidase (NOX2) reduces reactive oxygen species production in mouse lung

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List of Abbreviations:
ConA, Concanavalin A; MS, mass spectroscopy; NOX, NADPH oxidase; ROS, reactive oxygen species; PLA2, phospholipase A2; MJ33, 1-hexadecyl-3-trifluorethyglycero-sn-2-phosphomethanol; mPMVEC, mouse pulmonary microvascular endothelial cells; hPASMC, human pulmonary artery smooth muscle cells; PMN, polymorphonuclear leukocytes; STD, internal standard, IT, intratracheal, IV, intravenous; DCF, dichlorofluoroscin; HE, hydroethidine;
Ang II, angiotensin II; HRP, horseradish peroxidase; TBARS, thiobarbituric acid reactive substances
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Abstract

1-Hexadecyl-3-trifluoroethylglycero-sn-2-phosphomethanol (MJ33) is a fluorinated phospholipid analogue that inhibits the phospholipase A$_2$ (PLA$_2$) activity of peroxiredoxin 6 (Prdx6). Prdx6 PLA$_2$ activity is required for activation of NADPH oxidase (NOX2) and subsequent generation of reactive oxygen species (ROS) (Chatterjee et al., 2011). In vitro, MJ33 inhibited agonist-stimulated production of ROS by the isolated perfused mouse lung, lung microvascular endothelial cells, and polymorphonuclear leukocytes. MJ33 (0.02-0.5 µmol MJ33/kg body weight) in mixed unilamellar liposomes was administered to C57BL/6 mice by either intratracheal (IT) or intravenous (IV) routes. Lung MJ33 content, measured by liquid chromatography/ mass spectroscopy, showed uptake of 67-87% of the injected dose for IT and 23-42% for IV administration at 4h post-injection. PLA$_2$ activity of lung homogenates was markedly inhibited (>85%) at 4h post-administration. Both MJ33 content and PLA$_2$ activity gradually returned to near control levels over the subsequent 24-72h. Mice treated with MJ33 at 12.5-25 µmol/kg did not show changes (compared to control) in clinical symptomatology, body weight, hematocrit, and histology of lung, liver and kidney during a 30-50 day observation period. Thus, the toxic dose of MJ33 was > 25 µmol/kg while the PLA$_2$ inhibitory dose was ~0.02 µmol/kg indicating a high margin of safety. MJ33 administered to mice prior to lung isolation markedly reduced ROS production and tissue lipid and protein oxidation during ischemia followed by reperfusion. Thus, MJ33 could be useful as a therapeutic agent to prevent ROS-mediated tissue injury associated with lung inflammation or in harvested lungs prior to transplantation.
Introduction

The NADPH oxidases (NOXs) are a widely distributed family of proteins in mammalian organs that enzymatically generate reactive oxygen species (ROS) (Krause et al., 2012; Takac et al., 2012). NOX2, the original family member to be described, has been the best studied from both biochemical and physiological standpoints (Babior, 2002). ROS generated by NOX2 activity regulate many important cell and physiological processes and also have an important role in host defense (Ushio-Fukai and Alexander, 2004; Leto and Geiszt, 2006; Van Der Vliet, 2008). However, activation of NOX2 can be associated with the deleterious effects of inflammation (Takac et al., 2012).

Inflammation is a complex process that involves cellular recruitment and activation, release of various effector molecules, and additional mechanisms that contribute to the host response to trauma, infection, and other environmental insults. However, an overly exuberant inflammatory response can amplify the injury that occurs with a primary insult. Oxidant stress subsequent to generation of ROS is considered to be one of the mechanisms that may lead to tissue injury during inflammatory states. Molecular targets of oxidant stress include lipids, proteins, and DNA. Among other effects, peroxidation of membrane lipids represents a major threat to cellular integrity, oxidation of proteins can alter metabolic function, and oxidative damage to DNA can impair cell division and repair. Thus, modulation of oxidant stress can have important implications for clinical management of conditions associated with inflammation.

In this study, our focus was the lung which is especially at risk for inflammatory injury associated with sepsis or inhalation of toxic agents. Previous studies have examined the effect of
anti-oxidants in protecting lungs from ROS – mediated injury with mixed results. This latter approach has an obvious drawback since an ROS scavenger has to compete with tissue components for reaction with ROS, unavoidably resulting in some tissue oxidation. Thus, ROS scavenging is unlikely to be fully protective against cellular injury. On the other hand, targeting the pathways that generate ROS has theoretical appeal since it could be more effective in preventing tissue oxidation. Since NOX2 is a major source of ROS with inflammation, there have been some efforts devoted to investigation of inhibitors of this enzyme. A relatively recent review pointed out that most currently available NOX inhibitors have low selectivity, potency and bioavailability, precluding their therapeutic use and work in this area is continuing (Krause et al., 2012).

We recently demonstrated that peroxiredoxin 6 (Prdx6) is a key component of the NOX2 activation pathway in pulmonary endothelial cells and alveolar macrophages (Chatterjee et al., 2011) and a subsequent publication showed a similar requirement for a polymorphonuclear leukocyte (PMN) cell line (Ellison et al., 2012). Peroxiredoxins are a highly conserved family of peroxidases that are widely expressed throughout all kingdoms of life (Rhee et al., 2005). Prdx6 is a novel 1-cysteine member of the family that uniquely exhibits both glutathione peroxidase and phospholipase A2 (PLA2) activities (Manevich and Fisher, 2005; Schremmer et al., 2007; Fisher, 2011). These activities of Prdx6 play important roles in lung antioxidant defense and phospholipid metabolism. Activation of NOX2 is associated solely with the PLA2 activity of Prdx6 as demonstrated by rescue experiments in Prdx6 null cells (Chatterjee et al., 2011; Ellison et al., 2012). Phosphorylation of Prdx6 via a MAP kinase (Wu et al., 2009) followed by
translocation of phosphorylated Prdx6 to the cell membrane are crucial steps in the NOX2 activation pathway (Chatterjee et al., 2011).

The requirement of Prdx6 PLA₂ activity for activation of NOX2 provides a new target for modulation of ROS production. The present study proposes the use of 1-hexadecyl-3-(trifluoroethyl)-sn-glycero-2-phosphomethanol (MJ33), a known inhibitor of the PLA₂ activity of Prdx6 that does not inhibit its peroxidase activity (Fisher and Dodia, 1996; Kim et al., 1997). MJ33 (supplemental figure 1) is a fluorinated analogue of the sn-2 tetrahedral of a phospholipid transition state (Jain et al., 1991b) that shows relatively tight binding to Prdx6 (Manevich and Fisher, 2005) and inhibits the PLA₂ activity of phosphorylated recombinant Prdx6 by greater than 98% (Wu et al., 2009). We have recently demonstrated that use of a system to target MJ33 to the pulmonary endothelium of mice blocks the increases in VCAM expression and alveolar permeability following intratracheal administration of bacterial lipopolysaccharide (Hood et al., 2012).

In the present study, we evaluated the effect of MJ33 on the agonist-modulated activation of NOX2 activity in the intact isolated lung, in lung endothelial cells, and in PMN. Further, we determined the uptake of MJ33 by the intact lung, the in vivo dose-effect relationship for inhibition of PLA₂ activity and the potential toxicity of MJ33. Finally, we evaluated the effect of MJ33 on oxidative injury in a model of lung ischemia/reperfusion. These studies are oriented toward the potential use of MJ33 as a therapeutic agent.

