

Review Article

Medical Progress

INTERVENTIONAL PULMONOLOGY

LUIS M. SEIJO, M.D., AND DANIEL H. STERMAN, M.D.

INTERVENTIONAL pulmonology is a new field within pulmonary medicine focused on the use of advanced bronchoscopic and pleuroscopic techniques for the treatment of a spectrum of thoracic disorders ranging from tracheobronchial stenosis to pleural effusions associated with malignant tumors.¹⁻⁴

TECHNIQUES USED IN INTERVENTIONAL PULMONOLOGY

Rigid Bronchoscopic Debulking or Balloon Dilation

Therapeutic use of bronchoscopy began over 100 years ago with the development of rigid bronchoscopes. These instruments have beveled tips, which are ideal for coring through large tumors in the airways and for dilating strictures, and they have large internal diameters, which facilitate débridement of tumors, evacuation of clots, and ventilation.⁵ Despite advances in other adjunctive endoscopic techniques, rigid bronchoscopic recanalization remains the treatment of choice for life-threatening tracheobronchial obstruction (Fig. 1).

Balloon dilation has become an attractive alternative to dissection with a blunt rigid bronchoscope in less urgent cases of obstruction caused by malignant tumors. The technique is best used in combination with bronchoscopic laser therapy and placement of a tracheobronchial stent for the treatment of airway stenosis.⁶ Balloon bronchoplasty has also been used successfully to treat other disorders, including tuberculosis,⁷ fibrosing mediastinitis, and strictures associated with lung transplantation⁸ or prolonged intubation.⁶ It is less successful when used alone to treat stenosis accompanied by extrinsic airway compression and is contraindicated in patients with tracheobronchomalacia.^{8,9}

Although in the majority of cases, balloon dilation

is performed while the patient is under general anesthesia, treatment of certain airway lesions (e.g., short fibrotic strictures) can be accomplished with the use of a fiberoptic bronchoscope while the patient is under conscious sedation. Complications of balloon dilation of airway lesions include bronchospasm, chest pain, perforation of the airway, pneumothorax, and pneumomediastinum.^{6,8}

Endobronchial Laser Therapy

Perhaps the most widely known technique in interventional pulmonology is laser bronchoscopy. Lasers produce a beam of monochromatic, coherent light that can induce tissue vaporization, coagulation, hemostasis, and necrosis. Although primarily useful in the ablation of endoluminal malignant tumors, bronchoscopic laser therapy is also beneficial for the treatment of other tracheobronchial disorders, including inflammatory strictures, obstructive granulation tissue, amyloidosis, and benign tumors such as hamartomas.¹⁰⁻¹²

In 1976, Laforet et al. reported on endobronchial laser ablation of an obstructive neoplasm.¹³ Since then, several types of lasers have become available for the management of tracheobronchial obstruction. The carbon dioxide laser, used mainly by otolaryngologists, allows shallow penetration of tissue (to a depth of 0.1 to 0.5 mm) and highly precise cutting, but it has minimal hemostatic properties and must be used through a rigid bronchoscope.^{11,14} It is ideal for the management of laryngeal lesions. Interventional pulmonologists primarily use the neodymium:yttrium–aluminum–garnet (Nd:YAG) laser, which provides deeper penetration of tissue (to a depth of 3 to 5 mm), superior photocoagulation, and improved hemostasis, but with less precision in cutting. Photocoagulation with an Nd:YAG laser can be performed through a rigid or flexible bronchoscope, but rigid bronchoscopy remains the preferred method for the treatment of patients who have respiratory distress due to severe tracheobronchial obstruction.

The use of a laser in the tracheobronchial tree requires careful consideration of the anatomical location and configuration of the lesion. If the lesion is in close proximity to the esophagus or the pulmonary artery, endobronchial laser therapy poses a risk of fistula formation. Use of laser therapy in a patient with tracheobronchial narrowing due to extrinsic compression may result in perforation of the airway. In addition, prolonged obstruction of the airway (for more than six weeks) may lead to refractory atelectasis or bronchiectasis, minimizing the benefits of endobronchial recanalization.

From the Interventional Pulmonology Program, Division of Pulmonary, Allergy, and Critical Care Medicine, University of Pennsylvania Medical Center, Philadelphia. Address reprint requests to Dr. Sterman at the Interventional Pulmonology Program, Division of Pulmonary, Allergy, and Critical Care Medicine, Department of Medicine, Hospital of the University of Pennsylvania, 833 W. Gates Pavilion, 3400 Spruce St., Philadelphia, PA 19104, or at sterman@mail.med.upenn.edu.

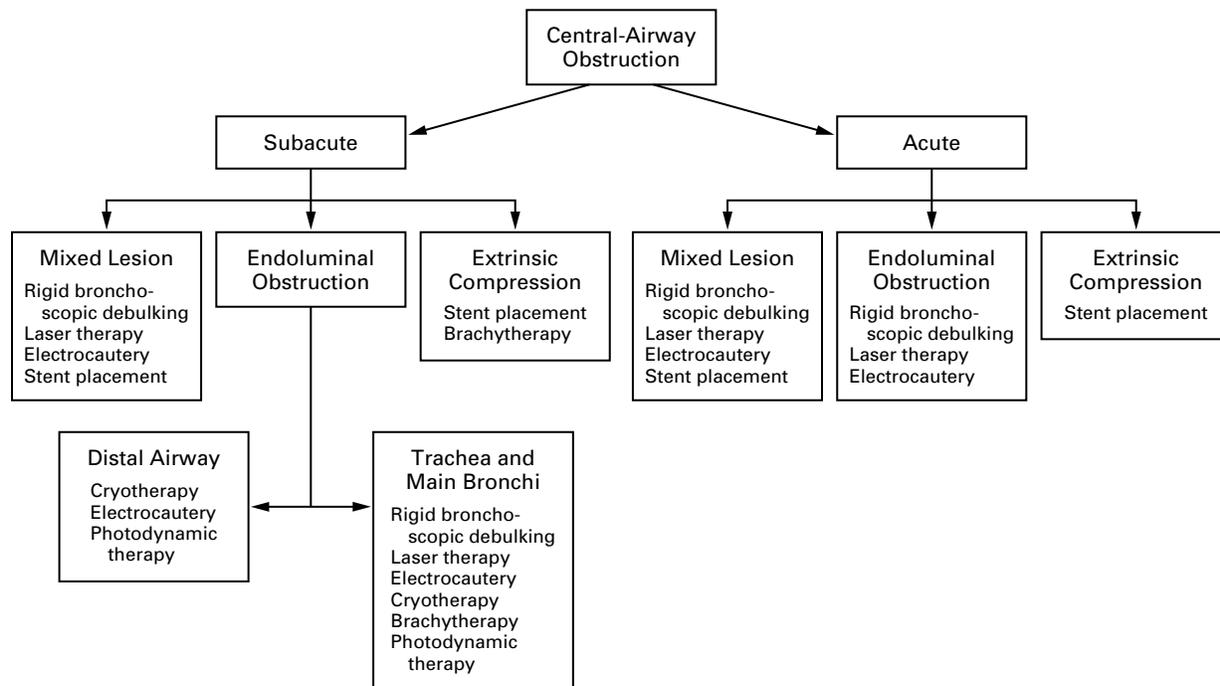


Figure 1. Algorithm for the Management of Central-Airway Obstruction Due to a Malignant Tumor.

