Decreased lung ischemia-reperfusion injury in rats after preoperative administration of cyclosporine and tacrolimus

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Objectives: Calcineurin inhibitors reduce experimental reperfusion injury in the liver, brain, heart, kidney, and small bowel. These studies were undertaken to determine whether these agents are similarly protective against lung ischemia-reperfusion injury.

Methods: Left lungs of male rats were rendered ischemic for 90 minutes and reperfused for as long as 4 hours. Treated animals received cyclosporine A (INN: ciclosporin; 1 or 5 mg/kg) or tacrolimus (0.2 mg/kg) 6 hours before ischemia, at reperfusion, or 2 hours after reperfusion. Injury was quantitated in terms of tissue polymorphonuclear leukocyte accumulation (myeloperoxidase content), vascular permeability (iodine 125–labeled bovine serum albumin extravasation), and bronchoalveolar lavage leukocyte content. Separate tissue samples were processed for nuclear protein and cytokine messenger RNA.

Results: Treatment with cyclosporine (5 mg/kg) or tacrolimus (0.2 mg/kg) 6 hours before reperfusion reduced lung vascular permeability by 54% and 56% relative to control animals (P < .03). The protective effects of cyclosporine and tacrolimus treatment before reperfusion correlated with 42% and 43% reductions in tissue polymorphonuclear leukocyte (myeloperoxidase) content (P < .008) and marked reductions in bronchoalveolar lavage leukocyte content. Separate tissue samples were processed for nuclear protein and cytokine messenger RNA.

Conclusion: Cyclosporine and tacrolimus treatment before reperfusion was protective against lung ischemia-reperfusion injury in rats. The mechanism of these protective effects may involve the inhibition of nuclear factor κB activation and expression of proinflammatory cytokine messenger RNA.

Lung reperfusion injury occurs in as many as 25% of human lung transplant recipients.1 Efforts to alleviate reperfusion injury have focused on altering native host defenses,2 improving preservation techniques,3 and minimizing ischemic times. However, lung reperfusion injury remains a problem and may increase the occurrence of acute and chronic rejection by enhancing allograft major histocompatibility complex II expression.

Cyclosporine A (CSA, INN: ciclosporin) and tacrolimus are immunosuppressive...
agents that have been used in solid-organ transplantation for 2 decades. The immunomodulatory effects of CSA and tacrolimus involve the inhibition of the protein tyrosine phosphatase calcineurin (Figure 1). CSA is a hydrophobic undecapeptide that forms a complex with the cytoplasmic protein cyclophilin that inhibits calcineurin. Tacrolimus is a macrolide antibiotic that binds to a similar cytoplasmic protein, FK-binding protein, and blocks calcineurin 100 times more effectively than does the CSA-cyclophilin complex. The inhibition of calcineurin in turn reduces the expression of a number of nuclear transcription factors, including NF-AT, Oct-1, AT-1, and NF-kB. Increase in NF-AT results in transcription of IL-2, powerful T-cell stimulant driving rejection. CSA and tacrolimus block this reaction by binding with cyclophilin and FK-binding protein to inhibit action of calcineurin.

Whereas inhibition of NF-AT may be central to the immunomodulatory effects of CSA and tacrolimus, blockade of NF-kB may have broader anti-inflammatory effects. NF-kB has a range of activities, including modulation of cell growth and death, regulation of immune function, and apparent effects on acute reperfusion injury. The ability of CSA and tacrolimus to block NF-kB activation during acute cardiac reperfusion injury has been demonstrated. CSA and tacrolimus have also been shown to reduce acute reperfusion injury of the liver and small bowel, but the mechanism mediating these protective effects has not been elucidated, and their impact on transcriptional regulation in this setting is unknown. Present no data exist regarding the effects of CSA or tacrolimus on lung ischemia-reperfusion injury.