Materials & Methods
Reagents

MJ33 is a fluorinated lipid analogue that was originally synthesized by Dr. Mahendra Jain, one of the co-authors of the present study. The compound is lipid soluble and somewhat soluble in aqueous media with a critical micelle concentration (CMC) of ~10 µM. This agent has been shown to have specificity for PLA$_2$, specifically pancreatic (type 1B) PLA$_2$ and Prdx6 PLA$_2$, but does not inhibit cytosolic (type IV) PLA$_2$, phospholipases C/D, or other lipases (Jain et al., 1991a; Fisher et al., 1992; Gelb et al., 1994; Kim et al., 1997). MJ33 is a chemically non-reactive compound that mimics the tetrahedral transition state of PLA$_2$ substrates and inhibits activity through its binding to the PLA$_2$ protein (Gelb et al., 1994). The inhibitor is effective at mol fraction (MJ33/ lipid substrate) of < 0.01 indicating a high binding affinity (Fisher et al., 1992). MJ33 inhibits NOX2 by preventing the generation of the lipid products of PLA$_2$ activity that are necessary for NOX2 activation (Chatterjee et al., 2011). At this time, a requirement for Prdx6 in the activation of other NOX isoforms has not been demonstrated. For the present studies, MJ33 (as the lithium salt) was purchased from Sigma Aldrich (St. Louis, MO).

Authentic lipids, including 1-heptadecanoyl-2-hydroxy-sn-glycero-3-phosphocholine (17:0 Lyso PC) used as an internal standard for mass spectroscopy, were purchased from Avanti Polar Lipids (Alabaster, MA). $^3$H- dipalmitoylphosphatidylcholine ($^3$H-DPPC) was purchased from (American Radiolabeled Chemicals, St. Louis, MO). Horseradish peroxidase (HRP) and the fluorescent dyes dihydrodichlorofluorescein (H$_2$DCF) diacetate, hydroethidine, and Amplex Red were obtained from Life Technol. (Carlsbad, CA). Angiotensin II (Ang II) was purchased from Bachem Bioscience (Torrance, CA). Ammonium acetate, methanol, chloroform, superoxide dismutase (SOD), and concanavalin A (Con A) were purchased from Sigma-Aldrich.
grade water and acetonitrile were obtained from Fisher Scientific (Fair Lawn, NJ). Gases were supplied by BOC Gases (Lebanon, NJ). A Kinetex C8 column was obtained from Phenomenex (Torrace, CA) and WST-1 was from Dojindo (Kumamoto, Japan).

**Cell isolation and culture**

These studies utilized murine wild type and NOX2 null pulmonary microvascular endothelial cells (mPMVEC), human pulmonary artery smooth muscle cells (hPASMC), murine polymorphonuclear leukocytes (PMN), and A549 cells. mPMVEC were primary isolates from mouse lungs as described previously (Chatterjee et al., 2006) and used at passages 7-12. A549, a lung carcinoma cell line, was obtained from the American Type Culture Collection (ATCC, Manassas, VA). mPMVEC and A549 human lung epithelial cells were grown at 37°C in DMEM-low glucose medium supplemented with 10% fetal bovine serum (FBS-Hyclone), 25mM HEPES, 5mM L-glutamine without antibiotics; for endothelial cells, growth supplement from bovine neural tissue was added at 1.5mg/100ml (E2759; Sigma Aldrich). Immortalized hPASMC, a gift from Dr. Elena Goncharova (Krymskaya et al., 2011), were grown in DMEM-low glucose medium supplemented with 5% fetal bovine serum, 2mM L-glutamine with antibiotics, and 1% ITS (insulin, transferrin, selenium).

Bone marrow PMN were isolated from mouse femurs as described previously (Nick et al., 2000). Briefly, mice were euthanized by intraperitoneal pentobarbital and fibias and femurs were surgically removed and flushed with Hanks balanced salt solution (minus Ca^{2+} and Mg^{2+}) using a 25 gauge needle. Marrow cords were suspended in isolation buffer and marrow cells were obtained by repeated aspiration (Lieber et al., 2004). The suspension was layered on a 3-step
Percoll gradient (72%, 64%, 52%). Following centrifugation at 1000xg for 30 min, cells at the interface of the 72% and 64% Percoll layers were removed and washed with isolation buffer prior to use.

Mice

Ten-week-old C57BL/6 wild-type and NOX2 (gp91phox) null mice were obtained from The Jackson Laboratory (Bar Harbor, ME). Animal care was in compliance with all regulations as set by the University of Pennsylvania Animal Care and Use Committee (IACUC) and by the Guide for the Care and Use of Laboratory Animals published by the US National Institutes of Health. Animals were kept under HEPA-filtered air in the vivarium maintained at 25 ± 3°C.

Preparation and administration of liposomes

Mixed unilamellar liposomes were prepared using authentic lipids with the mole fractional composition 0.5 DPPC, 0.25 egg phosphatidycholine (PC), 0.15 cholesterol, and 0.1 phosphatidylglycerol (PG); this mixture reflects the approximate composition of lung surfactant (Fisher et al., 2005). Liposomes were prepared by evaporating the mixture of lipids under N₂ gas overnight to form a film that was re-suspended with vigorous mixing in PBS with MJ33 at 1-10 mol %.

The mixture was frozen and thawed three times by exposure to liquid N₂ followed by incubation in a 50°C water bath, then extrusion at 50°C for 20 cycles through a 100 nm pore size filter (Avanti Polar Lipids Mini-Extruder). The recovery of lipid after the filtration step, based on [³H] label in DPPC, was 97± 2.3 % (mean ± SE, n=8) of the initial dpm. Liposome size (~100 nm) was confirmed by measurement of dynamic light scattering using a 90 Plus Particle Sizer.
All experiments with in vivo administration of MJ33 utilized the liposomal formulation as the drug carrier.

For IV injection, mice were placed in a restraining apparatus, the two lateral tail veins were dilated by placing the tail in warm water (40°C) for several min, and a 1ml syringe with a 26G3/8 needle was inserted into one of the veins. Liposomes were injected as a 100µl bolus. For IT injection, a midline neck incision was performed to expose the trachea of anesthetized mice. A small puncture hole between the third and the fourth ring in the trachea was made using fine micro-dissecting eye scissors and liposomes were infused directly using a 1ml syringe connected to PE-10-tubing. Liposomes were instilled as a 50µl bolus. A different IT method was used for toxicity studies in order to avoid a surgical incision. With animals in a Plexiglas restraining apparatus, a light-guided laryngoscope (45° angle) was used to visualize the trachea and liposomes were infused through a catheter.

**ROS production by cells**

mPMVEC or hPASMC in culture were treated with control liposomes or liposomes containing MJ33 (1 mol %) for 30 min and then incubated in the dark with an ROS-sensitive indicator dye, H_{2}DCF diacetate or hydroethidine (HE), for an additional 15 min. Cells were washed and treated with agonist to activate NOX2 and then ROS production by cells was measured by fluorescence microscopy. DCF fluorescence was measured at excitation/emission of 490/530 nm and HE fluorescence at 470/610 nm using an epifluorescence microscope fitted with a x20 objective (Nikon Diaphot TMD). Cells were randomly selected from the phase contrast images, and the intensity of cellular fluorescence was measured using Metamorph software (Molecular Devices,
Sunnyvale, CA). For each field, the intensity of 6–10 cells was averaged, and 4–5 fields were analyzed for each treatment condition. Additional experiments were performed by measuring total DCF fluorescence of mPMVEC grown in multiwell plates using a fluorescence microplate reader (FLX 800, Biotek, Winoski, VT). An increase in DCF or HE fluorescence reflects the oxidation of deacetylated H$_2$DCF while increased HE fluorescence reflects the generation of hydroethidium or other oxidation products.

