Video Article

Angiogenesis in the Ischemic Rat Lung

John Jenkins, Elizabeth Wagner

Johns Hopkins Asthma and Allergy Center, Division of Pulmonary and Critical Care Medicine, Johns Hopkins University

Correspondence to: Elizabeth Wagner at wagnerem@jhmi.edu

URL: <http://www.jove.com/video/50217>

DOI: [doi:10.3791/50217](https://doi.org/10.3791/50217)

Keywords: Medicine, Issue 72, Anatomy, Physiology, Biomedical Engineering, Pathology, Surgery, Lung, Lung Diseases, Lung Injury, Thoracic Surgical Procedures, Physiological Processes, Growth and Development, Respiratory System, Physiological Phenomena, angiogenesis, bronchial artery, blood vessels, arteries, rat, ischemia, intubation, artery ligation, thoracotomy, cannulation, animal model

Date Published: 2/8/2013

Citation: Jenkins, J., Wagner, E. Angiogenesis in the Ischemic Rat Lung. *J. Vis. Exp.* (72), e50217, doi:10.3791/50217 (2013).

Abstract

The adult lung is perfused by both the systemic bronchial artery and the entire venous return flowing through the pulmonary arteries. In most lung pathologies, it is the smaller systemic vasculature that responds to a need for enhanced lung perfusion and shows robust neovascularization. Pulmonary vascular ischemia induced by pulmonary artery obstruction has been shown to result in rapid systemic arterial angiogenesis in man as well as in several animal models. Although the histologic assessment of the time course of bronchial artery proliferation in rats was carefully described by Weibel¹, mechanisms responsible for this organized growth of new vessels are not clear. We provide surgical details of inducing left pulmonary artery ischemia in the rat that leads to bronchial neovascularization. Quantification of the extent of angiogenesis presents an additional challenge due to the presence of the two vascular beds within the lung. Methods to determine functional angiogenesis based on labeled microsphere injections are provided.

Video Link

The video component of this article can be found at <http://www.jove.com/video/50217/>

Introduction

Systemic angiogenesis in the lung is well-recognized. In disease states such as asthma², interstitial pulmonary fibrosis³, cancer⁴, and chronic pulmonary thromboembolism⁵, the systemic vasculature in and surrounding the lung proliferates and invades the pulmonary parenchyma. However, animal models to study this differential activation of the systemic rather than the pulmonary circulation are few. Perhaps the most reproducible model of systemic neovascularization in the lung of the adult mammal is that which occurs after inducing chronic pulmonary artery ischemia. The response to left pulmonary artery obstruction in humans⁶⁻⁷, dogs⁸, pigs⁹, sheep¹⁰, guinea pigs¹¹, rats^{1, 12, 13}, and mice¹⁴ is the rapid proliferation of the bronchial artery as well as intercostal arteries. The mechanisms responsible for systemic neovascularization of the lung after pulmonary ischemia are largely unknown and have not been widely studied. The time course of bronchial angiogenesis in the rat after left pulmonary artery obstruction has been carefully described in the histologic work of Weibel¹. Extending this work in the rat, our laboratory has focused on both the growth factors important in this process as well as the physiologic outcome of this neovascularity in the lung. Results demonstrate the CXC chemokine CINC-3 is elevated early after ischemia and treating rats with a neutralizing antibody to CXCR2, the receptor for CINC-3, attenuates angiogenesis¹³. The newly established bronchial vasculature 14 days after the onset of pulmonary ischemia was shown to be abnormal with significantly increased protein permeability¹⁵. Left lung function was not normal showing decreased diffusing capacity and a decrease in lung volume¹⁵. Although the neovascularity may have contributed to the preservation of lung tissue during chronic pulmonary ischemia, it appears not to be normal and may contribute to a sustained decrease in pulmonary function.