In vitro studies have delineated the mechanism by which CSA and tacrolimus may inhibit NF-kB activation. These studies suggest that inhibition of calcineurin blocks the degradation of I-kB, the cytoplasmic protein that binds NF-kB in its inactive form. In addition, processing of pro-form precursors of NF-kB to its functional heterodimeric form may be inhibited. The net result is a diminished concentration of active NF-kB in the nucleus, which leads to decreased production of messenger RNA for proinflammatory cytokines, neutrophil adhesion molecules, procoagulant proteins, and vasoconstrictive substances. The reduction in NF-kB activation and subsequent anti-inflammatory effects therefore appear to act through mechanisms that are distinct from the immunomodulation that occurs through NF-AT inhibition.

Studies of calcineurin inhibition during organ ischemia-reperfusion injury have methodologic problems that make clinical correlation difficult. Most animal studies investigating the effects of CSA and tacrolimus delivered the drugs during the week preceding the experiment to ensure that the animals had adequate drug levels before the ischemic insult. Although this strategy was protective in these mod-
els, it would be difficult to treat human transplant recipients accordingly, because the date of transplantation is determined by donor organ availability and therefore is unpredictable. One study that did administer the drug just before cardiac reperfusion demonstrated some protective effects that were dose dependent.5

We hypothesized that administration of CSA or tacrolimus would be protective in lung ischemia-reperfusion injury. Furthermore, treatment several hours before the ischemic insult might enhance the protective effects relative to treatment just before reperfusion or after it has begun. This would be akin to dosing patients as soon as an appropriate donor organ has been identified. In addition, we considered it important to investigate doses of CSA and tacrolimus that would be appropriate for human recipients. Previous animal studies have administered CSA at 10 to 25 mg/kg orally across 3 to 4 days or 5 mg/kg subcutaneously over 1 week.6-8 Such dosing schedules might be nephrotoxic or otherwise detrimental if applied to human recipients. Finally, knowing that CSA and tacrolimus attenuate reperfusion injury in heart, liver, and small bowel, and that in vitro studies indicate that these effects may be mediated through NF-κB blockade, we hypothesized that protective effects of these drugs would be associated with a demonstrable reduction in NF-κB activation.

Material and Methods
Reagents
Commercial preparations of CSA (Novartis Pharmaceuticals Corporation, East Hanover, NJ) and tacrolimus (Fujisawa Healthcare, Inc, Deerfield, Ill) were obtained from the inpatient pharmacy at the University of Washington. All other reagents were purchased from Sigma (St Louis, Mo) unless otherwise specified.

Animal Model
Pathogen-free adult male Long-Evans rats (Simonsen Laboratories, Inc, Gilroy, Calif) weighing 280 to 320 g were used for all experiments. The University of Washington Animal Care Committee approved all experimental protocols. Animals were anesthetized with 30 to 35 mg of intraperitoneally administered pentobarbital, shaved, and prepared for surgery. A 14-gauge angiocatheter was inserted into the trachea through a midline neck incision and secured with a 4–0 braided suture. Animals were then placed on a Harvard Rodent Ventilator (Harvard Apparatus, Inc, Holliston, Mass) with an inspired oxygen content of 60%, a rate of 80 breaths/min, and 2 cm H2O of positive end-expiratory pressure. Maximal peak pressures were maintained below 10 cm H2O. All animals received 0.4 mg atropine intramuscularly, and anesthesia was maintained with inhaled halothane. Dissection was conducted with an operating microscope, and a warming blanket was placed underneath each animal throughout the experiment.

A left anterolateral thoracotomy in the fifth intercostal space was performed. The left lung was mobilizedatraumatically, and the inferior pulmonary ligament was divided sharply. Animals then received 50 units of intravenous heparin in saline solution (total volume 500 μL). Five minutes after heparin was administered, the left pulmonary artery, veins, and main stem bronchus were occluded with a noncrushing microvascular clamp. During the experiment the lungs were kept moist with periodic application of warm isotonic sodium chloride solution, and the incision was covered to minimize evaporative losses. The period of ischemia was constant at 90 minutes. At the end of the ischemic period the clamp was removed from the hilum, and the lung was allowed to ventilate and reperfuse for periods as long as 4 hours. Animals received subcutaneously 0.5 mL warm saline solution per hour to maintain hydration during the experiment. At the end of the reperfusion period a midline incision from the neck to the pubis was made to allow access to the chest and abdomen. Blood samples were obtained from the inferior vena cava just before the animal was killed. The heart-lung block was rapidly excised, and the pulmonary circulation was flushed through the main pulmonary artery with 20 mL isotonic sodium chloride solution. The lungs were then separated from the heart and mediastinal tissues and analyzed as outlined here. A time-matched sham control group underwent the same procedure except that the microvascular clamp was not applied to the hilum (thoracotomy alone).