Superoxide (O$_2^-$) production by PMN was measured colorimetrically by the change in absorbance with reduction of the tetrazolium salt, WST-1 (Tan and Berridge, 2000). WST is reduced by O$_2^-$ to a soluble blue formazan dye (Marshall et al., 1995). PMN (5x10$^5$) were equilibrated for 15 min at 37°C without or with MJ33 (1 mol %) in liposomes and then WST (500µM) and Con A (250µg/ml) were added to the incubation medium. Where indicated, superoxide dismutase (SOD, 100 U) was added to the incubation medium. The reduction of WST-1 was measured at 450 nm using a Cary 50 Bio UV-Visible spectrophotometer (Agilent, Foster City, CA). SOD-sensitive O$_2^-$ production was calculated from results in the presence and absence of SOD using a molar extinction coefficient of 37,000 M$^{-1}$ cm$^{-1}$ (Tan and Berridge, 2000).

**Isolated lung perfusion**

Mice were anesthetized with intra-peritoneal (IP) sodium pentobarbital at 50 mg/kg body weight. Liposomes with or without MJ33 (1 mol %) were instilled into the lung through an endotracheal catheter placed at the level of the tracheal carina. After 15 min, the lungs were cleared of blood, removed from the thorax, and placed in a chamber for continuous perfusion as described previously (Fisher et al., 1992; Fisher et al., 2005). Lungs were continuously ventilated through...
a tracheal cannula with air containing 5% CO₂ and perfused with recirculating Krebs-Ringer bicarbonate solution supplemented with 10 mM glucose and 3% bovine serum albumin. ROS generation was measured by the addition of Amplex Red (25 µM) plus HRP (25 µg/ml) to the perfusate (Chatterjee et al., 2011). Amplex Red does not permeate the cell membrane and thus detects extracellular H₂O₂. Ang II (50 µM) was added to the lung perfusate as a NOX2 agonist. Aliquots of the perfusate were removed at 15 min intervals, and fluorescence intensity of the oxidized Amplex Red product was measured (excitation/emission, 545/610 nm) using a spectrofluorimeter (Photon Technology International, Birmingham, NJ). At the end of the perfusion period, the lung was removed from the chamber and dried in an oven to constant weight. Oxidation of Amplex Red was calculated as nmol per gram dry weight of lung using an extinction coefficient of 54,000 cm⁻¹ M⁻¹. The slope of the line for Amplex red oxidation vs. time was calculated by the least mean squares method.

Isolated lungs from wild type and NOX2 null mice also were studied for ischemia-reperfusion injury. In some mice, MJ33 (4 nmol) was administered IV 15 min prior to harvest of the lung. The isolated perfused and ventilated lung was equilibrated for 30 min and then subjected to ischemia by stopping the perfusion pump while ventilation continued; this model results in ischemia without tissue hypoxia (Eckenhoff et al., 1992; Al-Mehdi et al., 1998; Zhang et al., 2005). The perfusion pump was restarted after 60 min of ischemia; reperfusion was continued for another 60 min at which time the lungs were harvested for measurement of lung oxidant injury as assessed by lipid peroxidation (thiobarbituric acid reactive substances [TBARS], 8-isoprostanes) and protein oxidation (protein carbonyls). These procedures for analysis of lung oxidative injury have been reported previously (Liu et al., 2010). ROS production during reperfusion was
measured by addition of Amplex Red (plus HRP) to the perfusion fluid during the initial equilibration period as described above.

Assay for PLA$_2$ activity

To measure the effect of MJ33 on PLA$_2$ activity in vivo, MJ33 in liposomes was injected via intratracheal (IT) or intravenous (IV) routes. Both male and female mice were used for these studies. Mice were sacrificed at intervals by exsanguination under pentobarbital anesthesia and lungs were surgically removed, cleared of blood, and then homogenized for analysis of PLA$_2$ activity. The substrate for measurement of PLA$_2$ activity was mixed unilamellar liposomes containing tracer $[^{3}H]$-DPPC (4,400 dpm/nmol) incubated with lung homogenate at 37°C for 1hr under acidic, Ca$^{2+}$-free conditions (40 mM sodium acetate: 5 mM EDTA buffer pH 4.0) (Chen et al., 2000). The reaction was stopped by addition of chloroform/methanol (1:2) and lipids were extracted by the Bligh and Dyer method (Bligh and Dyer, 1959). Radiolabeled free fatty acids were separated by a two-step thin layer chromatography procedure using hexane/ether/acetic acid as a solvent system along with authentic palmitic acid as a standard. The free fatty acid spots were identified using I$_2$ vapor, scraped from the plate, and analyzed by scintillation counting using an internal standard for quench correction. Values for recovered dpm were corrected for blank values obtained in the absence of enzyme. PLA$_2$ activity was calculated based on the specific radioactivity of DPPC (Chen et al., 2000).

Detection of MJ33 in lung tissue by mass spectroscopy:

MJ33 was measured in lungs from mice that had been injected with varying amounts of MJ33 via an IT or IV route. Animals were sacrificed at varying times post-MJ33 injection. Lungs were cleared of blood by perfusion through the pulmonary artery and homogenized for analysis of
MJ33 levels using a triple quadrupole mass spectrometer (Supplemental figure 1A). In some experiments, lungs were lavaged prior to homogenization by instillation and aspiration (x5) of 800 µl cold PBS. For extraction of lipids, lung homogenate (0.5ml), broncho-alveolar lavage fluid, or equivalent H₂O as a control was transferred to a 10ml glass test tube containing 1ml of PBS buffer (1M, pH 6.8) that was spiked with 2 ng of 17:0 Lyso PC. The latter was chosen as an internal standard (STD) since it is not an endogenous phospholipid, as confirmed by the lack of a corresponding peak in the MS transition. After adding 2 ml of methanol to each tube, the samples were vortexed for 30 s, and 4 ml of chloroform was added to the samples and shaken for 30 min. After centrifugation at 2500 rpm for 5 min, the organic layer was transferred to a new clean glass tube, then evaporated to dryness under nitrogen, reconstituted in 100µL methanol, and maintained at 4°C in the auto-sampler tray before analysis.

Reverse-phase chromatography for LC/MS measurements was performed using a Waters Alliance 2690 HPLC system (Waters Corp., Milford, MA) with gradient elution at 30°C in the linear mode with a Kinetex C8 column (100× 2.6 mm i.d., 2.6 µm) using a flow rate of 0.2 ml/min. Solvent A was water and solvent B was 95% acetonitrile, each with 10 mM ammonium acetate. The linear gradient was as follows: 30% B at 0 min, 30% B at 2 min, 100% B at 8 min, 100% B at 16 min, 30% B at 16 min, and 30% B at 25 min.

Mass spectrometry was conducted on a Thermo Finnigan TSQ Quantum Ultra AM mass spectrometer (Thermo Fisher, San Jose, CA) equipped with an electrospray ionization (ESI) source operated in the positive ion mode. Unit resolution was maintained for both parent and product ions for multiple reaction monitoring (MRM) analyses. Operating conditions were:
spray voltage 3500V, vaporizer temperature 300°C, and heated capillary temperature 280°C.
Nitrogen was used for the sheath gas and auxiliary gas set at 25 and 10 (in arbitrary units),
respectively. Collision induced dissociation (CID) was performed using argon as the collision
gas at 1.5 mTorr in the second (rf-only) quadrupole. An additional DC offset voltage was
applied to the region of the second multipole ion guide (Q0) at 5 V. The following SRM
transitions were monitored; MJ33, $m/z$ 493 → 113 (collision energy 18eV) as a qualifier, $m/z$ 493 → 119 (collision energy 15eV) as a quantifier, 17:0 Lyso PC, $m/z$ 510 → 104 (collision energy 18eV) as a qualifier and $m/z$ 510 → 184 (collision energy 15eV) as a quantifier. The mass spectrum of MJ33 showed the fragments 381, 139, and 113. A linear calibration curve was constructed with authentic MJ33 in the range 0.1 ng/ml to 10 ng/ml ($y=0.004x + 0.00007$, $r^2 = 0.999$). No peak corresponding to MJ33 was detected in the control samples. All data analysis was performed using Xcalibur software, version 2.0 SR2 (Thermo Fisher) from raw mass spectral data.