Obstruction of the trachea and main bronchi by advanced primary or metastatic lung cancer is often a medical emergency necessitating immediate intervention. In many cases, surgical resection is impossible because of the advanced stage of the tumor or the presence of severe underlying lung disease. Therefore, interventional procedures are necessary to reestablish the patency of the airway and prevent overt respiratory failure. This treatment algorithm addresses both acute and subacute central-airway obstruction, as well as the location and type of lesion — characteristics that can be determined by computed tomography and diagnostic bronchoscopy. Endobronchial stent placement is generally reserved for airway lesions associated with some degree of extrinsic compression, although patients who have residual mucosal tumors after recanalization may also benefit from the placement of a stent. Cryotherapy, brachytherapy, and photodynamic therapy are not used to treat acute airway obstruction because of the delay in achieving tumor necrosis with these approaches. Endobronchial laser therapy is not recommended for distal airway lesions because of the risk of delayed perforation of the bronchus and adjacent structures.

Although endobronchial laser therapy is generally safe and well tolerated, it may be complicated by cardiac arrhythmias,¹⁵ perforation of the airway, pneumothorax, hemorrhage, hypoxemia, or endobronchial fire (ignition of the bronchoscope or endotracheal tube).^{4,16-18} In rare cases, pulmonary edema or fatal pulmonary venous gas embolism has been reported.¹⁹⁻²¹ Patients with endotracheal tubes and those who require high concentrations of supplemental oxygen are at increased risk for endobronchial fire. Fortunately, the overall risk is less than 0.1 percent.^{11,22} The overall rate of mortality attributable to endoscopic laser therapy is quite low, not exceeding 0.3 to 0.5 percent in several large series.^{16,17,22,23}

In a study of Nd:YAG laser therapy in approximately 1800 patients with endobronchial obstruction due to malignant tumors, recanalization rates exceeded 90 percent for tumors located in the trachea, main-stem bronchi, and right intermediate bronchus.⁴ The treat-

ment was less successful in patients with peripheral lesions (recanalization rate, 50 to 70 percent) or with associated extrinsic airway compression. Serious complications, such as hemorrhage, pneumothorax, or cardiorespiratory failure, occurred in less than 3 percent of all patients.

Several retrospective case series have compared survival rates among patients undergoing laser therapy for advanced lung cancer with survival rates among historical controls.^{24,25} Laser therapy was associated with increased survival rates among patients with lung cancer who required emergency treatment to restore airway patency.²⁴ In addition, Stanopoulos et al. reported successful weaning from mechanical ventilation after endoscopic laser debulking in 9 of 17 patients with advanced lung cancer (53 percent) who presented in respiratory failure.²⁶ Colt and Harrell reported similar findings when laser therapy was combined with placement of an endobronchial stent.²⁷

Photocoagulation with an Nd:YAG laser is an invaluable treatment for patients with airway obstruction due to benign endoluminal tumors. In a study reported by Shah and colleagues, complete resection was achieved with a single laser treatment in 62 percent of patients with benign tumors.¹⁰ Other investigators have reported similar findings, with rates of complete resection ranging from 50 percent to 80 percent.²⁸⁻³¹

Endobronchial Cryotherapy and Electrocautery

Cryotherapy and electrocautery are excellent, cost-effective alternatives to laser therapy for the management of tracheobronchial obstruction. The effects of electrocautery on tissue are similar to those of the Nd:YAG laser,³² whereas cryotherapy probes induce tissue necrosis through hypothermic cellular crystallization and microthrombosis.³³ As with the Nd:YAG laser, both electrocautery and cryotherapy can be administered through a rigid or flexible bronchoscope. Flexible probes are used more often, however, because they do not require the administration of general anesthesia, with its associated risks.

Cryotherapy and electrocautery have been used successfully to relieve airway obstruction caused by benign tracheobronchial tumors, polyps, and granulation tissue.³⁴ These techniques may be superior to lasers for distal lesions because of the lower risk of airway perforation. Similarly, carcinoma in situ and mucosal dysplasia may be adequately treated with cryotherapy or electrocautery alone, although multiple treatments may be required for optimal results.^{33,35} Cryotherapy is a safe treatment for infiltrative lesions of the airway,³⁶ and according to anecdotal reports, it has proved beneficial in patients with post-transplantation anastomotic stenosis³⁷ and in those with foreign-body aspiration.³⁸

Endobronchial cryotherapy is generally not effective for paucicellular lesions that are relatively impervious to freezing, such as fibrotic stenoses, cartilaginous or bony lesions, and lipomas.³⁹ Furthermore, endobronchial cryotherapy, unlike either laser therapy or electrocautery, cannot be used to achieve rapid relief of symptomatic airway obstruction.^{34,39} The most common serious complication of both electrocautery and cryotherapy is hemorrhage. The estimated incidence of clinically significant bleeding in patients treated with electrocautery is 2.5 percent.³²

Endobronchial Brachytherapy

Brachytherapy is the treatment of tumors with radiation delivered internally through implanted radioactive seeds or inserted wires. This technique ensures the delivery of a maximal therapeutic dose of radiation to the tumor with a minimal effect on normal surrounding tissues.⁴⁰ Endobronchial brachytherapy involves the bronchoscopic insertion of a thin, hollow catheter through a malignant obstruction under fluoroscopic guidance.³³ A radioactive implant is then inserted into

the catheter and left in position for a predetermined period (2 to 40 hours, depending on the dose rate).

In 1922, Yankauer reported the use of rigid bronchoscopic brachytherapy for the palliation of airway obstruction due to malignant tumors.⁴⁰ Modern techniques, including the use of flexible bronchoscopes, polyethylene afterloading catheters, and iridium-192 implants, were first described in 1983.⁴¹ Since the development of techniques involving a high dose rate in the 1980s,⁴² endobronchial brachytherapy has become a particularly attractive option for outpatient treatment.

Relief of airway obstruction is the primary goal of endobronchial brachytherapy, although curative treatment may be attempted in conjunction with external-beam radiation in selected patients.^{43,44} Brachytherapy is safest and most effective for central airway lesions, although in one study, small peripheral tumors proved to be more responsive than bulkier central tumors.⁴⁵ Among patients with obstruction due to malignant tumors, rates of recanalization range from 60 percent to 90 percent, with decreased dyspnea, cessation of hemoptysis, and relief of cough in most cases.^{40,46-48} Brachytherapy has also been used for the prevention and treatment of airway stenosis related to recurrent growth of granulation tissue in patients with lung transplants.⁴⁹

Endobronchial brachytherapy may require multiple treatments to be effective. It is generally used as an adjunct to either photocoagulation with an Nd:YAG laser or conventional external-beam irradiation in an effort to achieve both rapid and sustained recanalization in patients with obstruction due to malignant tumors.^{47,50,51} Brachytherapy may also be administered in conjunction with the placement of an endobronchial stent in patients with extrinsic compression of the airways due to malignant tumors.

Serious complications of brachytherapy include massive hemoptysis and fistula formation.^{52,53} Because of the risk of fatal hemorrhage, every effort should be made to rule out the involvement of central vessels before treatment is administered. The incidence of serious complications varies widely, with rates as low as 0 to 10 percent in some of the largest studies^{40,54,55} and as high as 30 to 42 percent in two small studies.^{56,57}

Photodynamic Therapy

Photodynamic therapy is currently approved by the Food and Drug Administration for the palliation of airway obstruction caused by malignant tumors and as an alternative to surgery in selected patients with minimally invasive central lung cancer.^{58,59} Photodynamic therapy works on the principle that certain compounds, such as hematoporphyrin derivatives, function as photosensitizing agents, rendering malignant cells susceptible to damage from monochromatic light.⁶⁰ The selective effect of photodynamic therapy on malignant cells is thought to be due to the greater up-

take and retention of photosensitizing agents in neoplastic cells than in normal cells.⁶¹ This effect appears to be most pronounced approximately 24 to 48 hours after infusion of the photosensitizing agent. For this reason, bronchoscopic treatment of target lesions is often performed one to two days after the agent has been injected.⁶² Subsequent bronchoscopy is often required to débride necrotic tissue.⁵⁹