Treated animals received CSA intravenously at 1 mg/kg or 5 mg/kg or tacrolimus intravenously at 0.2 mg/kg. Drugs were mixed in 500 μL of sterile phosphate-buffered saline solution just before administration and then injected through the penile vein 5 minutes before reperfusion or 2 hours after removal of the hilar clamp. Additional groups of animals were treated with intravenous CSA or tacrolimus 6 hours before the beginning of the experiment. A control group of animals received cremaphor vehicle (the solvent for CSA and tacrolimus) at matched time points.

Lung Permeability Index
To quantitate lung injury related to ischemia and reperfusion, a lung permeability index was determined in the following manner. Iodine 125–radiolabeled bovine serum albumin (125I-BSA) was obtained from NEN Life Science Products, Inc (Boston, Mass). Before use of the 125I-BSA in vivo, serial dilutions were performed to obtain an activity of 800,000 counts/min (cpm) per dose. This volume of 125I-BSA, approximately 2 μL of the stock solution, was then brought to a final volume of 500 μL in a 1% BSA and phosphate-buffered saline solution. Five minutes before removal of the hilar clamp or at an equivalent time in the sham control group, the 125I-BSA preparation was intravenously injected. Immediately before the animal was killed, 1 mL blood was drawn from the inferior vena cava. The heart-lung block was then excised and flushed as described previously. The radioactivity was then quantitated for the left and right lungs and for inferior vena cava blood with a gamma counter. The permeability index was then expressed as the ratio of the cpm in the left lung to the cpm in 1.0 mL of inferior vena caval blood as follows: permeability index = cpm in left lung/cpm in 1.0 mL blood. This ratio corrected for any variation in systemic blood levels of radioactivity and provided a reproducible measure of lung microvascular permeability.

Myeloperoxidase Assay
Tissue myeloperoxidase (MPO) content was used to quantitate neutrophil accumulation in the lungs. After flushing, the pulmonary circulation the lungs were homogenized for 60 seconds in a solution of 0.5% hexadecyltrimethylammonium bromide and...
5-mmol/L ethylenediaminetetraacetic acid (EDTA) in 50-mmol/L potassium-phosphate buffer (pH 6.0). Samples were then sonicated for 40 seconds in four 10-second bursts. The homogenized tissue was maintained on ice between all tissue processing steps. Samples were then centrifuged at 2300g for 30 minutes at 4°C and the supernatants were recovered. Assay buffer was composed of 0.0005% hydrogen peroxide and 0.167-mol/L o-dianisidine dihydrochloride in 100-mmol/L potassium phosphate buffer (pH 6.0). A 50-μL aliquot of each sample was mixed with 1.45 mL of assay buffer, and the change in absorbance at 460 nm over 1 minute was recorded.

**Bronchoalveolar Lavage**

Additional animals underwent bronchoalveolar lavage (BAL) at the time they were killed. Through an extended median sternotomy, a 14-gauge angiocatheter was placed and the lungs were lavaged individually with 3.0 mL cold sterile saline solution. Individual lung BAL analysis was accomplished by clamping the contralateral hilum. At least 80% of the instilled fluid was recovered from each lung. This fluid was centrifuged (1500g for 8 minutes at 4°C) to pellet the cells. The red blood cells were lysed, and the pellet was resuspended in isotonic sodium chloride solution. Cells were then counted and a differential count was determined with a hemocytometer (Hauser Scientific Partnership, Horsham, Pa).