**Proliferation, cell viability, and clonogenic survival in response to MJ33:**
For proliferation studies, T25 cell culture flasks were each plated with 1x10^5 cells; MJ33 was added to the plates on the next day. The rate of proliferation was determined on a daily basis for up to 10 days. Attached cells were made into a single suspension by incubation with 0.25% trypsin/EDTA for 3-5min, and counted to obtain the total cell number on each plate. Cell viability was assessed by a trypan blue exclusion assay (Lee et al., 2000). Clonogenic survival studies were performed as described previously (Lee and Shogen, 2008). Briefly, 7ml of 15% FBS-DMEM (low glucose at 1g/L) was added to 6-well plates with $10^2$ - $10^4$ cells that were incubated at 37°C for 5 h to allow for cell attachment and then incubated for various durations.
with MJ33 at 0-25 µM. Cultures were fixed with 99.5% isopropyl alcohol, stained with 1% crystal violet and counted. Colonies with more than 50 cells were scored as positive. The surviving fraction (SF) was calculated by: # of colonies counted/ # of cells plated x (PE/100). To evaluate a potential role of cellular contact inhibition on MJ33 cytotoxicity for mPMVEC, feeder cells (heavily x-irradiated at 30 Gy with a $^{137}$Cs source, rendering them unable to generate clones) were added at $10^3$, $10^4$, or $10^5$ cells per well to generate confluent or plateau-phase growth patterns.

**Toxicity studies in mice**

Toxicity of MJ33 after its administration by IV or IT routes was evaluated by daily measurement of animal weight and observation of the level of spontaneous activity during the subsequent 60 days, measurement of hematocrit at 30 days after dosing, and histological examination of lungs, liver, and kidneys at 10 days after MJ33 administration. Death was not used as an end point in accordance with policies of the University of Pennsylvania IACUC. Only female mice were used for these studies because of the difficulties of housing male mice together for prolonged periods.

**Measurement of hematocrit**

To measure the hematocrit, a capillary pipette containing anticoagulant was inserted in the lateral canthus of the eye and 40 µl blood was collected from the retro-orbital sinus. The capillary tubes were centrifuged for 10 min at 12,500 rpm (Autocrit Ultra3, Clay Adams), and the separate bands of hematocrit were read on a hematocrit scale.
Histological evaluation

Histological evaluation of lungs, liver, and kidneys of mice was performed after administration of MJ33 (500 nmol) via IV or IT routes. Anesthetized mice were exsanguinated 10 days post-treatment by cutting the caudal vena cava. A cannula was inserted into the trachea and fixed with a ligature. Lungs and heart were removed en bloc, and lungs were inflated via the cannula by gentle infusion of fixative (10% phosphate buffered formalin, pH 7.0) over 5 min to reach a constant fluid pressure of 25 cm H₂O. The trachea was tied with a ligature, the lungs were placed in a glass vial containing 10% formalin and were kept on ice for 24 h. Livers and left kidneys also were collected and fixed in formalin. All tissue samples were processed by the Pathology Core at the Children’s Hospital of Philadelphia (Abramson Research Center, Philadelphia, PA, USA). After fixation, tissues were embedded in paraffin and blocks were sectioned and stained with hematoxylin and eosin. The tissue sections were examined independently by 2 different investigators and a result was determined by consensus.

Statistical analysis

All measured values are presented as the mean ± standard error (SE) for each group. ANOVA with post-hoc Tukey test (SysStat, San Jose, CA) was used for multiple group comparisons. Differences within a group before and after treatment were evaluated using a paired t-test. Statistical significance was set at 95% (p < 0.05).

Results

Effect of MJ33 on agonist-induced ROS production
The effect of MJ33 on agonist-induced ROS generation was evaluated in isolated lungs of mice with Amplex Red (plus HRP) added to the perfusate to detect H$_2$O$_2$. Unilamellar liposomes containing MJ33 at 0, 0.2 or 4 nmol were injected IV into anesthetized mice and the lungs were isolated 15 min later for \textit{in vitro} perfusion. Amplex red oxidation was plotted vs. time of perfusion (Fig. 1A). The addition of Ang II to the perfusate resulted in approximately 7-fold increase in the rate of Amplex Red oxidation (Fig. 1A). The effect of Ang II was inhibited by approximately 25% by 0.2 nmol MJ33 and was essentially abolished in lungs that had been pretreated with 4 nmol MJ33. This result indicates ROS production in the presence of Ang II and a dose-dependent inhibition by treatment with MJ33. Similar inhibition of Amplex Red oxidation was observed after pretreatment with 4 nmol MJ33 by the IT route (not shown). ROS production following Ang II treatment of lungs from NOX2 null mice was decreased by > 90 % as compared to wild type lungs indicating that NOX2 is the predominant source of ROS in this model (Fig. 1A). However, ROS production by NOX2 null lungs following Ang II was greater (P < 0.05) than control (evaluated at 60 min of perfusion) suggesting another (non-NOX2) albeit minor (~9%) source of Ang II-stimulated ROS production. There was no significant difference in Ang II stimulated ROS production between the wild type treated with 4nmol MJ33 and the NOX2 null lungs, suggesting that this dose of MJ33 totally inhibited NOX2-dependent ROS production but had no effect on the NOX2-independent ROS production (Fig. 1A).

The effect of MJ33 on ROS production was evaluated further with mPMVEC in monolayer culture. ROS production was determined by oxidation of DCF or HE as assessed by fluorescence microscopy. mPMVEC that were stimulated by Ang II showed increased DCF fluorescence (Fig. 1B) or HE fluorescence (not shown) that was abolished by pre-treatment of cells with MJ33.
Similar results were obtained for the effect of MJ33 on ROS production by mPMVEC by measuring cellular fluorescence with a multiplate reader (not shown). Significant ROS production was not seen following Ang II treatment of NOX2 null mPMVEC (Fig. 1B). PMN isolated from mouse bone marrow also showed a robust respiratory burst, as measured by WST reduction in response to Con A (250µg/ml) (Fig. 1C). Reduction of WST reached a plateau value in ~4 min. This stimulated O$_2^-$ generation was inhibited by pre-treatment of PMN with MJ33 (4 µM). The rate of Con A-stimulated O$_2^-$ generation, calculated from the change of WST-1 absorbance during the linearly increasing portion of the curve and corrected for the residual after SOD treatment was 484 ± 9 pmol/min/10$^6$ cells for control cells and 104 ± 74 for cells pre-treated with MJ33 (mean ± SE, n=3), a 79% reduction in the presence of MJ33. There was essentially no SOD-insensitive O$_2^-$ production by these cells (Fig. 1C). PMN from NOX2 null mice showed minimal WST-1 reduction following Con A (41±18 pmol/min/10$^6$ cells), indicating NOX2 as the predominant (>90%) source of O$_2^-$ under these experimental conditions (Figure 1C). MJ33 had no effect on this residual O$_2^-$ production by NOX2 null PMN (not shown).

The results for Amplex red oxidation by the isolated lung stimulated with Ang II indicated that about 10% of ROS production was through non-NOX2 sources. While there are several possible enzymatic sources for non-NOX2 ROS production, NOX1 would be a likely possibility as this is the major source of ROS induced by AngII in vascular smooth muscle (Lassegue et al., 2001). We thus evaluated the possible effect of MJ33 on AngII-stimulated ROS production by hPASMC. These cells showed an increase in ROS production in the presence of Ang II that was less robust than the response of mPMVEC (Supplemental Fig. 2). ROS production by these cells
was unaffected by the presence of MJ33 indicating that NOX1 does not require the PLA$_2$ activity of Prdx6 for activation.