Ideal candidates for photodynamic therapy include patients with airway obstruction due to malignant polypoid endobronchial masses, with minimal extrinsic airway compression, and patients with minimally invasive tumors of the central airways.⁶² Metastatic tumors have also been treated successfully with photodynamic therapy.⁶³⁻⁶⁵ In patients with bulky tumors, endobronchial photodynamic therapy may substantially reduce the obstruction, with objective increases in spirometric measurements and subjective improvements in dyspnea and the quality of life.^{65,66} Complications of photodynamic therapy include increased photosensitivity of the skin³⁵ and hemoptysis resulting from extensive tumor necrosis.⁶⁷

Tracheobronchial Stenting

The medical term “stent” was first used to denote a device that supported the healing of gingival grafts, developed by the British dentist Charles R. Stent.⁶⁸ The term has since been used to refer to any device designed to maintain the integrity of hollow tubular structures, such as the coronary arteries and the esophagus. Anecdotal reports of attempts to use stents in the tracheobronchial tree date back to 1915.⁶⁹ The Montgomery T tube, designed in the 1960s, was the first reliable, dedicated airway stent.⁷⁰ However, stent implantation in the lower trachea and bronchi did not become standard medical practice until Dumon's 1990 report on the safety and ease of placement of a dedicated airway stent made of silicone.⁷¹

There are two main types of endobronchial stents in use today: tube stents made of silicone or plastic, and expandable metallic stents.⁶⁸ Silicone stents, including the Dumon stent, are usually placed with the use of a rigid bronchoscope while the patient is under general anesthesia.^{4,72} These stents are inexpensive and easy to remove from the airway; they provide protection from tumor ingrowth and cause minimal irritation to adjacent normal tissues.^{4,68,72-75} Potential complications of silicone stents include migration, formation of granulation tissue, and inspissation of secretions. Bifurcated stents are also available for the palliation of main carinal lesions (Fig. 2). These stents have been effective in the management of carinal compression associated with malignant tumors, tracheoesophageal fistulas, and tracheobronchomalacia.⁷⁶

Unlike silicone stents, metal stents can be placed with the use of a flexible bronchoscope, and they are less likely to migrate and are more likely to preserve normal mucociliary clearance.^{68,69,77} Metal stents re-

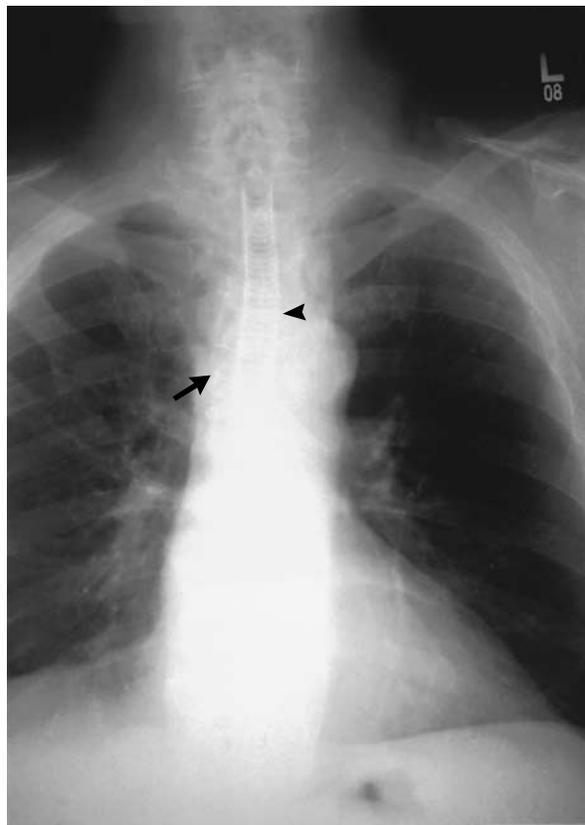


Figure 2. Posteroanterior Chest Radiograph of a 66-Year-Old Man with Diffuse Malacia of the Trachea and Main Bronchi Due to Mediastinitis after Coronary-Artery Bypass Surgery.

A bifurcated stent made of silicone and stainless steel (arrowhead) was inserted into the airway at the level of the main carina (arrow). This so-called dynamic stent is anatomically similar to the native trachea, with stainless-steel struts to support the anterior and lateral walls and a soft, pliable posterior membrane to allow effective clearance of secretions. The mediastinum is radiopaque because of the patient's mediastinitis.

main fairly expensive, however, and if they are misplaced in the airway, rigid bronchoscopy is often required for their removal.^{69,72} In addition, mucosal inflammation and the formation of granulation tissue are common at the proximal and distal ends of metal stents, and endoscopic intervention may be required to restore airway patency.

Endobronchial stents have a critical role in a multimodal endoscopic approach to both benign and malignant stenoses of the airways.⁷² Stenosis due to locally advanced bronchogenic carcinoma, for example, can be treated with a combination of endoscopic laser therapy and stent implantation in order to prevent respiratory failure (Fig. 3 and 4). Stent placement can also be used to maintain airway patency after endobronchial brachytherapy⁴⁰ or can be combined with laser therapy and balloon dilation in the endoscopic management of fibrotic strictures.⁷⁸

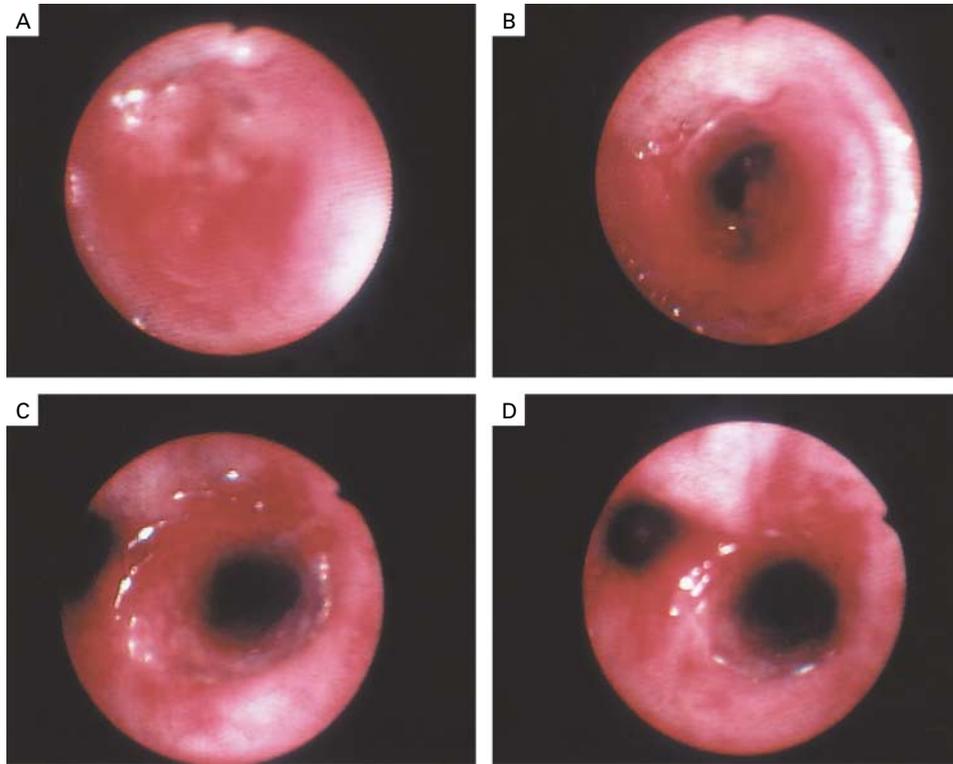


Figure 3. Bronchoscopic Images Obtained after External-Beam Radiation Therapy in a 67-Year-Old Man with Stage IIIA Non-Small-Cell Carcinoma of the Left Upper Lobe.