Perhaps one of the most curious aspects of this model relates to the spatial distribution of proliferating blood vessels. Despite the release of growth factors within the pulmonary parenchyma due to ischemia, the neovascularity originates in relatively large upstream bronchial arteries. The normal bronchial artery arises as a small branch from the aorta and invades the airway tree at the carina. Thus the mechanism by which growth factors induce the initial phase of arteriogenesis is not clear. We suggest that the rat, with a vascular anatomy similar to humans, provides a unique opportunity to study the mechanisms responsible for systemic angiogenesis during pulmonary ischemia. Although complete obstruction of the left pulmonary artery is a rare occurrence in human subjects, increased bronchial vascularity appears to be similarly induced in patients whatever the site and size of pulmonary artery obstruction¹⁶. Thus, we provide a detailed description of the surgical approach to ligate the left pulmonary artery in rats and a means to quantify the magnitude of angiogenesis.

Protocol

All protocols performed on rats have been approved by the Johns Hopkins University Animal Care and Use Committee and in accordance with NIH guidelines. Whenever possible, the animal should be surgically prepped in an area separate from the surgical area to minimize contamination of the surgical site.

1. Anesthesia/analgesia

1. Place rat (Sprague Dawley male rats, 125-150 g; Harlan, Indianapolis, IN) in an induction chamber infused with 3% isoflurane.
2. Place anesthetized rat on surgery board attached to nose cone and ventilator with 3% isoflurane anesthetic. Use sterile technique for all procedures. Drape rat to ensure sterile operating field.

2. Intubation

1. Immobilize appendages in supine position using surgical tape.
2. Remove rat from nose cone, extend tongue with padded forceps.
3. Use 14 gauge intracath with blunted metal stilet as guide. Slide behind tongue, into trachea.
4. Remove metal stilet leaving white plastic intracath in trachea. Ensure the rat is breathing and air is flowing through the tube.
5. Replace nose cone with direct adapter to catheter, connect rat to ventilator (90 breaths/min; 8 ml/kg tidal volume; Rodent Ventilator Model 683, Harvard Apparatus, Holliston, Massachusetts).
6. Apply Puralube (Butler Schein, Dublin, OH) veterinary ointment on eyes.

3. Thoracotomy

1. Place rat right side down, immobilize appendages using surgical tape.
2. Shave left side ribcage area.
3. Remove excess fur with small vacuum.
4. Make sterile field by wiping down with alcohol followed by Povidone-Iodine swabstick (Dynarex Corporation, Orangeburg, NY). Repeat this process two more times (for a total of three scrubs).
5. Make transverse incision with sterile dissecting scissors or sterile scalpel in center of field.
6. Blunt dissect through layers of tissue and fat down to ribs (last layer is thin membrane covering ribs).
7. Count ribs to determine 3rd intercostal space.
8. Using sterile 45 ° Graefe forceps, make blunt incision between 3rd and 4th rib.
9. Insert rib separators, pull gently creating open void with full visualization of lung, place tape on sutures to keep open.