**Uptake and retention of MJ33 in the lung**

As the initial step to evaluate potential use of MJ33 *in vivo*, the uptake and retention of MJ33 by the lung were determined after IT or IV administration of the agent. MJ33 accumulation in the lung homogenate and bronchoalveolar lavage fluid was evaluated by mass spectrometry (MS) under positive electron spray ionization (ESI) conditions. The mass spectrum of authentic MJ33 exhibited an intense parent ion at $m/z$ 493 corresponding to the parent M-H$^+$ (not shown); the MS/MS spectrum indicated the product ions $m/z$ 139 and $m/z$ 113 (Supplemental Figure 1). Since there is no isotopically labeled MJ33 commercially available, we chose 17:0 lysoPC as an internal standard (STD). MS/MS analysis of the lysoPC standard showed the characteristic PC fragment, $m/z$ 184 (not shown). The calibration curve used to quantify MJ33 in lung tissue was constructed using the area ratio of the MJ33 to the area ratio of the STD. To confirm the presence of MJ33 in the lung homogenate, we used liquid chromatography (LC)-MS/MS. The MJ33 detected in the lung homogenate had the same MH$^+$ ion as authentic MJ33 and the same product ions, $m/z$ 139 and $m/z$ 113. The internal STD was used to confirm the relative retention time of the MJ33 detected in lung homogenate vs. the relative retention time of authentic MJ33.

In control lung (liposomes only, no MJ33), a peak corresponding to MJ33 was not seen (Figure 2A, top) but the presence of a peak corresponding to the STD (Figure 2A, bottom) indicated the effective extraction of lipids from the sample. There were significant peaks corresponding to MJ33 and the STD after IV (Figure 2B) or IT (Figure 2C) injection of 0.4 nmol MJ33. At 4 h
after IV injection of 0.4, 4, or 10 nmol MJ33, 23-42% of the administered dose was found in the lung (Table 1). The retention in the lung at 4h-post IT injection of MJ33 was 67-87% of the injected dose (0.4-10 nmol). The percentage of injected MJ33 present in the lung declined slightly at 24 and 48 h after dosing but MJ33 was undetectable in the lung after either IV or IT administration at 72 h-post treatment (Table 1). MJ33 was not detected in the bronchoalveolar lavage fluid at 4h after IT administration of 0.4 nmol (not shown). These results indicate that the MJ33 was retained in the lung for a considerable period and was either intracellular or tightly bound to the cell surface.

**Inhibition of lung PLA2 activity**

The effect of MJ33 on lung PLA2 activity following IT or IV administration was evaluated using dosing protocols similar to those used for MS analysis. The enzymatic assay was performed at pH 4 in the absence of Ca2+, conditions that are relatively specific for the PLA2 activity of lung Prdx6 (Fisher et al., 1992; Fisher and Dodia, 1996). The patterns for inhibition of lung PLA2 activity were similar for MJ33 administered by either the IV (Fig. 3A) or IT (Fig. 3B) route. There was a marked (~85%) decrease in lung PLA2 activity at the 4 h time point that was similar for all 3 doses (0.4, 4, 10 nmol) of MJ33. However, the persistence of the inhibitory effect of MJ33 beyond 4h and the degree of recovery were dose dependent. For doses of 4 and 10 nmol, there was only slight recovery at 24 h and partial recovery at 48 h post-treatment but activity had returned close to the control values at 72 h post-treatment. This pattern of effect on PLA2 activity agreed reasonably well with the pharmacokinetics of MJ33 in lung tissue as evaluated by MS/MS (Table 1). There were no differences between male and female mice in the time-related effect of MJ33 on lung PLA2 activity.
In vitro cytotoxicity of MJ33

Cytotoxicity of MJ33 was evaluated with A549 human lung epithelial cells and mPMVEC during the exponential growth-phase and with confluence. Plating efficiency (PE), calculated as the percentage of seeded cells that grow into macroscopic colonies in the absence of MJ33, was ~45% for mPMVEC and ~55% for A549 cells. There was no effect of 10-25 µM MJ33 on A549 cell division during a 10 day observation period (Fig 4A). Survival of A549 cells exposed continuously during the exponential growth phase to 5-25 µM MJ33 was unaffected as determined by colony formation (clonogenic assay) at 2 or 7 days of exposure (Fig. 4B). We have shown previously that acute exposure (<24 h) to MJ33 at 10-25 µM did not reduce the viability of mPMVEC, as measured by a MTT assay (Lien et al., 2012), and the present study showed no decrease in survival for confluent cells exposed for 24 h to 10 or 20 µM MJ33 (Fig. 4C). A clonogenic survival assay with exposures of mPMVEC for up to 7 days also was used to assess cytotoxicity. Continuous exposure to 5-10 µM MJ33 for 24 h had no effect on survival of exponentially growing mPMVEC, although survival was decreased with longer exposure to MJ33 at concentrations > 5 µM (Fig. 4C). The presence of a feeder cell layer resulted in a statistically significant (P < 0.05) but relatively modest (~10%) increase of mPMVEC survival on exposure to 20 µM MJ33 for 24h (not shown). For recombinant Prdx6 assayed in vitro, PLA₂ activity was inhibited by 50% at 0.3 µM and essentially 100% at 0.4 µM MJ33 (not shown). Thus, these studies with primary lung endothelial cells and a lung cancer cell line show a lack of toxicity for confluent cells with MJ33 up to 25 µM while MJ33 at concentrations >5 µM appeared to be cytotoxic to the primary lung endothelial cells, but not the lung cancer cell line,
during the exponential growth phase. The toxic concentration of MJ33 significantly exceeds the concentration required for inhibition of PLA₂ activity.

_In vivo toxicity_

Toxicity studies of intact mice evaluated the effect of MJ33 on their activity level, body weight, hematocrit, and histology of lungs, liver, and kidneys. Mice (n=5 per group) received a single dose of 250, 500 or 2500 nmol MJ33 (~12.5-125 µmol/kg body weight) by the IV or IT route followed by a 7 week observation period (5 weeks for the higher dose). By clinical observation, mice in all 3 treatment groups exhibited generally normal behavior; some mice appeared to be slightly less active during the initial 1-3 days after MJ33 administration but then recovered during the subsequent days. Clinical manifestations such as hyperactivity, aggressive behavior, alopecia, dehydration, salivation, nasal discharge, constipation, or diarrhea were not observed and none of the mice died or had to be euthanized because of poor clinical appearance. A small weight loss was seen for control mice at the end of 1 week of observation that was possibly related to the experimental procedures; these mice during subsequent weeks progressively gained weight to surpass the initial body weight by week 4 (Figure 5A, B). The initial weight loss was slightly greater in most groups of mice dosed with MJ33, but the mice subsequently resumed their normal rate of weight gain. By 5 weeks post injection, the mean weights for all groups except 2500 nmol IT were similar and exceeded their initial weight (Fig. 5 A, B). None of the changes in weight for the single dose groups were statistically different from control (P > 0.05).

MJ33 was also given in repeated doses of 2.5 µmol IV daily for 4 days. The mean body weight in these mice decreased by about 12% compared to the initial weight in the first week of
treatment and did not show a subsequent recovery as seen with the single injection condition (Figure 5A). The difference in body weight beyond 3 weeks compared to control was statistically significant (p < 0.05). Like the single dose studies, clinical symptomology with the multiple dose regimen was not observed.

The hematocrit (Hct), the volume percentage of erythrocytes in blood, was measured prior to treatment and at 30 days after 0, 250, or 500 nmol MJ33 IT or IV. There was no significant effect of MJ33 on hematocrit at these doses (Table 2).

Tissue sections of lungs, liver, and kidneys were examined by 2 observers with good agreement between their respective analyses. The lungs of MJ33-treated and control animals appeared similar with no evidence of alveolar edema, vascular congestion, inflammation, or destruction or fibrosis of alveolar septae (Figure 6). Likewise, no histological differences were found in the liver or kidneys in the MJ33 treated groups as compared to control (Supplemental Fig. 3).