**Electrophoretic Mobility Shift Assay**

Additional left lungs were snap frozen at the end of the experiment. These lungs were stored at -80°C. Lung RNA was isolated in guanidine thiocyanate, with two rounds of acid phenol and chloroform extraction and alcohol precipitation. RNA integrity was confirmed by agarose gel electrophoresis and quantitated by optical density measurements at 260 nm. RNA from each lung was evaluated with the Riboquant system (BD PharMingen, San Diego, Calif); rat cytokine template was used for detection of cytokines. In vitro transcription was carried out in transcription buffer supplemented with uridine triphosphate labeled with α-phosphorus 32 (3000 Ci/mmol; Amersham Pharmacia Biotech, Inc, Piscataway, NJ) and T7 RNA polymerase. After deoxyribonuclease I treatment, the riboprobe was isolated by phenol and chloroform extraction and ammonium and ethanol precipitation. Labeling efficiency was determined by measuring Chernov activity in a scintillation counter. Each riboprobe was diluted to the optimal activity (defined by the manufacturer), added to 20 μg of kidney RNA, heated to 90°C, allowed to cool to 56°C, and annealed overnight. After ribonuclease and proteinase K treatment, protected RNA hybrids were purified by phenol and chloroform extraction and ammonium acetate and ethanol precipitation and then separated by electrophoresis on 5% polyacrylamide/8-mol/L urea gels. Gels were dried and subjected to autoradiography with Kodak Biomax MS2 Film (Eastman Kodak Company, Rochester, NY). Triplet samples for each specimen were analyzed. Densitometric analysis was done with ImageJ software (version 1.2).

**Statistical Analysis**

All data were presented as mean ± SEM unless otherwise designated. Comparisons between multiple groups were made with analysis of variance. Comparisons between individual groups were made with a 2-tailed Student t test.

**Results**

Four different control groups were generated. Negative control animals were those that did not undergo any surgical manipulation. Sham (thoracotomy only) control animals were placed on the ventilator and underwent a left thoracotomy and as long as 5.5 hours of mechanical ventilation without occlusion of the hilum. A third group underwent thoracotomy and the occlusion of left hilum for 90 minutes. These animals were killed before any reperfusion. The final group was subjected to 90 minutes of ischemia followed by 4 hours of reperfusion (Figure 2). The permeability index in animals undergoing thoracotomy alone (0.18 ± 0.02) was double that seen in unmanipulated lungs (0.09 ± 0.006, P < .04). Animals that were subjected to 90 minutes of ischemia without reperfusion demonstrated increases in permeability that were not significantly different from the sham thoracotomy control group (0.18 ± 0.02 vs 0.22 ± 0.005, P = .20). However, a marked increase in permeability was seen in animals that underwent 4 hours of reperfusion after 90 minutes of ischemia (0.75 ± 0.01, P < .001 relative to unmanipulated control group).

Positive control animals that received phosphate-buffered saline solution had a mean permeability index of 0.75 ± 0.01, and those receiving Cremophor vehicle had a...
mean index of 0.76 ± 0.03 (Figure 3). Animals that received CSA or tacrolimus 2 hours after reperfusion did not demonstrate any reduction in vascular permeability relative to positive control animals. In contrast, treatment at the time of reperfusion (Figure 3) reduced permeability by 17% in animals receiving CSA at 5 mg/kg (P < .03) and by 18% in animals treated with tacrolimus at 0.2 mg/kg (P < .04). Administration of low-dose CSA (1 mg/kg) at reperfusion provided no protection relative to positive control animals. Treatment with tacrolimus or CSA at either dose 6 hours before the initiation of the experiment decreased lung injury significantly (Figure 3). The permeability index in animals pretreated with CSA at 1 mg/kg was reduced by 33% (P < .008); with CSA at 5 mg/kg the reduction was 54% (P < .004), and with tacrolimus at 0.2 mg/kg the reduction was 56% (P < .003). Thus advance treatment with CSA and tacrolimus offered greater and more consistent protection than did treatment at the time of reperfusion or 2 hours after reperfusion had begun.

MPO Activity

MPO activity was measured in lungs from unmanipulated control animals and in lungs from animals subjected to 90 minutes of ischemia and reperfusion (P < .001). Animals that underwent thoracotomy alone and ischemia alone also had statistically significant increases in permeability relative to unmanipulated control animals (P < .04; n = 4 in all experimental groups).