Atelectasis of the left upper lobe developed after radiation therapy. The left-upper-lobe bronchus was completely obstructed by friable tissue (Panel A). Endobronchial biopsy revealed only granulation tissue, with no evidence of tumor. After recanalization with an Nd:YAG laser, the distal airways were visible (Panel B). There was residual fibrotic narrowing of the orifice of the left upper lobe. An uncovered, expandable metal stent (Wallstent, Schneider, Plymouth, Minn.) was inserted into the bronchus (Panel C), with full expansion after balloon dilation (Panel D).

Most studies of the efficacy of endobronchial stent placement have had impressive results. Dumon and colleagues reported excellent clinical outcomes and few complications with the use of silicone stents in patients with extrinsic airway compression due to malignant tumors but a lower success rate among patients with tracheal stenosis caused by other disorders.⁷⁹ The most common complications in this study were migration of the stent, formation of granulation tissue, and inspissation of secretions. Rates of success, broadly defined as symptomatic relief, in smaller studies have ranged between 78 percent and 98 percent.^{68,80,81} In two small studies of patients who had been intubated because of respiratory failure due to unresectable tracheobronchial and mediastinal disease, stent placement facilitated extubation in nearly all the patients.^{82,83}

The benefits of stent placement appear to persist in patients who survive for a period of several months or years after implantation. Long-term follow-up data, however, are limited to benign disease,^{74,75,84-88} since the mean follow-up period in patients with airway

compression due to malignant tumors does not usually exceed three to four months.^{79,89,90} Some authors have reported poor long-term results with the use of metal stents in patients with fibroinflammatory stenosis due to nonmalignant disorders.⁹¹ In addition, there have been case reports of massive hemorrhage associated with the use of stents in patients with extrinsic compression attributable to aneurysmal dilatation or congenital malformations of the aorta.⁷⁶

Pleuroscopy

This procedure, also known as medical thoracoscopy, was described in Europe by Jacobeus, a pulmonologist who adapted a rigid cytoscope for insertion into the pleural cavity.⁹² Pleuroscopy was initially used for the diagnosis and management of tuberculous lung disease, but it was abandoned once effective antimycobacterial drugs became widely available.⁹³ The recent emphasis on minimally invasive surgical techniques has led to a renewed interest in the procedure.

Pleuroscopy as performed by interventional pulmo-

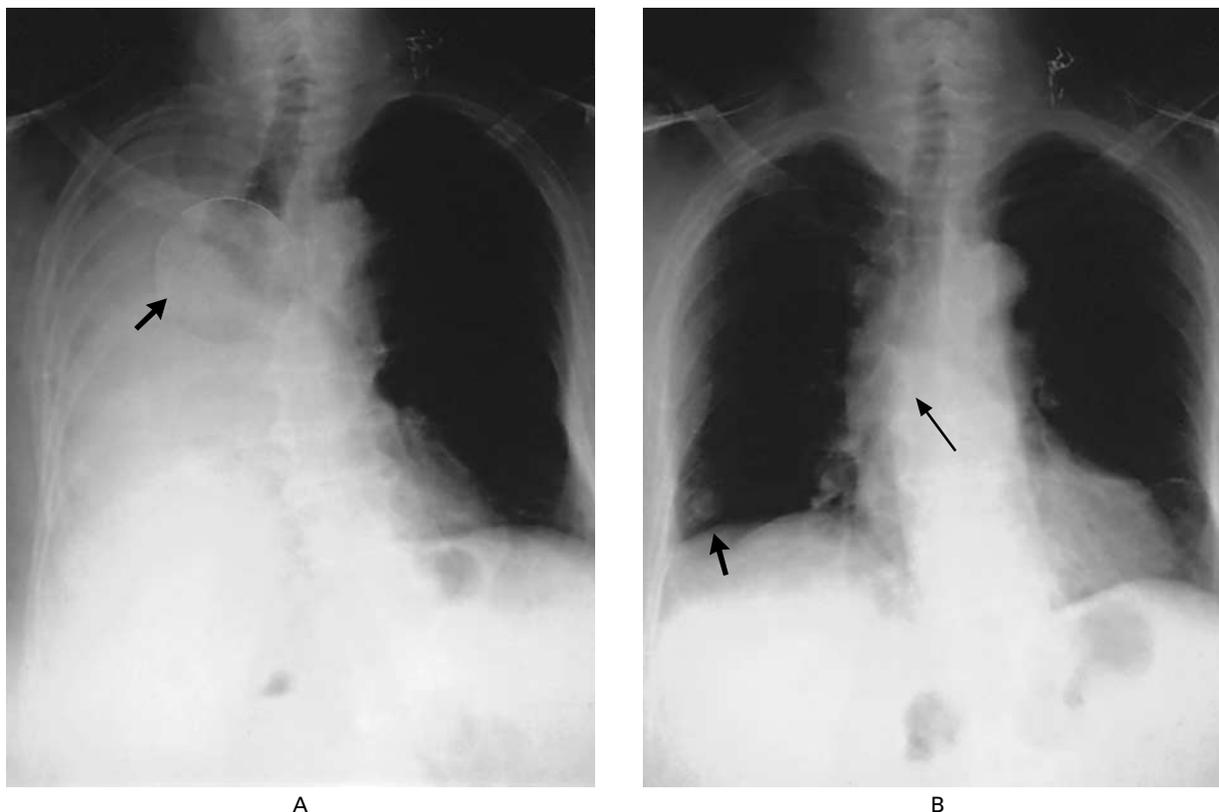


Figure 4. Chest Radiographs of a 66-Year-Old Woman with Severe Dyspnea and Hemoptysis and a History of Endometrial Carcinoma. Complete atelectasis of the right lung was noted, with an ipsilateral shift of the trachea and mediastinum and a bulging mass in the region of the right main bronchus (Panel A, magnified area, arrow). Diagnostic bronchoscopy showed a fungating tumor emanating from the orifice of the right main bronchus and extending into the distal trachea. Nd:YAG-laser photocoagulation was performed, with debulking of the endobronchial tumor and insertion of a metal stent (Wallstent) in the right main bronchus (Panel B, long arrow). A metastatic focus is visible in the inferolateral aspect of the right lower lobe (short arrow).

nologists differs from video-assisted thoracic surgery, in that local anesthesia and conscious sedation are most often used in lieu of general anesthesia, a single thoracic puncture is made rather than multiple incisions, and the procedure can be safely performed in an ambulatory care setting.^{94,95} Although pleuroscopy is primarily used for the diagnosis and management of pleural disorders, it can also be used to perform lung biopsy⁹⁶ and manage spontaneous pneumothorax.⁹⁷ Mortality rates associated with pleuroscopy are extremely low, ranging from 0.01 percent to 0.24 percent.⁹⁸⁻¹⁰⁰ Complications of the procedure include bleeding, persistent pneumothorax, and intercostal nerve or vessel injury.⁹⁴

COMMON CLINICAL APPLICATIONS

Endoluminal Airway Obstruction

Endoluminal obstruction of the tracheobronchial tree may result from various benign and malignant

processes. The most common cause of endobronchial obstruction is advanced bronchogenic carcinoma. In patients with inoperable tumors of the central airways, restoration of airway patency may provide palliation and may even prolong life, particularly in the case of impending respiratory failure.²⁴

Signs and symptoms of central airway obstruction vary but often include wheezing, cough, stridor, hoarseness, hemoptysis, and chest pain. A careful pre-treatment evaluation should be performed to distinguish symptoms attributable to focal tracheobronchial lesions from those related to underlying diffuse airflow obstruction, parenchymal lung disease, or both. Mild-to-moderate tracheal stenosis, for example, may contribute only marginally to the degree of dyspnea experienced by a patient with severe chronic obstructive lung disease. Although pulmonary-function testing and thoracic imaging techniques such as computed tomography (CT)^{101,102} and magnetic resonance imaging¹⁰³ may be useful in the evaluation of a patient

with suspected obstruction of the central airway, bronchoscopy, either rigid or flexible, remains the diagnostic gold standard.¹

The bronchoscopic approach to the management of endoluminal obstruction depends on the location of the lesion, the presence or absence of associated extrinsic compression, and the degree of clinical urgency (Fig. 1). Rigid-bronchoscopic debulking, with adjunctive laser therapy or electrocautery, is recommended when airway recanalization must be performed on an emergency basis. If endobronchial obstruction is accompanied by marked extrinsic compression, the placement of a stent may be beneficial.