Effect of MJ33 on lung ischemia/reperfusion (I/R) injury

ROS production measured by Amplex red oxidation was increased significantly during the reperfusion phase of I/R. (ROS production during the ischemic period cannot be measured by this method.) The rate of ROS production with reperfusion (Fig. 7) was approximately 50% greater than the rate observed in the presence of Ang II (Fig. 1). The rate of ROS production with reperfusion was decreased by 66% vs. control (continuous perfusion) in the presence of MJ33. ROS production was decreased further (~80% vs. control) with NOX2 null lungs indicating that the major fraction but not all of NOX2-dependent ROS production was inhibited
by MJ33. Approximately 20% of ROS production appeared to be NOX2 – independent in this model of lung injury.

Oxidative stress associated with I/R was evaluated by measuring indices of lung lipid peroxidation (TBARS, 8-isoprostanes) and protein oxidation (protein carbonyls). Altered lung permeability was evaluated by the wet to dry weight ratio. Control was the isolated wild type lung that was continuously perfused for 2 h (Table 3). Values for continuously perfused NOX2 null lungs (not shown) were essentially identical to the control wild type. In wild type mice, the I/R protocol resulted in a significant (p<0.05) increase of 81-110% in the biochemical indices of oxidative stress. The wet/dry ratio of the lung also was significantly increased following I/R. These changes were largely abolished by pretreatment of the mice with MJ33. I/R in the NOX2 null lungs indicated that the bulk of the oxidative damage was due to activation of the NOX2 system. However, the values obtained for I/R with NOX2 null lungs were significantly greater than control indicating that ~20% of the lung injury with I/R could be attributed to non-NOX2 sources of ROS (Table 3).

Discussion

Previous studies with mPMVEC, lung alveolar macrophages and PMN have shown that the PLA$_2$ activity of Prdx6 is required for activation of NOX2 and the subsequent production of ROS (Chatterjee et al., 2011; Ellison et al., 2012). We have proposed that inhibiting PLA$_2$ activity could be useful therapeutically as a means to limit the activation of NOX2 and ROS production that are in part responsible for cell injury associated with lung inflammation (Kratzer et al., 2012; Min et al., 2012). In this study, we have evaluated MJ33, a known inhibitor of the
PLA₂ activity of Prdx6 (Fisher and Dodia, 1996; Kim et al., 1997), as an inhibitor of NOX2 activation. As shown in the present study, MJ33 does prevent the activation of NOX2 and generation of ROS. An important advantage of MJ33 is that this agent does not inhibit the peroxidase activity of Prdx6 (Fisher and Dodia, 1996) as this activity has been shown to be important for anti-oxidant defense (Wang et al., 2004; Wang et al., 2006a; Wang et al., 2006b; Wang et al., 2008; Hood et al., 2012; Lien et al., 2012). This dichotomy is consistent with the clearly distinct active sites for the two enzymatic activities (Chen et al., 2000; Fisher, 2011). Thus, the important role of Prdx6 as a peroxide scavenger and its function in anti-oxidant defense is preserved in the presence of MJ33.

We first tested the effect of MJ33 on ROS generation by model systems including the isolated perfused mouse lung and mPMVEC in culture that were stimulated by Ang II, and isolated bone marrow PMN that were stimulated with Con A. Agonist-treated PMN from NOX2 null mice showed scant ROS production in the absence of MJ33, indicating NOX2 as the predominant source of ROS; we have previously shown NOX2 as the predominant source of ROS in Ang II-treated mPMVEC (Chatterjee et al., 2011). Study of NOX2 null lungs indicated that approximately 90% of ROS production by Ang II–stimulated lungs was via NOX2. The addition of MJ33 to the lung perfusate or the cell incubation medium markedly inhibited the agonist-induced ROS production in each of these systems. MJ33 inhibits the activation of NOX2 by preventing the generation of lipid products of PLA₂ activity (Chatterjee et al., 2011). Studies with pulmonary artery smooth muscle cells where NOX1 is the predominant source of ROS following Ang II treatment (Lassegue et al., 2001) indicate that MJ33 does not inhibit this NOX
isoform. These results suggest that MJ33 could be useful as a biologically-active agent to inhibit NOX2-mediated ROS generation in vivo.

To better understand possible therapeutic applications of MJ33 in an intact animal, we measured the inhibitory effect of MJ33 on lung PLA₂ activity with increasing concentration of the agent delivered by either IT or IV routes. PLA₂ activity in the lung homogenate was significantly reduced to ~15% of the control level by MJ33 delivered by either route when measured at 4 h after exposure, and remained significantly reduced at 4-24 h post treatment. This 85% reduction in PLA₂ activity (measured at pH 4 in Ca²⁺ free buffer) with MJ33 is similar to the effect of Prdx6 “knock-out”, indicating that the residual activity reflects some PLA₂ source other than Prdx6 (Fisher et al., 2005). Varying doses of Prdx6 from 0.4 nmol to 10 nmol (IT and IV) were equally effective in inhibiting enzyme activity, but the recovery time of PLA₂ activity was dose dependent. Based on these observations, a daily treatment with 4 nmol MJ33 (0.2 µmol/kg body wt) in the mouse could effectively maintain near total inhibition of lung Prdx6 PLA₂ activity.

We overcame the inability to label MJ33 with fluorophores or other tags by developing a method using liquid chromatography coupled with tandem mass spectroscopy to assay the pharmacodynamics of MJ33. The method used three parameters to positively identify MJ33 in lung homogenate: first, demonstrating the same parent ion as authentic MJ33 (m/z 493); second, demonstrating another identical product ion as authentic MJ33 (m/z 493); and third, demonstrating the same relative retention time as the internal standard and authentic MJ33 (7.3min). The LC-MS method was linear in the range 0.2-10 ng/mL and the limit of detection was 0.1 ng/mL.
After an IT injection, the percentage of injected MJ33 remaining in the lung at 4 h was 67-87% for doses of 0.4 -10 nmol. Uptake by the lung at 4h after IV injection of MJ33 was 23-42 % of the injected dose; this presumably reflects a first pass effect of the injected lipophilic agent. MJ33 after either IT or IV injection gradually disappeared from the lung during the subsequent 72h. The fate of the MJ33 after leaving the lung is not known but the agent could be stored in the fat or excreted since it is unlikely to be metabolized.

The measurement of lung MJ33 showed generally good agreement with our data for MJ33-induced inhibition of PLA2 activity. Thus, either the IT or IV routes of administration appear to be satisfactory to inhibit lung Prdx6 PLA2 activity. The optimal route for clinical use could depend on the etiology of the inflammatory lung disease. This study has not determined the effect of route of administration on the distribution of MJ33 amongst the various cells that comprise the lung. Conceivably the distribution could favor endothelium after IV administration and epithelium after IT dosing but that remains to be determined.

After showing effectiveness as an enzyme inhibitor in vivo, the next step was to evaluate potential toxicity. The only biological effects thus far described for MJ33 is its ability to inhibit several forms of PLA2, mainly pancreatic (secreted) PLA2 and Prdx6 PLA2 (Jain et al., 1991b; Kim et al., 1997; Chen et al., 2000). Inhibition of the pancreatic enzyme is unlikely to be toxic over the short term and mice null for this enzyme seem to do quite well (Huggins et al., 2002). Mice that are null for Prdx6 reproduce normally and have normal growth rates and life spans (Mo et al., 2003; Wang et al., 2003) although they show defective lung lipid metabolism with
age-dependent accumulation of surfactant-related lipids (Fisher et al., 2005). Prdx6 null mice also showed increased oxidant sensitivity related to the loss of the Prdx6 peroxidase activity (Wang et al., 2003; Wang et al., 2004; Wang et al., 2006a), but this function of Prdx6 is unaffected by MJ33 (Fisher et al., 1999). Thus, a toxic effect of MJ33, if found, would likely be due to the chemical itself and unrelated to its inhibition of PLA₂ activity.