Reperfusion was associated with a steady increase in MPO activity that was detectable at 2 hours after reperfusion and marked by 3 (0.26 ± 0.01) and 4 hours (0.41 ± 0.04, P < .002; Figure 4). In an additional experiment with an extended time course, the tissue MPO content did not continue to increase at 6 or 8 hours and began to decrease at 12 hours (data not shown). MPO content thus appeared to peak after 4 hours of reperfusion.

MPO analysis was performed on lungs from animals that received CSA at 5 mg/kg 6 hours before ischemia, at the time of reperfusion, or 2 hours after the onset of reperfusion. MPO was also assessed in lungs from animals that received tacrolimus at 0.2 mg/kg 6 hours before ischemia (Figure 5). A statistically significant decrease in MPO content was seen only among animals that were treated in advance with CSA or tacrolimus. Animals treated with CSA or tacrolimus 6 hours before ischemia demonstrated 38% and 36% decreases in tissue neutrophil accumulation, respectively (P < .008). Treatment with CSA or tacrolimus at the time of reperfusion or 2 hours after the onset of reperfusion was not associated with a reduction in lung MPO.

**BAL Leukocyte Content**

BAL analysis was undertaken to evaluate leukocyte content in the distal airways and alveolar compartment. Baseline BAL leukocyte counts were recorded in four groups: unmanipulated control group, sham thoracotomy control group, ischemia-alone control group (90 minutes of ischemia alone), and reperfusion injury group (90 minutes of ischemia followed by 4 hours of reperfusion; Figure 6). Animals undergoing sham thoracotomy or ischemia alone did not demonstrate a significant increase in BAL leukocyte content. However, a statistically significant increase in BAL leukocyte content was seen after 4 hours of reperfusion (P < .005). The predominant cell type at the end of the reperfusion period in this group was the neutrophil, whereas all the other groups demonstrated mainly alveolar macrophages (>95%).

As seen with tissue neutrophil accumulation, BAL cell count was reduced only in animals treated before ischemia with CSA or tacrolimus (Figure 6). Treatment at the time of reperfusion or 2 hours later did not reduce BAL leukocyte count (162 ± 14 and 146 ± 15, respectively). In animals receiving CSA or tacrolimus 6 hours before ischemia, the BAL leukocyte count was 32% lower than the positive control values (P < .01).

**Electrophoretic Mobility Shift Assay Analysis for NF-κB**

Shown in Figure 7 is a representative electrophoretic mobility shift assay (EMSA) that details the expression of NF-κB in the nucleus of control and treated left lungs. There was no NF-κB detected in the negative control animals. In contrast, significant activation of NF-κB was apparent in the animals that underwent ischemia and reperfusion. The band specific for NF-κB was delineated with a nonradiolabeled (cold) oligonucleotide. The location of this band is demonstrated in the lane marked “cold competition.” Animals that received treatment before ischemia with CSA and tacrolimus appeared to have attenuated NF-κB expression in the nucleus. These differences are quantitated in the densitometric analysis depicted below the autoradiograph.
Ribonuclease Protection Assay

Ribonuclease protection assays were performed on lung extracts from four experimental groups, unmanipulated negative control animals, animals that underwent 90 minutes of ischemia and 4 hours of reperfusion, and animals that underwent ischemia and reperfusion after advance treatment with CSA at 5 mg/kg or tacrolimus at 0.2 mg/kg (Figure 8). In unmanipulated control animals there were few transcriptional products in the left lung. In contrast, animals that underwent ischemia and reperfusion of the left lung demonstrated significantly increased messenger RNA expressions for a variety of cytokines (IL-3, IL-5, IL-1β, tumor necrosis factor α (TNF-α), IL-2, and interferon γ). The housekeeping genes demonstrated similar activities in all lanes and are shown at the bottom of the gel. This indicates that loading was equal, and relative differences in cytokine messenger RNA expressions assessed by densitometry are therefore meaningful and valid. In injured lungs from animals treated before ischemia with CSA or tacrolimus, there were decreased expressions of several inflammatory and...
regulatory cytokines relative to positive control lungs. These differences were quantitated in the densitometric analysis shown in Figure 9.