4. Left Pulmonary Artery Ligation

1. With 90 ° Graefe forceps, move left lung back with right hand.
2. Using Dumont pattern #5 straight forceps, grab left pulmonary artery and airway with left hand. Left pulmonary artery will lay on top of airway. Ensure that forceps are directly perpendicular to the table. Additionally, it is most effective to pick up the left pulmonary artery/ left mainstem bronchus at the most distal position (closest to parenchyma). Push the ventilating left lung aside with this left hand maneuver.
3. Use Dumont pattern #5 45 ° curved forceps to separate the left pulmonary artery from the left mainstem bronchus at their natural borders. This separation line appears thin and white between the two individual structures.
4. Separate directly under artery without going through the vessel; smoothly slide the forceps tips held together along the separation.
5. Continue until tips of forceps have visibly separated the left pulmonary artery and the left mainstem bronchus. Only a small point of tip needs to be fully through. If blood can be visualized on the tip of the forceps, then it is not fully through and ligation should not be attempted. Once through the space, the left pulmonary artery will lay on the curve of the forceps. Hold in this position.
6. Gently release straight blunt forceps grip (left hand) and grab piece of a pre-cut suture (~2-3 inches; polypropylene suture size 6-0; Myco Medical, Cary, NC).
7. Open curved forceps cradling left pulmonary artery and grab suture. Gently pull suture through space between left pulmonary artery and left mainstem bronchus in an upward motion relative to the curve of the forceps.
8. Tie down occluding the left pulmonary with a square knot, carefully snip remainder of suture.
9. Close ribs using blunt forceps to hold rib and suture twice with polypropylene (blue monofilament) size 4-0 attached to a 19 mm, 3/8 circle reverse cutting needle (Myco Medical, Cary, NC) in a hemostat, being careful not to suture skin (only ribs).
10. Complete a loose square knot, inflate the lung, place on positive end-expiratory pressure (PEEP; 2-5 cmH₂O), hyperinflate lung then secure knot tightly, and make another full knot before snipping remainder of suture. Remove from PEEP and visualize for 30 sec to ensure that the lung does not collapse. Apply 5 drops Bupivacaine (APP Pharmaceuticals, Schaumburg, IL).
11. Close skin by placing tissue glue on wound and push skin together using back end of cotton tip applicator. Give Bupivacaine (2.0 mg/kg subcutaneous) at the site of incision subsequently every 8 hr for 24 hr or until the animal resumes normal activity. As the subcutaneous layer in this area is very thin, it is not easy to suture on its own. When the tissue glue is applied and the skin is pushed together it also closes the subcutaneous layer.
12. Turn off isoflurane gas but continue to ventilate the rat for 1-2 min on room air until voluntary movement returns. Disconnect the tracheal tube from the ventilator and ensure that the rat is breathing spontaneously before removing it.

13. Wipe off Puralube from eyes with cotton swab and monitor movement and recovery. Inject buprenorphine hydrochloride (0.05 mg/kg intraperitoneal, Butler Schein, Dublin OH). Continue delivery of analgesic every 12 hr for 48 hr after surgery.

5. Left Carotid Artery Cannulation

To assess the magnitude of bronchial perfusion of the ischemic left lung at desired time points after left pulmonary artery ligation, inject labeled microspheres through the left carotid artery into the aortic arch. Prepare rats as above 1-2.

1. Cut midline along neck, blunt dissect to reveal trachea and left carotid artery (microsphere injection site).
2. Insert catheter filled with heparinized saline, blunt tip of PE20 tubing (Becton Dickinson, Sparks MD) into vessel connected to 25 g needle, 4-way stopcock, 1 ml syringe.
3. Place bottle of microspheres (15 μ m crimson polystyrene fluorescent microspheres, 1 x 10⁶ spheres/ml; Invitrogen, Eugene, OR) in water sonicator for 30 sec.
4. Remove bottle, vortex and draw up 0.5 ml (500,000 microspheres) into a 1 ml Hamilton glass syringe (Hamilton Company, Reno, NV) through a 20 g needle.
5. Attach Hamilton syringe to 4-way stopcock and infuse microspheres with syringe pump (rate: 500 μ l/min; Genie Plus, Kent Scientific, Torrington, CT).
6. Remove Hamilton syringe and flush apparatus with 1 ml of heparized saline at 500 μ l/min.
7. Perform full chest thoracotomy and exsanguinate the rat by severing the inferior vena cava.
8. Remove the left lung and other tissues of interest.
9. To extract microspheres from tissue, take the entire left lung of the rat after exsanguination and place it in 2M KOH (4-6 ml). Place in a 55 °C water bath and leave overnight for tissue digestion. Add Tween 80 (0.25%) to wash the beads, vortex (10 sec) and centrifuge (2,000 rpm; 20 °C for 10 min). Remove the supernatant, add 2-ethoxyethyl acetate (1 ml), vortex and let stand for 1 hr. Vortex the suspension and centrifuge (2,000 rpm; 20 °C for 10 min). Remove the 2-ethoxyethyl acetate aqueous layer containing the fluorescence, place in a cuvette and measure using a Hitachi F-2500 fluorescence spectrophotometer (excitation 612/emission 618; Digilab, Holliston, MA).