Assay of cellular proliferation and clonogenic survival of a lung cancer epithelial cell line and primary lung endothelial cells were used to evaluate potential cytotoxicity of MJ33. Toxicity of MJ33 at concentrations up to 20-25 µM was not demonstrated for A549 cells in the rapid growth phase, for confluent A549 cells, or for confluent mPMVEC. mPMVEC that were treated with MJ33 for increasing time of exposure (up to 7d) showed inhibition of clonogenic potential at concentrations > 5 µM. Based on a dose of 4 nmol IT to inhibit lung PLA₂ activity for 24 h, one can estimate an average peak MJ33 concentration in the lung of < 20 µM, a dose that is non-toxic for confluent cells.

To assess the potential toxicity of MJ33 in intact mice, MJ33 was administered far above the 0.4-4 nmol dosages that fully inhibit lung Prdx6 PLA₂ activity. Mice treated with a single dose of MJ33 at up to 2500 nmol showed neither overt clinical signs of toxicity nor weight changes that were significantly different than control. There was no change in hematocrit or lung histology following treatment. Based on these results, MJ33 appears to be a reliable non-toxic inhibitor of NOX2 activation. Considering that 0.4 nmol to the mouse fully inhibited PLA₂ activity and 2500 nmol in a single dose was not toxic, MJ33 appears to have a safety margin of at least 6000, although multiple doses of MJ33 (10 µmol total dose) did result in non-lethal toxicity.
The potential utility of MJ33 as an agent to prevent oxidative stress was evaluated in a model of lung I/R injury. The lung injury in I/R is recognized in part as a manifestation of oxidative stress associated with NOX2 activation (Eckenhoff et al., 1992; Dodd-O and Pearse, 2000; Zhang et al., 2005; Brandes et al., 2010). The role of oxidative stress with I/R in the current study was demonstrated by increased ROS production, increased tissue lipid peroxidation, increased protein oxidation, and increased lung permeability. These manifestations were markedly reduced by pretreatment of mice with MJ33. These results suggest potential clinical utility for MJ33 in conditions where pretreatment in anticipation of I/R is a possibility. A clear example would be pretreatment of donor lungs in anticipation of lung transportation.

Our study based on NOX2 null mice indicates that ~80% of the ROS production and oxidative injury with lung I/R is due to NOX2 activation and that there was no significant difference between NOX2 null mice and wild type mice pre-treated with MJ33. The ROS production that was not from NOX2 could derive from NOX1 activation, the mitochondrial respiratory chain or xanthine oxidase among other sources and does not appear to be inhibited by MJ33. NOX3 and NOX5 are not present in rodent lung and NOX4 is constitutively active so they should not contribute to the increase in ROS production with 2 h I/R. Of note, Prdx6 can also function as an ROS scavenger through its peroxidase activity, and this activity is not inhibited by MJ33.

Based on its low toxicity, MJ33 would appear to offer a major advantage over currently available NOX2 inhibitors (Jaquet et al., 2009; Krause et al., 2012). Among these agents, diphenyleneiodonium (DPI), a commonly used inhibitor of NOX2 in vitro, is not specific and its
toxicity related to general inhibition of flavoproteins would preclude its use as a therapeutic agent (Gatley and Martin, 1979; Jaquet et al., 2009). Apocynin is perhaps the most specific inhibitor described to date. Importantly this agent can ameliorate lung injury associated with ischemia-reperfusion, hemorrhagic shock, and gram negative sepsis (Wang, 1994; Dodd-O and Pearse, 2000; Zhou et al., 2002; Abdelrahman et al., 2005; Zhu et al., 2008; Yang et al., 2009). However, the mechanism for action of this inhibitor is not clear and the margin between its therapeutic and toxic doses after systemic administration is relatively narrow (Tang et al., 2008). Several other drawbacks to the use of apocynin have been noted including observations that it initially stimulates ROS production and that its activation requires ROS plus a peroxidase (Brandes et al., 2010). Recently, a TAT-peptide inhibitor of PKCδ has been shown to block NOX2 activation and protect against sepsis-induced lung injury (Kilpatrick et al., 2010), but studies of toxicity and specificity for the agent have not yet been reported and clinical use of the TAT-peptide may be problematic. Although these other inhibitors of NOX2 have limited clinical utility, their effectiveness supports the concept that inhibition of this enzymatic pathway can have a beneficial effect on lung inflammation.

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Authorship Contributions

Participated in research design: Lee, Chatterjee, Zagorski, Feinstein, Fisher
Conducted experiments: Lee, Dodia, Chatterjee, Zagorski

Contributed new reagent or analytic tools: Mesaros, Blair, Jain

Performed data analysis: Lee, Dodia, Chatterjee, Mesaros

Contributed to the writing of the manuscript: Lee, Mesaros, Fisher

Declaration of Conflicting Interest

The authors declare no conflicts of interest with respect to the authorship and/or publication of this article.
References


adenine dinucleotide phosphate (NADPH) oxidase inhibitor, in isolated perfused rat lung.

Footnotes

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Figure Legends

Figure 1. Inhibition of agonist-induced ROS production by MJ33. A. Effect of MJ33 on Ang II-induced ROS generation by isolated perfused wild type (WT) and NOX2 null mouse lungs as measured by Amplex Red oxidation. Liposomes containing 0, 0.2, or 4 nmol MJ33 were administered intravenously over 1 min to the intact mouse. The lung was isolated 15 min later, placed in the temperature controlled (37°C) chamber, and perfused in the absence or presence of Ang II (50 µM). Perfusate samples were taken at 15 min intervals and analyzed for oxidized Amplex Red; results are plotted vs. time of perfusion. The numbers in parentheses indicate the slope (in nmol/g dry wt/min) of the lines calculated by the least mean squares method. Results are mean ± SE for n=3 (in some cases, the SE bars are within the plotted points). The slope of the line for NOX2 null mice minus Ang II (not shown) was essentially identical to the control. B. ROS production by isolated mouse pulmonary microvascular endothelial cells as indicated by epifluorescence microscopy for DCF. Cells were incubated for 30 min with liposomes with or without MJ33 (1 mol %), then loaded with H₂DCF diacetate (5µM) and stimulated with 10 µM Ang II. Integrated fluorescence of 10-15 cells per field was quantitated using Metamorph software. Basal conditions represent no Ang II. DCF fluorescence intensities are expressed in arbitrary units. Values are mean ± S.E. for n=3. *P<0.05 vs the corresponding basal. C. Effect of MJ33 in liposomes (1 mol %) on the respiratory burst of mouse bone marrow PMN stimulated with concanavalin A (Con A, 250 µg/ml). ROS production was evaluated at 30 s intervals by the change in absorbance due to reduction of WST-1. The control was Con A stimulation in the presence of SOD. This figure is representative for n=3.
Figure 2. Measurement of MJ33 content in the lung using liquid chromatography (LC)/mass spectroscopy (MS). MJ33 (0.4 nmol in liposomes) was injected intravenously (IV, Panel B) or into the trachea (IT, panel C) in the intact mouse. The lungs were isolated 4 h later, cleared of blood and homogenized. An internal standard (STD), 17:0 lysophosphatidylcholine, was added to the lung homogenate and the lipids were extracted. The chromatograms are shown in the upper panels for MJ33 and in the lower panels for the STD. The elution time for all MJ33 and STD peaks was 7.3 min. MJ33 was detected from m/z 493 → 139 and the STD from m/z 510 → 184. Liposomes without MJ33 were injected IT as a baseline control (Panel A). The integrated peak area for each panel is indicated.