The complexity of a lesion is equally important in determining the best approach to resection. Tracheal webs, for example, are often managed by laser resection alone, whereas complex fibrotic strictures may warrant the combination of rigid-bronchoscopic or balloon dilation, laser resection, and stent placement (Fig. 3 and 4). For focal tracheal stenoses in low-risk patients, surgical resection and primary reanastomosis should remain the first-line therapy.¹⁰⁴

Extrinsic Airway Compression

Extrinsic airway compression usually results from malignant involvement of structures adjacent to the central airways, such as mediastinal lymph nodes or the esophagus, but it may be associated with a benign process, such as fibrosing mediastinitis,¹⁰⁵ tuberculosis,¹⁰⁶ aneurysmal dilatation of the aorta,⁷⁶ or sarcoidosis.¹⁰⁷ The clinical signs and symptoms of extrinsic airway compression often mimic those of endobronchial obstruction. The diagnosis is established on the basis of bronchoscopic evidence of marked narrowing of the airway in the absence of an endobronchial mass. Chest CT has an important adjunctive role in identifying anatomical structures external to the narrowed lumen.

Therapeutic options in the management of extrinsic airway compression are limited. Ablative endoscopic approaches such as laser therapy, cryotherapy, photodynamic therapy, and electrocautery are contraindicated because of the risk of airway perforation. Although some patients with malignant disease may benefit from endobronchial brachytherapy, tracheobronchial stent placement is the palliative treatment of choice for patients with symptomatic extrinsic airway compression.

Tracheobronchomalacia

Diffuse or focal tracheobronchomalacia is perhaps the most challenging disorder encountered by the interventional pulmonologist. Cartilaginous tracheobronchomalacia, as seen in patients with post-intubation injury or relapsing chondritis,¹⁰⁸ reflects a loss of the structural integrity of the trachea or mainstem bronchi due to destruction of the airway's cartilaginous rings. Membranous, or crescentic, trache-

obronchomalacia is manifested by airway collapse during exhalation as a result of laxity of the membranous portion of the trachea and main bronchi and is usually seen in patients with long-standing chronic obstructive pulmonary disease.¹⁰⁹ Focal tracheobronchomalacia may be a complication of long-standing intubation⁶⁸ or an anastomotic complication after lung transplantation.¹¹⁰ Tracheobronchomalacia is best diagnosed on the basis of fiberoptic bronchoscopy, with the patient breathing spontaneously, although dynamic CT scanning, with images obtained on inspiration and expiration, is often helpful.

The endoscopic treatment of choice for patients with diffuse tracheobronchomalacia is the insertion of a silicone or expandable metal stent. This intervention is more likely to be successful in patients with the cartilaginous type of tracheobronchomalacia than in those with the membranous type. For many patients with focal tracheobronchomalacia, surgery is the best therapeutic option. An alternative treatment for selected patients with diffuse tracheobronchomalacia is the pneumatic stent afforded by noninvasive ventilatory techniques such as nasal continuous positive airway pressure.^{111,112}

Pleural Effusions Due to Malignant Tumors

Large, symptomatic pleural effusions resulting from primary or secondary malignant processes affect substantial numbers of patients with primary breast or lung cancer at some point during the course of their illness.³ Such patients often present with dyspnea and chest pain. Although thoracentesis can confirm the underlying diagnosis and temporarily alleviate dyspnea, repeated thoracentesis does not provide long-term control of respiratory symptoms. In fact, pleural effusions due to malignant tumors recur an average of four or five days after thoracentesis.¹¹³ Similarly, tube thoracostomy alone is ineffective for the long-term control of malignant effusions.¹¹⁴

The interventional pulmonologist can offer several therapeutic options to patients with symptomatic, recurrent pleural effusions. Patients who have complete lung reexpansion after thoracentesis but in whom pleural fluid reaccumulates rapidly may be good candidates for tube thoracostomy, followed by chemical pleurodesis with doxycycline, bleomycin, or talc slurry.¹¹⁵⁻¹¹⁷ Pleurodesis with talc slurry has proved effective in preventing the reaccumulation of pleural fluid in 72 percent to 95 percent of cases.^{115,118-120} Alternatively, patients may opt for outpatient pleurodesis with the insertion of a small-caliber chest tube outfitted with a one-way valve, which allows for drainage at home.¹²¹ Semipermanent, implantable pleural catheters such as the Pleurx catheter (Denver Biomedical, Golden, Colo.) can induce sufficient pleural irritation by themselves to achieve pleurodesis in approximately 46 percent of patients.¹²²

Alternatively, pleuroscopy may be performed. This

procedure is the gold standard in the diagnosis of pleural effusions caused by malignant disorders, with a sensitivity of more than 90 percent,^{123,124} as compared with 60 to 80 percent for large-volume thoracentesis and closed pleural biopsy.⁹² Talc pleurodesis with the use of pleuroscopy, which can be performed during a brief hospital stay, with minimal side effects, provides effective control of pleural effusions due to malignant conditions in 71 to 100 percent of patients.^{123,125-127} Pleuroscopic talc poudrage may be unsuccessful in achieving pleurodesis in patients with lung parenchyma that cannot reexpand because of bulky pleural disease, long-standing exudative effusions, or a large, obstructive endobronchial mass.^{120,128,129} Patients with complex effusions and multiple adhesions may benefit from traditional video-assisted thoracic surgery.

CONCLUSIONS

In the past decade, tremendous technical advances have been made in thoracic endoscopy and the adjunctive use of lasers, stents, and electrocautery probes. Although primarily palliative, interventional pulmonology may soon provide definitive alternatives to surgery for conditions such as early-stage lung cancer and benign endobronchial tumors. Advances in molecular biology and immunology will undoubtedly lead to further progress in this new discipline. Ultimately, interventional pulmonologists may use bronchoscopic and pleuroscopic techniques to administer gene therapy or immunotherapy, or both, for the treatment and prevention of disorders as diverse as cystic fibrosis and lung cancer.

We are indebted to Dr. John Hansen-Flaschen for his helpful comments and to Drs. Michael Unger and Larry Kaiser for their guidance.