Representative Results

Vascular cast: Results of the effects of left pulmonary artery ischemia in the rat are depicted in **Figure 1**. Shown is a methacrylate cast of the bronchial vasculature and the extensive vascularity of the left airway tree 28 days after LPAL. To obtain this cast, the systemic vasculature was injected with a methacrylate mixture (red), retrograde into the descending aorta and the trachea was cannulated and injected with a white silicon based material. This vascular cast provides a remarkable visualization of bronchial angiogenesis in the lung. Especially noteworthy are the large, tortuous upstream bronchial vessels of the left lung. Although not shown in this image, the right bronchus is essentially devoid of large vessels except for the normal, lone, right bronchial artery.

Histology: Changes in airway vascularization visualized in left lung histologic sections are shown in **Figure 2**. Bronchial vessels observed 3 days and 14 days after LPAL are shown. **Figure 2A** shows an airway section from the left lung 3 days after LPAL. Note prominent bronchial vessels located within the airway wall. The inset is from a serial section of 2A stained for PCNA (proliferating cell nuclear antigen). Note the PCNA positive endothelial cells lining the bronchial vessel. **Figure 2B** shows a left lung section from a rat 14 days after LPAL. Note the increased size of the bronchial vessels at this time point and the withered pulmonary artery.

Functional angiogenesis: To assess the magnitude of systemic perfusion of the left lung through the bronchial arteries, fluorescent microspheres were infused into the occluded left carotid artery such that all microspheres mixed in the aortic arch as part of the cardiac output. Fluorescence from this size of microsphere (15 μ m) was virtually undetected in the left lung of naïve rats. However there was a substantial, significant and consistent level measured in the left lung from rats studied 14 days after LPAL.

Figure 1. Red methacrylate cast of the tortuous neovasculature associated with the left bronchus (white) 28 days after left pulmonary artery ligation (LPAL).

Figure 2A. Airway section from left rat lung 3 days after left pulmonary artery (PA) ligation showing bronchial vessels adjacent to a larger pulmonary artery. Inset shows a serial section stained with PCNA to identify proliferating endothelial cells of the bronchial artery (BA). Distance bar indicates 200 μ m.

Figure 2B. Left lung section taken from rat 14 days after LPAL. Note the substantial increase in the size of the bronchial arteries. Distance bar indicates 500 μ m.

Figure 3. Analysis of fluorescent microspheres lodged in the left lung of naïve rats (n=2 rats) and 14 days after left pulmonary artery ligation (LPAL; n=3 rats).

Discussion

Left pulmonary artery ligation in all species tested leads to robust systemic neovascularization of the ischemic lung. We have presented the details of the surgical approach in a rat model. Our results produced by vascular casting, histopathology, and *in vivo* labeling demonstrate that bronchial arteries proliferate and perfuse the pulmonary parenchyma. Thus, the mechanisms of bronchial angiogenesis can be studied in an animal model that parallels the human condition of chronic pulmonary thromboembolism. Furthermore, this clinically relevant small animal model provides an opportunity to study the form and function of neovascularization where human experimentation is not tenable.