Discussion
The utility of CSA and tacrolimus for induction and maintenance immunosuppressive therapy after solid-organ transplantation has been demonstrated during the past 20 years. The immunomodulatory effects are mediated through transcriptional blockade of IL-2 production. These studies explored the possibility that these calcineurin inhibitors regulate inflammatory effects through a different transcriptional mechanism. One isolated study demonstrated that administration of CSA after reperfu-

Figure 5. MPO activity in treated animals. Tissue neutrophil accumulation was modestly decreased in animals receiving CSA at 5 mg/kg 2 hours after reperfusion and just before reperfusion, but decrease was not significant (ns). Animals that received CSA at 5 mg/kg and tacrolimus (TAC) at 0.2 mg/kg 6 hours before experiment demonstrated 38% and 36% decreases in MPO content, respectively. These were statistically significant (P < .008). Each experimental group included 4 animals.

Figure 6. BAL cell count increased significantly in those animals undergoing ischemia followed by reperfusion relative to unmanipulated control animals (P < .005). Small increases in cell count were seen with thoracotomy alone and with ischemia alone, but these increases were not significant. Predominant cell type in animals was alveolar macrophage, and most cells in unperfused lungs were neutrophils. Each experimental group included 6 animals. No relative decrease in alveolar cell count was seen in animals that received CSA at 5 mg/kg after 2 hours of reperfusion or at time of reperfusion. However, significant decrease in alveolar leukocyte accumulation was seen in animals that received CSA at 5 mg/kg and tacrolimus (TAC) at 0.2 mg/kg 6 hours before experiment, resulting in 32% increase in BAL cell counts. Asterisk indicates P < .01.
isolation was protective against cardiac ischemia-reperfusion injury. This group studied isolated rabbit hearts that were perfused with buffer on a Langendorff apparatus. The ischemic time was limited to 30 minutes, and CSA was given 10 minutes into reperfusion. This is consistent with our finding that some protection was afforded by higher dose of CSA given at the time of reperfusion. However, in our study lung vascular permeability was not decreased in animals receiving CSA or tacrolimus after the onset of reperfusion (Figure 3). This is supportive of a transcriptional regulatory mechanism of action. Specifically, CSA and tacrolimus may modulate ischemia and reperfusion through blockade of NF-κB. Ischemia alone or ischemia followed by a short period of reperfusion activates NF-κB. Thus the blockade of NF-κB activation would be most effective if the intervention were to be performed before ischemia.

Protection from reperfusion injury is correlated with a decreased expression of NF-κB, as seen in the EMSA in Figure 7. Selective blockade of TNF-α with a polyclonal rat anti–TNF-α antibody afforded 45% protection from reperfusion injury, and combined blockade with anti–TNF-α and anti–IL-1β antibodies provided 60% protection (unpublished data). The degree of protection offered by CSA or TAC pretreatment (54%-56%) is therefore impressive. Greater protection may be difficult to achieve because of incomplete transcriptional blockade or redundancy in proinflammatory mediators released in the lung.

The protection from vascular injury seen with calcineurin inhibitors administered 6 hours before ischemia was correlated with a decrease in neutrophil accumulation in both lung tissue and the alveolar compartment. Only animals treated before ischemia with CSA or tacrolimus demonstrated significant reductions in tissue neutrophil accumulation. These findings support conclusions of earlier studies regarding the ability of CSA and tacrolimus to reduce polymorphonuclear neutrophil accumulation in reperfused heart and liver.