REFERENCES

1. Mehta AC, Harris RJ, De Boer GE. Endoscopic management of benign airway stenosis. *Clin Chest Med* 1995;16:401-13.
2. Baharloo F, Veyckemans F, Francis C, Bietlot MP, Rodenstein DO. Tracheobronchial foreign bodies: presentation and management in children and adults. *Chest* 1999;115:1357-62.
3. Colt HG. Thoracoscopic management of malignant pleural effusions. *Clin Chest Med* 1995;16:505-18.
4. Cavaliere S, Venuta F, Foccoli P, Toninelli C, La Face B. Endoscopic treatment of malignant airway obstructions in 2,008 patients. *Chest* 1996;110:1536-42. [Erratum, *Chest* 1997;111:1476.]
5. Helmers RA, Sanderson DR. Rigid bronchoscopy: the forgotten art. *Clin Chest Med* 1995;16:393-9.
6. Noppen M, Schlessler M, Meysman M, D'Haese J, Peche R, Vincken W. Bronchoscopic balloon dilatation in the combined management of postintubation stenosis of the trachea in adults. *Chest* 1997;112:1136-40.
7. Nakamura K, Terada N, Ohi M, Matsushita T, Kato N, Nakagawa T. Tuberculous bronchial stenosis: treatment with balloon bronchoplasty. *AJR Am J Roentgenol* 1991;157:1187-8.
8. Sheski FD, Mathur PN. Long-term results of fiberoptic bronchoscopic balloon dilation in the management of benign tracheobronchial stenosis. *Chest* 1998;114:796-800.
9. Ferretti G, Jouvan FB, Thony F, Pison C, Coulomb M. Benign noninflammatory bronchial stenosis: treatment with balloon dilation. *Radiology* 1995;196:831-4.
10. Shah H, Garbe L, Nussbaum E, Dumon JF, Chiodera PL, Cavaliere S. Benign tumors of the tracheobronchial tree: endoscopic characteristics and role of laser resection. *Chest* 1995;107:1744-51.
11. Turner JF Jr, Wang KP. Endobronchial laser therapy. *Clin Chest Med* 1999;20:107-22.
12. Mares DC, Broderick LS, Cummings OW, et al. Tracheobronchial amyloidosis: a review of clinical and radiographic characteristics, bronchoscopic diagnosis, and management. *J Bronchol* 1998;5:147-55.
13. Laforet EG, Berger RL, Vaughan CW. Carcinoma obstructing the trachea: treatment by laser resection. *N Engl J Med* 1976;294:941.
14. Rebeiz EE, Shapsay SM, Ingrams DR. Laser applications in the tracheobronchial tree. *Otolaryngol Clin North Am* 1996;29:987-1003.
15. Hanowell LH, Martin WR, Savelle JE, Foppiano LE. Complications of general anesthesia for Nd:YAG laser resection of endobronchial tumors. *Chest* 1991;99:72-6.
16. Cavaliere S, Foccoli P, Toninelli C, et al. Nd:YAG laser therapy in lung cancer: an 11-year experience with 2,253 applications in 1,585 patients. *J Bronchol* 1994;1:105-11.
17. Ramser ER, Beamis JF Jr. Laser bronchoscopy. *Clin Chest Med* 1995;16:415-26.
18. Denton RA, Dedhia HV, Abrons HL, Jain PR, Lapp NL, Teba L. Long-term survival after endobronchial fire during treatment of severe malignant airway obstruction with the Nd:YAG laser. *Chest* 1988;94:1086-8.
19. Miro AM, Shivaram U, Finch PJ. Noncardiogenic pulmonary edema following laser therapy of a tracheal neoplasm. *Chest* 1989;96:1430-1.
20. Dullye KK, Kaspar D, Ramsay MA, Giesecke AH. Laser treatment of endobronchial lesions. *Anesthesiology* 1997;86:1387-90.
21. Lang NP, Wait GM, Read RR. Cardio-cerebrovascular complications from Nd:YAG laser treatment of lung cancer. *Am J Surg* 1991;162:629-32.
22. Brutinel WM, Cortese DA, Edell ES, McDougall JC, Prakash UB. Complications of Nd:YAG laser therapy. *Chest* 1988;94:902-3.
23. Dumon JF, Shapshay S, Bourcereau J, et al. Principles for safety in application of neodymium-YAG laser in bronchology. *Chest* 1984;86:163-8.
24. Desai SJ, Mehta AC, VanderBrug Medendorp S, Golish JA, Ahmad M. Survival experience following Nd:YAG laser photoresection for primary bronchogenic carcinoma. *Chest* 1988;94:939-44.
25. Macha HN, Becker KO, Kemmer HP. Pattern of failure and survival in endobronchial laser resection: a matched pair study. *Chest* 1994;105:1668-72.
26. Stanopoulos IT, Beamis JF Jr, Martinez FJ, Vergos K, Shapshay SM. Laser bronchoscopy in respiratory failure from malignant airway obstruction. *Crit Care Med* 1993;21:386-91.
27. Colt HG, Harrell JH. Therapeutic rigid bronchoscopy allows level of care changes in patients with acute respiratory failure from central airways obstruction. *Chest* 1997;112:202-6.
28. Otto W, Szlenk Z, Paczkowski P, Gackowski W, Najnigier B, Karwowski A. Endoscopic laser treatment of benign tracheal stenoses. *Otolaryngol Head Neck Surg* 1995;113:211-4.
29. Brutinel WM, Cortese DA, McDougall JC, Gillio RG, Bergstrahl EJ. A two-year experience with the neodymium-YAG laser in endobronchial obstruction. *Chest* 1987;91:159-65.
30. Toty L, Personne C, Colchen A, Vourc'h G. Bronchoscopic management of tracheal lesions using the neodymium yttrium aluminum garnet laser. *Thorax* 1981;36:175-8.
31. Personne C, Colchen A, Leroy M, Vourc'h G, Toty L. Indications and technique for endoscopic laser resections in bronchology: a critical analysis based upon 2,284 resections. *J Thorac Cardiovasc Surg* 1986;91:710-5.
32. Homasson JP. Endobronchial electrocautery. *Semin Respir Crit Care Med* 1997;18:535-43.
33. Sheski FD, Mathur PN. Cryotherapy, electrocautery, and brachytherapy. *Clin Chest Med* 1999;20:123-38.
34. Marasso A, Gallo E, Massaglia GM, Onoscuri M, Bernardi V. Cryosurgery in bronchoscopic treatment of tracheobronchial stenosis: indications, limits, personal experience. *Chest* 1993;103:472-4.
35. van Boxem TJ, Venmans BJ, Schramel FM, et al. Radiographically occult lung cancer treated with fiberoptic bronchoscopic electrocautery: a pilot study of a simple and inexpensive technique. *Eur Respir J* 1998;11:169-72.
36. Mathur PN, Wolf KM, Busk ME, Briete WM, Datzman M. Fiberoptic bronchoscopic cryotherapy in the management of tracheobronchial obstruction. *Chest* 1996;110:718-23.
37. Maiwand MO, Homasson JP. Cryotherapy for tracheobronchial disorders. *Clin Chest Med* 1995;16:427-43. [Erratum, *Clin Chest Med* 1995;16:ix.]
38. Roden S, Homasson JP. Une nouvelle indication de la cryothérapie endobronchique: l'extraction de corps étrangers. *Presse Med* 1989;18:897.
39. Homasson JP. Bronchoscopic cryotherapy. *J Bronchol* 1995;2:145-9.
40. Villanueva AG, Lo TC, Beamis JF Jr. Endobronchial brachytherapy. *Clin Chest Med* 1995;16:445-54.
41. Mendiondo OA, Dillon M, Beach JL. Endobronchial irradiation in the treatment of recurrent non-oat cell bronchogenic carcinoma. *J Ky Med Assoc* 1983;81:287-90.
42. Seagren SL, Harrell JH, Horn RA. High dose rate intraluminal irradiation in recurrent endobronchial carcinoma. *Chest* 1985;88:810-4.