The surgical procedures listed provide sufficient detail such that an animal technician with surgical experience should be able to quickly master these techniques. Although our observations have been predominantly in Sprague Dawley male rats (75-100 g), other strains are equally appropriate for study. We have not systematically evaluated the age-dependence of the extent of bronchial angiogenesis. We have also provided details for the cannulation of the left carotid artery. We have used the measurement of fluorescent microspheres infused retrograde into the left carotid artery that mix in the aorta and subsequently lodge in the left lung as a marker for systemic perfusion of the ischemic lung and consequently, functional angiogenesis. We acknowledge that this measurement may be variable due to changes in cardiac output during the time of infusion. However, simultaneous measurement of systemic blood pressure continuously in rats before and during this measurement procedure indicated no changes in cardiovascular status. Furthermore, results presented in **Figure 3** demonstrate a small acceptable range

of variability within a group of rats. Interestingly, the naïve animals showed a minimally detectable level of fluorescence suggesting a very low bronchial perfusion in normal rats. Using microspheres of a smaller size may allow for detection of a different basal level of perfusion.

Our results show a well-developed vasculature by 28 days after LPAL. The methacrylate cast of the bronchial vessels associated with the left bronchus provides a remarkable view of the extensive and tortuous neovasculature that forms in response to pulmonary ischemia. Perhaps less dramatic but consistent with the vascular cast are the images obtained from histologic sections of the left lung 14 days after LPAL (**Figure 2B**). Rats were exsanguinated prior to tissue harvest for the section shown. However, the growth of the bronchial vessels is especially obvious when comparing the size of bronchial vessels to histological sections of left lung obtained 3 days after LPAL (**Figure 2A**). Bronchial endothelium shows clear signs of proliferation as assessed using PCNA staining. Thus the process of angiogenesis begins early after the onset pulmonary ischemia.

Our methods to determine the extent of the neovasculature have ranged from early histologic assessment of proliferating bronchial endothelium to microsphere measurement of fully functioning, perfusing bronchial vessels and vascular casting. Our intent is to provide a perspective of the process of bronchial angiogenesis over time. Proliferating vessel counting requires precision of the evaluator, adequate sample size, and an assumption that vessels associated with the airways are the primary locus of early angiogenesis. Assessment of microspheres lodging in the lung requires a fully developed angiogenic vasculature, consequently a late phenomenon. Additionally, the method assumes adequate mixing of spheres during infusion, no changes in cardiac output during the procedure, and that changes in the number of lodged spheres reflects changes in the number of perfusing vessels and not changes in vasoreactivity. We acknowledge the continued challenge to better define the process of systemic angiogenesis in an organ that normally has two distinct vascular beds and pulmonary networks appear to remain patent during ischemia. Endothelial labels in the rat have not demonstrated a unique angiogenic phenotype separating the two vasculatures histochemically. *In vivo* imaging is hampered by both ventilatory and cardiac motion. We continue to search for additional methods to quantify the extent of neovascularization with greater precision. The challenge remains to identify a small but growing systemic vascular bed among other non-perfusing pulmonary capillary networks.

The mechanisms responsible for ischemia-induced angiogenesis are not fully understood. Past work implicates the CXC-chemokines as playing a role at some stage in the overall process¹³. Because the left lung in this model is ischemic but fully ventilated, hypoxia-inducible factors are unlikely to play a role in the systemic neovascularization that takes place. It should be emphasized that this situation is unlike other organs where ischemia is accompanied by tissue hypoxia. Thus the growth factors responsible for complete bronchial vessel growth within the lung need to be completely defined. Although the growth of new vessels appears to be essential for the preservation of ischemic tissue, angiogenic bronchial vessels have been shown to be dilated, pro-inflammatory, and hyperpermeable^{15, 17}. The chronic consequences of these abnormal vessels include persistent airway narrowing due to vasodilation and airway edema¹⁸.

Additional questions related to this model pertain to the site of growth factor release and the responding vessels. Despite pulmonary ischemia, it is the systemic bronchial vessels that respond to the ischemic stimulus and not other pulmonary vessels within the lung. It is not clear how the ischemic stimulus is conveyed to upstream bronchial vessels. Despite a multitude of growth factors likely released by ischemic parenchymal tissue, how vessels originating from the aorta are induced to proliferate uniquely to the left lung is not clear. Future experimentation is required to address these questions.