The EMSA suggests that NF-κB activation was blocked in the animals that received CSA or tacrolimus before ischemia. This is the first direct demonstration of NF-κB blockade by calcineurin inhibitors in an intact animal model. Others have assumed that NF-κB is involved because IκB activity is preserved with tacrolimus treatment. Multiple investigators have established that NF-κB activation is decreased by calcineurin inhibitors in vitro. This phenomenon has been investigated specifically in human bronchial epithelial cells and suggests that these events may be important in the development of acute lung inflammation. Several mechanisms through which calcineurin in-
Inhibitors may decrease NF-κB activity have been postulated. Calcineurin affects cellular kinetics and activation through its action as a calcium- and calmodulin-dependent serine and threonine phosphatase. It is believed that calcineurin blockade results in the inactivation of the 20S proteasome. The 20S proteasome is the site of degradation of IκB, the cytoplasmic inhibitor of NF-κB. Calcineurin also appears to facilitate processing of NF-κB proforms (p105) to the active heterodimer (p65). The inactivation of the 20S proteasome by calcineurin inhibitors thus results in the decreased production of active NF-κB and the preservation of cytoplasmic stores of IκB.

NF-κB blockade by CSA and tacrolimus correlates with the decreased expressions of phlogistic cytokine messenger RNA seen with the ribonuclease protection assay. Pretranscriptional blockade of NF-κB provided by CSA and tacrolimus would explain the diminution in expression of a wide variety of cytokines. TNF-α expression is regulated through some NF-κB-dependent mechanism. NF-κB activation has also been linked to the production of intracellular adhesion molecule 1, P-selectin, and IL-8. The decreased production of these adhesion molecules, chemokines, and cytokines seen with calcineurin inhibitors therefore may result in the decreased leukocyte accumulation seen in the tissue and alveolar compartment of the lung (Figures 5 and 6). Ultimately, these effects may reflect the amelioration of tissue injury seen with CSA and tacrolimus treatment.

The protection provided by calcineurin inhibitors has also been associated with inhibition of the mitochondrial transition pore opening and with variable effects on nitric oxide and endothelin metabolism. These mechanisms may also be relevant in a model of lung ischemia-reperfusion, and further investigation is needed to elucidate the relevance of these pathways in this model. Most studies of the mitochondrial transition pore have been done in models of liver and heart reperfusion injury. Mitochondrial transition pore changes may be more important in these tissues than in the lung, because heart and liver appear to have a higher concentration of mitochondria.

Protection afforded by calcineurin inhibitors may also be mediated through amelioration of postischemic vasoconstriction. Tacrolimus appears to preserve the activity of inducible nitric oxide synthase during rat brain ischemia-reperfusion injury, and both CSA and tacrolimus may suppress endothelin 1 production in rat small-bowel ischemia-reperfusion. These mechanisms may be particularly relevant to the pulmonary circulation, where amelioration of postischemic vasoconstriction may improve ventilation-perfusion mismatch and overall lung function.

CSA and tacrolimus can cause vasoconstriction in the renal vascular bed. If such effects were seen in the pulmonary circulation, alteration of Starling forces would limit the edema seen with reperfusion injury. To characterize the acute effects of CSA and tacrolimus on pulmonary vasmotor tone, pulmonary arterial pressures were measured in rats after administration of the calcineurin inhibitors. Pulmonary arterial pressures were not altered in animals receiving CSA or tacrolimus for 30 minutes after administration of the drug (data not shown). It thus appears that the...
The potential for use of CSA or tacrolimus as advance treatment for human lung transplant recipients is contingent on the ability to deliver an adequate dose of calcineurin inhibitor at the appropriate time. Other studies have administered nephrotoxic doses of CSA (10–25 mg/kg) to animals to demonstrate protective effects.\(^6\)\(^-\)\(^8\) In this study we treated the study animals with clinically relevant doses of CSA (1 and 5 mg/kg intravenously) and tacrolimus (0.2 mg/kg).
Significant protection was associated with advanced treatment at both dose levels. CSA at 1 mg/kg is a dose that is commonly used for patients with impaired creatinine clearance, so the doses used in these studies appear to be both clinically relevant and safe. Ultimately the effectiveness of this particular treatment strategy needs to be systematically studied in human beings.

In conclusion, advance treatment of rats with CSA and tacrolimus decreased lung ischemia-reperfusion injury. The protection afforded by calcineurin inhibitors was associated with a decrease in neutrophil accumulation in the lung tissue and the alveolar compartment. The mechanism mediating protection may involve the blockade of the transcription factor NF-κB, and this blockade appears to result in a decreased expression of proinflammatory cytokines.