43. Sutedja G, Baris G, van Zandwijk N, Postmus PE. High-dose rate brachytherapy has a curative potential in patients with intraluminal squamous cell lung cancer. *Respiration* 1994;61:167-8.
44. Saito M, Yokoyama A, Kurita Y, Uematsu T, Miyao H, Fujimori K. Treatment of roentgenographically occult endobronchial carcinoma with external beam radiotherapy and intraluminal low dose rate brachytherapy. *Int J Radiat Oncol Biol Phys* 1996;34:1029-35.
45. Ofiara L, Roman T, Schwartzman K, Levy RD. Local determinants of response to endobronchial high-dose brachytherapy in bronchogenic carcinoma. *Chest* 1997;112:946-53.
46. Pisch J, Villamena PC, Harvey JC, Rosenblatt E, Mishra S, Beattie EJ. High dose-rate endobronchial irradiation in malignant airway obstruction. *Chest* 1993;104:721-5.
47. Hernandez P, Gursahaney A, Roman T, et al. High dose rate brachytherapy for the local control of endobronchial carcinoma following external irradiation. *Thorax* 1996;51:354-8.
48. Nori D, Allison R, Kaplan B, Samala E, Osian A, Karbowitz S. High dose-rate intraluminal irradiation in bronchogenic carcinoma: technique and results. *Chest* 1993;104:1006-11.
49. Kennedy AS, Sonett JR, Orens JB, King K. High dose rate brachytherapy to prevent recurrent benign hyperplasia in lung transplant bronchi: theoretical and clinical considerations. *J Heart Lung Transplant* 2000;19:155-9.
50. Miller JJ Jr, Phillips TW. Neodymium:YAG laser and brachytherapy in the management of inoperable bronchogenic carcinoma. *Ann Thorac Surg* 1990;50:190-6.
51. Shea JM, Allen RP, Tharratt RS, Chan AL, Siefkin AD. Survival of patients undergoing Nd:YAG laser therapy compared with Nd:YAG laser therapy and brachytherapy for malignant airway disease. *Chest* 1993;103:1028-31.
52. Khanavkar B, Stern P, Alberti W, Nakhosteen JA. Complications associated with brachytherapy alone or with laser in lung cancer. *Chest* 1991;99:1062-5.
53. Langendijk JA, Tjwa MK, de Jong JM, ten Velde GP, Wouters EF. Massive haemoptysis after radiotherapy in inoperable non-small cell lung carcinoma: is endobronchial brachytherapy really a risk factor? *Radiother Oncol* 1998;49:175-83.
54. Speiser B. Advantages of high dose rate remote afterloading systems: physics or biology. *Int J Radiat Oncol Biol Phys* 1991;20:1133-5.
55. Zajac AJ, Kohn ML, Heiser D, Peters JW. High-dose rate intraluminal brachytherapy in the treatment of endobronchial malignancy: work in progress. *Radiology* 1993;187:571-5.
56. Sutedja G, Baris G, Schaake-Koning C, van Zandwijk N. High dose rate brachytherapy in patients with local recurrences after radiotherapy of non-small cell lung cancer. *Int J Radiat Oncol Biol Phys* 1992;24:551-3.
57. Suh JH, Dass KK, Pagliaccio L, et al. Endobronchial radiation therapy with or without neodymium yttrium aluminum garnet laser resection for managing malignant airway obstruction. *Cancer* 1994;73:2583-8.
58. Cortese DA, Edell ES, Kinsey JH. Photodynamic therapy for early stage squamous cell carcinoma of the lung. *Mayo Clin Proc* 1997;72:595-602.
59. Balchum OJ, Doiron DR. Photoradiation therapy of endobronchial lung cancer: large obstructing tumors, nonobstructing tumors, and early-stage bronchial cancer lesions. *Clin Chest Med* 1985;6:255-75.
60. Edell ES, Cortese DA. Bronchoscopic phototherapy with hematoporphyrin derivative for treatment of localized bronchogenic carcinoma: a 5-year experience. *Mayo Clin Proc* 1987;62:8-14.
61. Gomer CJ, Dougherty TJ. Determination of [3H]- and [14C]hematoporphyrin derivative distribution in malignant and normal tissue. *Cancer Res* 1979;39:146-51.
62. Edell ES, Cortese DA. Photodynamic therapy: its use in the management of bronchogenic carcinoma. *Clin Chest Med* 1995;16:455-63.
63. Sutedja T, Baas P, Stewart F, van Zandwijk N. A pilot study of photodynamic therapy in patients with inoperable non-small cell lung cancer. *Eur J Cancer* 1992;28A:1370-3.
64. LoCicero J III, Metzendorf M, Almgren C. Photodynamic therapy in the palliation of late stage obstructing non-small cell lung cancer. *Chest* 1990;98:97-100.
65. McCaughan JS Jr. Survival after photodynamic therapy to non-pulmonary metastatic endobronchial tumors. *Lasers Surg Med* 1999;24:194-201.
66. Moghissi K, Dixon K, Stringer M, Freeman T, Thorpe A, Brown S. The place of bronchoscopic photodynamic therapy in advanced unresectable lung cancer: experience of 100 cases. *Eur J Cardiothorac Surg* 1999;15:1-6.
67. Taber S, Buschemeyer WC III, Fingar VH, Wieman TJ. The treatment of malignant endobronchial obstruction with laser ablation. *Surgery* 1999;126:730-3.
68. Mehta AC, Dasgupta A. Airway stents. *Clin Chest Med* 1999;20:139-51.
69. Becker HD. Stenting of the central airways. *J Bronchol* 1995;2:98-106.
70. Montgomery WW. T-tube tracheal stent. *Arch Otolaryngol* 1965;82:320-1.
71. Dumon JF. A dedicated tracheobronchial stent. *Chest* 1990;97:328-32.
72. Colt HG, Dumon JF. Airway stents: present and future. *Clin Chest Med* 1995;16:465-78.
73. Wassermann K, Koch A, Muller-Ehmsen J, Reuter M, Michel O, Eckel HE. Clinical and laboratory evaluation of a new thin-walled self-expanding tracheobronchial silicone stent: progress and pitfalls. *J Thorac Cardiovasc Surg* 1997;114:527-34.
74. Martinez-Ballarín JI, Diaz-Jimenez JP, Castro MJ, Moya JA. Silicone stents in the management of benign tracheobronchial stenoses: tolerance and early results in 63 patients. *Chest* 1996;109:626-9.
75. Diaz-Jimenez JP, Muñoz E, Ballarín J, Kovitz K, Pressas E. Silicone stents in the management of obstructive tracheobronchial lesions: 2 year experience. *J Bronchol* 1994;1:15-8.
76. Freitag L, Tekolf E, Stamatis G, Greschuchna D. Clinical evaluation of a new bifurcated dynamic airway stent: a 5-year experience with 135 patients. *Thorac Cardiovasc Surg* 1997;45:6-12.
77. Dasgupta A, Dolmatch BL, Abi-Saleh WJ, Mathur PN, Mehta AC. Self-expandable metallic airway stent insertion employing flexible bronchoscopy: preliminary results. *Chest* 1998;114:106-9.
78. Shapshay MS, Beamis JF Jr, Hybels RL, Bohigian RK. Endoscopic treatment of subglottic and tracheal stenosis by radial laser incision and dilation. *Ann Otol Rhinol Laryngol* 1987;96:661-4.
79. Dumon JF, Cavaliere S, Diaz-Jimenez JP, et al. Seven year experience with the Dumon prosthesis. *J Bronchol* 1996;3:6-10.
80. Carasco CH, Nesbitt JC, Charnsangavej C, et al. Management of tracheal and bronchial stenoses with the Gianturco stent. *Ann Thorac Surg* 1994;58:1012-7.
81. Hauck RW, Lembeck RM, Emslander HP, Schomig A. Implantation of Accuflex and Strecker stents in malignant bronchial stenoses by flexible bronchoscopy. *Chest* 1997;112:134-44.
82. Shaffer JP, Allen JN. The use of expandable metal stents to facilitate extubation in patients with large airway obstruction. *Chest* 1998;114:1378-82.
83. Zannini P, Melloni G, Chiesa G, Caretta A. Self-expanding stents in the treatment of tracheobronchial obstruction. *Chest* 1994;106:86-90.
84. Shapshay SM, Beamis JF Jr, Dumon JF. Total cervical tracheal stenosis: treatment by laser, dilation, and stenting. *Ann Otol Rhinol Laryngol* 1989;98:890-5.
85. Simpson GT, Strong MS, Healy GB, Shapshay SM, Vaughan CW. Predictive factors of success or failure in the endoscopic management of laryngeal and tracheal stenosis. *Ann Otol Rhinol Laryngol* 1982;91:384-8.
86. Rousseau H, Dahan M, Lauque D, et al. Self-expandable prostheses in the tracheobronchial tree. *Radiology* 1993;188:199-203.
87. Strausz J. Management of postintubation tracheal stenosis with stent implantation. *J Bronchol* 1997;4:294-6.
88. Sarodia BD, Dasgupta A, Mehta AC. Management of airway manifestations of relapsing polychondritis: case reports and review of literature. *Chest* 1999;116:1669-75.
89. Monnier P, Mudry A, Stanzel F, et al. The use of the covered Wallstent for the palliative treatment of inoperable tracheobronchial cancers: a prospective, multicenter study. *Chest* 1996;110:1161-8.
90. Bolliger CT, Heitz M, Hauser R, Probst R, Perruchoud AP. An airway Wallstent for the treatment of tracheobronchial malignancies. *Thorax* 1996;51:1127-9.
91. Nashef SA, Dromer C, Velly JF, Labrousse L, Couraud L. Expanding wire stents in benign tracheobronchial disease: indications and complications. *Ann Thorac Surg* 1992;54:937-40.
92. Mares DC, Mathur PN. Medical thoracoscopy: the pulmonologist's perspective. *Semin Respir Crit Care Med* 1997;18:603-15.
93. Smythe WR, Kaiser LR. History of thoracoscopic surgery. In: Kaiser LR, Daniel TM, eds. *Thoracoscopic surgery*. Boston: Little, Brown, 1993: 1-16.
94. Colt HG. Thoracoscopy: window to the pleural space. *Chest* 1999;116:1409-15.
95. Mathur PN, Astoul P, Boutin C. Medical thoracoscopy: technical details. *Clin Chest Med* 1995;16:479-86.
96. Mathur P, Loddenkemper R. Medical thoracoscopy: role in pleural and lung diseases. *Clin Chest Med* 1995;16:487-96.
97. Boutin C, Astoul P, Rey F, Mathur PN. Thoracoscopy in the diagnosis and treatment of spontaneous pneumothorax. *Clin Chest Med* 1995;16:497-503.
98. Boutin C, Viallat JR, Cargnino P, Farisse P, Choux R. La thoracoscopie en 1980: revue générale. *Poumon Coeur* 1981;37:11-9.
99. Viskum K, Enk B. Complications of thoracoscopy. *Poumon Coeur* 1981;37:25-8.
100. Inderbitzi RG, Grillet MP. Risk and hazards of video-thoracoscopic surgery: a collective review. *Eur J Cardiothorac Surg* 1996;10:483-9.