Figure 3. PLA₂ activity of lung homogenate from male and female mice measured at varying times after A. intravenous (IV) or B. intratracheal (IT) injection of MJ33 (0-10 nmol) in liposomes. PLA₂ activity was measured by assay in Ca²⁺-free medium, pH 7. Results are mean ± SE for n=3-5. *P < 0.05 vs. control (no MJ33) for each time point.

Figure 4. Effect of MJ33 on the proliferation of cells in culture. A. Proliferation of A549 cells during exposure to 0, 10, or 25 µM MJ33 during a 10 day incubation. B. Clonogenic assay of A549 cells after a 2 or 7 day exposure to MJ33 at 0-25 µM. At the end of exposure, the number of colonies were counted and expressed as a fraction of the number of cells that were originally plated. C. Clonogenic assay of mouse pulmonary microvascular endothelial cells treated with 0 to 25 µM MJ33 for 1, 2, or 7 days. Cells were studied during the exponential growth phase or at confluence (confl-24h). Mean ± S.E. for n=3-4.
Figure 5. Change in body weight for mice that were administered 250, 500, or 2500 nmol MJ33 in liposomes A. intravenously (IV) or B. intratracheal (IT). An additional IV group received multiple doses (IV) at 2500 nmol per day for 4 days. Mouse body weights were measured at weekly intervals. Values are mean ± SE for n=5 for each group. Initial body weight of mice was 21.5 ±0.15 g (n=45). No deaths occurred during the observation periods. * P<0.05 compared to the corresponding control (no MJ33).

Figure 6. Effect of MJ33 on lung histology. The mice were administered MJ33, 500 nmol in liposomes or blank liposomes (control) by either IT or IV routes. The sections were stained with hematoxylin and eosin. A. scale bar 200 µm; B. scale bar 50 µm.

Figure 7. ROS production during reperfusion in mouse lungs following 60 min ischemia. ROS production was measured by the Amplex Red method as described in Fig. 1. The dye was added to the perfusate during the 15 min equilibration period prior to ischemia. A zero reperfusion time perfusate sample was used for background subtraction. Values represent mean ± SE for n=3. The numbers in parentheses indicate the slope of the line (nmol/g dry wt/min) calculated by least mean squares.
Table 1. Lung uptake of MJ33 at increasing time after IV or IT injection as determined by mass spectroscopy

<table>
<thead>
<tr>
<th>Time Post-injection (h)</th>
<th>Amount injected nmol</th>
<th>Uptake %, of injected dose IT</th>
<th>Uptake %, of injected dose IV</th>
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<td></td>
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<td>4</td>
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<tr>
<td></td>
<td></td>
<td>0.4</td>
<td>68±4 *</td>
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<td></td>
<td></td>
<td>4</td>
<td>87</td>
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<td>10</td>
<td>67</td>
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<td>0.4</td>
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<td>10</td>
<td>nd</td>
<td>nd</td>
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</table>

The amount of MJ33 was calculated based on the area ratios (Fig. 2) using a calibration curve constructed with the same amount of internal standard and variable amount of MJ33. nd, none detected, *Mean±range for n=2; all other values are single measurements.
Table 2. Hematocrit levels at 30 d after IV or IT injection of MJ33

<table>
<thead>
<tr>
<th>MJ33 dose (nmol)</th>
<th>Hematocrit (mean±SE)</th>
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<tbody>
<tr>
<td></td>
<td>IT</td>
<td>IV</td>
<td></td>
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<tr>
<td>control*</td>
<td>47.7±0.3</td>
<td>45.8±0.9</td>
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</tr>
<tr>
<td>250</td>
<td>47.2±0.4</td>
<td>46±0.3</td>
<td></td>
</tr>
<tr>
<td>500</td>
<td>46.4±0.7</td>
<td>47.0±0.4</td>
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</table>

Pretreatment hematocrit was 47.4±0.2. Values are mean±SE for n=5.

*Blank liposomes with MJ33.
Table 3. Effect of MJ33 on indices of oxidative stress associated with lung ischemia/reperfusion (I/R)

<table>
<thead>
<tr>
<th></th>
<th>TBARS</th>
<th>8-isoprostanes</th>
<th>Protein carbonyls</th>
<th>Lung weight wet/dry</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>74.6±2.5*†</td>
<td>210 ±7*†</td>
<td>5.07±0.19*†</td>
<td>5.85±0.04*†</td>
</tr>
<tr>
<td>I/R, wild type</td>
<td>136±3.0†</td>
<td>440± 23†</td>
<td>9.25±0.12†</td>
<td>7.35±0.07†</td>
</tr>
<tr>
<td>I/R + MJ33, wild type</td>
<td>95.5±2.1*</td>
<td>310±5**†</td>
<td>7.00±0.01**†</td>
<td>6.35±0.02*</td>
</tr>
<tr>
<td>I/R, NOX2 null</td>
<td>82.3±1.8*†</td>
<td>250±5*</td>
<td>6.09±0.07*</td>
<td>6.16±0.007*</td>
</tr>
</tbody>
</table>

Studies were done with isolated lungs from wild type or NOX2 null mice. Control = continuous perfusion. I/R = 1 h ischemia followed by 1 h reperfusion. TBARS, thiobarbituric and reactive substance. Results are mean ± SE for n=3. *P<0.05 vs I/R, wild type; † P < 0.05 vs. I/R, NOX2 null.
Figure 1.

A.

![Graph showing the effect of different treatments on Amplex Red oxidation.

- WT (Control) (0.013)
- + Ang II (0.092)
- + Ang II + 0.2 nmol MJ33 (0.071)
- + Ang II + 4 nmol MJ33 (0.017)
- gp91phoxnull + Ang II (0.020)
Figure 1.

B. WT          Nox2 null

Basal

Ang II

MJ33 + AngII

Relative fluorescence intensity

WT

NOX2 null

0

2

4

6

8

Basal AngII MJ33 + AngII

*
Figure 1.

![Normalized WST-1 absorbance](image)

- **Con A**
- **Con A + SOD**
- **Con A + MJ33**
- **Con A + Nox2 null**

Normalized WST-1 absorbance vs. Time (min)
Figure 2

A

no MJ 33

Peak Area
739

STD

Peak Area
9783329

Time (min)

B

MJ 33-IV

Peak Area
80863

STD

Peak Area
26377214

Time (min)

C

MJ 33-IT

Peak area
298490

STD

Peak area
48595729

Time (min)
Figure 3.

A.

Hours post treatment with IV MJ33

aiPLA2 activity (nmol/hr/mg protein)

- 0 nmol
- 0.4 nmol
- 4 nmol
- 10 nmol

* denotes significant difference.
Figure 3.

B.

![Graph showing aiPLA$_2$ activity against hours post treatment with IT MJ33](image)

- **aiPLA$_2$ activity (nmol/hr/mg protein)**
- **Hours post treatment with IT MJ33**

- **Graph Key**
  - Open circle: 0 nmol
  - Black square: 0.4 nmol
  - Black circle: 4 nmol
  - Black triangle: 10 nmol

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Figure 4.

A.

Number of cells

Days after MJ33 treatment

- control
- MJ33-10uM
- MJ33-25uM
Figure 4.

B.

Surviving fraction

- $48h$
- $7days$

MJ33 concentration (uM)

0 5 10 15 20 25

0.1

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Figure 5

A.

Change in body weight (g)

-4
-3
-2
-1
0
1
2

weeks post-treatment with IV MJ33

0 nmol
250 nmol
500 nmol
2500 nmol
2500 x4 nmol

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Figure 5

B.

Change in body weight (g)

weeks post-treatment with IT MJ33
Figure 6

A. This article has not been copyedited and formatted. The final version may differ from this version.

IT

Control

+MJ33

IV

200 μm

200 μm
Figure 6

B. Control +MJ33

IT

IV

Scale bar: 50 μm
Figure 7.