References


Discussion

Dr Stuart W. Jamieson (San Diego, Calif). Few here will remember that for the first clinical lung transplants, 20 years ago now, all the donors were brought into the adjoining room for harvesting of the heart-lung block. Needless to say this caused considerable tactical problems, particularly in flying bodies across state lines. It was clear to everybody that better understanding of lung injury would be required to make clinical lung transplantation an effective therapy. In late 1982 we started to use modified Euro-Collins solution for lung protection, and the first long-distance lung procurement was done in December that year. Our group in San Diego has continued to use that solution exactly as formulated then, and we are comfortable with ischemic times as long as 8 hours. But most things could probably be improved, and this is important work that is well presented. It confirms that, as with other organs that are devastated by reperfusion injury, calcium-dependent signaling is an important component in the development of lung ischemia-reperfusion injury, which has been characterized here by alterations in lung vascular permeability, tissue MPO accumulation, and leukocyte transendothelial migration. There is also reduction in the expression of cytokine messenger RNA, which suggests an inhibitory effect on the calcineurin-dependent expression of NF-κB. That calcineurin inhibitors were ineffective when administered at the time of or after reperfusion further suggests interruption of an early event in the evolution of reperfusion injury, if not of a response to hypoxia itself.

I would appreciate your comments on some areas. Do you have direct evidence regarding the site of the calcineurin effect, whether this is on leukocytes, endothelial cells, or both? If the effect is
restricted to leukocytes, then most teams are probably already benefitting from this therapy, because the recipient has usually been already treated some hours before reperfusion. But if the effect is primarily on the vascular endothelium, then this would suggest that the donor ought to be treated.

Significant immunomodulation is unlikely to occur with single doses of CSA or FK-506, and you make the point that these agents exhibit differential antiinflammatory and immunomodulatory effects. Do you think that this dichotomy in effects could be related to different sensitivities to the drugs of discrete populations of leukocytes? On a related issue, what do you believe is the key effector cell in this reaction?

Finally, the ribonuclease protection assay demonstrated a generalized decrease in inflammatory cytokine content with both drugs. Have you demonstrated functional roles for these cytokines in lung ischemia-reperfusion injury?

Dr Krishnadasan. Thank you for those insightful comments. With respect to your first question, about calcineurin’s effects on leukocytes and endothelial cells, the in vitro data to date suggests, and studies have mostly concentrated on T lymphocytes, that CSA and tacrolimus do have effects at high doses on those types of cells. For inflammatory cells, such as alveolar macrophages and neutrophils, the lower doses of CSA and tacrolimus seem to be effective at blocking NF-kB activation.

The interesting question of course is what happens to the endothelial cell, and obviously in our whole-organ preparations we cannot sort out what is happening in individual cells. Mulligan’s laboratory has turned the opposite way and looked at what is going on in vitro, and it appears that in fact the endothelial cell does play an important role in these reactions. Verrier’s laboratory for the last 10 or 15 years also has been doing work indicating that the endothelial cell may be the first cell that is activated. This moves to your second question about the potential effector cell.

We believe that the effector cell in the lung is probably the alveolar macrophage. It is situated in a position that would make it ideally suited to driving the reaction, and in fact when you compare pulmonary artery endothelial cells’ and alveolar macrophages’ effects on hypoxia and reoxygenation in vitro, we find that alveolar macrophages have a much more robust response to hypoxic stimulus. This makes sense, because the alveolar macrophage is a monocytic cell that produces all the relevant cytokines that we discussed. Again, this dovetails into your third question regarding the ribonuclease protection assay and the functional roles of the cytokines.

Much of my first year in the laboratory was spent defining functional roles for various cytokines, among them TNF-α, IL-1β, some of the chemokines, macrophage inflammatory protein-1α, MIP-2, membrane cofactor protein 1, and some T helper 2 cytokines such as IL-4 and IL-10. It appears that the most protection that we are afforded by single cytokine blockade is about a 45% decrease in lung permeability, which if you compare that with the data that I presented today suggests that the effects of CSA and tacrolimus are more proximal in the cascade and are pretty remarkable compared with single-cytokine blockade.