Disclosures

No conflicts of interest declared.

Acknowledgements

We acknowledge the work of Dr. Adlah Sukkar, M.D. in assisting with casting of the lung. This work has been funded by NHLBI, HL088005.

References

1. Weibel, E.R. Early stages in the development of collateral circulation to the lung in the rat. *Circulation Research*. **8**, 353-76 (1960).
2. Li, X. & Wilson, J.W. Increased vascularity of the bronchial mucosa in mild asthma. *Am. J. Respir. Crit. Care Med.* **156**, 229-33 (1997).
3. Turner-Warwick, M. Precapillary systemic-pulmonary anastomoses. *Thorax*. **18**, 225-37 (1963).
4. Muller, K.M. & Meyer-Schwickerath, M. Bronchial arteries in various stages of bronchogenic carcinoma. *Pathol. Res. Pract.* **163**, 34-46 (1978).
5. Remy-Jardin, M., Duhamel, A., *et al.* Systemic Collateral Supply in Patients with Chronic Thromboembolic and Primary Pulmonary Hypertension: Assessment with Multi-Detector Row Helical CT Angiography. *Radiology*., 274-81 (2005).
6. Karsner, H. & Ghoreyeb, A. Studies in infarction: The circulation in experimental pulmonary embolism. *J. Exp. Med.* **18**, 507-22 (1913).
7. Endrys, J., Hayat, N., *et al.* Comparison of bronchopulmonary collaterals and collateral blood flow in patients with chronic thromboembolic and primary pulmonary hypertension. *Heart*. **78**, 171-6 (1997).
8. Virchow, V. Über die Standpunkte in den Wissenschaftlichen Medizin. *Virchow Archiv*. **1**, 1-19 (1847).
9. Fadel, E., Mazmanian, G.M., *et al.* Lung reperfusion injury after chronic or acute unilateral pulmonary artery occlusion. *Am. J. Respir. Crit. Care Med.* **157**, 1294-300 (1998).
10. Charan, N.B. & Carvalho, P. Angiogenesis in bronchial circulatory system after unilateral pulmonary artery obstruction. *J. Appl. Physiol.* **82**, 284-91 (1997).
11. Shi, W., Hu, F., *et al.* Altered reactivity of pulmonary vessels in postobstructive pulmonary vasculopathy. *J. Appl. Physiol.* **88**, 17-25 (2000).
12. Shi, W., Giaid, A., *et al.* Increased reactivity to endothelin of pulmonary arteries in long-term post-obstructive pulmonary vasculopathy in rats. *Pulm. Pharmacol. Ther.* **11**, 189-96 (1998).
13. Sukkar, A., Jenkins, J., *et al.* Inhibition of CXCR2 Attenuates Bronchial Angiogenesis in the Ischemic Rat Lung. *J. Appl. Physiol.* **104**, 1470-5 (2008).

14. Mitzner, W., Lee, W., *et al.* Angiogenesis in the mouse lung. *Am. J. Pathol.* **157**, 93-101 (2000).
15. Wagner, E.M., Jenkins, J., *et al.* Lung and vascular function during chronic severe pulmonary ischemia. *J. Appl. Physiol.* **110**, 538-44 (2011).
16. Remy-Jardin, M., Bouaziz, N., *et al.* Bronchial and nonbronchial systemic arteries at multi-detector row CT angiography: comparison with conventional angiography. *Radiology.* **233**, 741-9 (2004).
17. Baluk, P., Tammela, T., *et al.* Pathogenesis of persistent lymphatic vessel hyperplasia in chronic airway inflammation. *J. Clin. Invest.* **115**, 247-57 (2005).
18. Bailey, S.R., Boustany, S., *et al.* Airway vascular reactivity and vascularisation in human chronic airway disease. *Pulm. Pharmacol. Ther.* **22**, 417-25 (2009).