- 101.** Vining DJ, Liu K, Choplin RH, Haponik EF. Virtual bronchoscopy: relationships of virtual reality endobronchial simulations to actual bronchoscopic findings. *Chest* 1996;109:549-53.
- 102.** Rapp-Bernhardt U, Welte T, Budinger M, Bernhardt TM. Comparison of three-dimensional virtual endoscopy with bronchoscopy in patients with oesophageal carcinoma infiltrating the tracheobronchial tree. *Br J Radiol* 1998;71:1271-8.
- 103.** Suto Y, Tanabe Y. Evaluation of tracheal collapsibility in patients with tracheomalacia using dynamic MR imaging during coughing. *AJR Am J Roentgenol* 1998;171:393-4.
- 104.** Wanamaker JR, Eliachar I. An overview of treatment options for lower airway obstruction. *Otolaryngol Clin North Am* 1995;28:751-70.
- 105.** Kalweit G, Huwer H, Straub U, Gams E. Mediastinal compression syndromes due to idiopathic fibrosing mediastinitis — report of three cases and review of the literature. *Thorac Cardiovasc Surg* 1996;44:105-9.
- 106.** Worthington MG, Brink JG, Odell JA, et al. Surgical relief of acute airway obstruction due to primary tuberculosis. *Ann Thorac Surg* 1993;56:1054-62.
- 107.** Mendelson DS, Norton K, Cohen BA, Brown LK, Rabinowitz JG. Bronchial compression: an unusual manifestation of sarcoidosis. *J Comput Assist Tomogr* 1983;7:892-4.
- 108.** Masaoka A, Yamakawa Y, Niwa H, et al. Pediatric and adult tracheobronchomalacia. *Eur J Cardiothorac Surg* 1996;10:87-92.
- 109.** Jokinen K, Palva T, Sutinen S, Nuutinen J. Acquired tracheobronchomalacia. *Ann Clin Res* 1977;9:52-7.
- 110.** Susanto I, Peters JI, Levine SM, Sako EY, Anzueto A, Bryan CL. Use of balloon-expandable metallic stents in the management of bronchial stenosis and bronchomalacia after lung transplantation. *Chest* 1998;114:1330-5.
- 111.** Ferguson G, Benoist J. Nasal continuous positive airway pressure in the treatment of tracheobronchomalacia. *Am Rev Respir Dis* 1993;147:457-61.
- 112.** Adliff M, Ngato D, Keshavjee S, Brenaman S, Granton JT. Treatment of diffuse tracheomalacia secondary to relapsing polychondritis with continuous positive airway pressure. *Chest* 1997;112:1701-4.
- 113.** Anderson CB, Philpott GW, Ferguson TB. The treatment of malignant pleural effusions. *Cancer* 1974;33:916-22.
- 114.** Izbicki R, Weyhing BT III, Baker L, Caoili EM, Vaitkevicius VK. Pleural effusion in cancer patients: a prospective randomized study of pleural drainage with the addition of radioactive phosphorous to the pleural space vs. pleural drainage alone. *Cancer* 1975;36:1511-8.
- 115.** Marom EM, Patz EF Jr, Erasmus JJ, McAdams HP, Goodman PC, Herndon JE. Malignant pleural effusions: treatment with small-bore-catheter thoracostomy and talc pleurodesis. *Radiology* 1999;210:277-81.
- 116.** Zimmer PW, Hill M, Casey K, Harvey E, Low DE. Prospective randomized trial of talc slurry vs bleomycin in pleurodesis for symptomatic malignant pleural effusions. *Chest* 1997;112:430-4.
- 117.** Turler A, Gawenda M, Walter M. Palliative iodized talc pleurodesis with instillation via tube thoracostomy. *Support Care Cancer* 1997;5:61-3.
- 118.** Bloom AI, Wilson MW, Kerlan RK Jr, Gordon RL, LaBerge JM. Talc pleurodesis through small-bore percutaneous tubes. *Cardiovasc Intervent Radiol* 1999;22:433-6.
- 119.** Milanez RC, Vargas FS, Filomeno LB, et al. Intrapleural talc for the treatment of malignant pleural effusions secondary to breast cancer. *Cancer* 1995;75:2688-92.
- 120.** Sanchez-Armengol A, Rodriguez-Panadero F. Survival and talc pleurodesis in metastatic pleural carcinoma, revisited: report of 125 cases. *Chest* 1993;104:1482-5.
- 121.** Patz EF Jr. Malignant pleural effusions: recent advances and ambulatory sclerotherapy. *Chest* 1998;113:Suppl:74S-77S.
- 122.** Putnam JB Jr, Light RW, Rodriguez RM, et al. A randomized comparison of indwelling pleural catheter and doxycycline pleurodesis in the management of malignant pleural effusions. *Cancer* 1999;86:1992-9.
- 123.** Colt HG. Thoracoscopy: a prospective study of safety and outcome. *Chest* 1995;108:324-9.
- 124.** Harris RJ, Kavuru MS, Mehta AC, et al. The impact of thoracoscopy on the management of pleural disease. *Chest* 1995;107:845-52.
- 125.** Mares DC, Mathur PN. Medical thoracoscopic talc pleurodesis for chylothorax due to lymphoma: a case series. *Chest* 1998;114:731-5.
- 126.** Danby CA, Adebonojo SA, Moritz DM. Video-assisted talc pleurodesis for malignant pleural effusions utilizing local anesthesia and I.V. sedation. *Chest* 1998;113:739-42.
- 127.** Panebianco V, Calanducci F, Poli A, et al. Pleuroscopy and talc pleurodesis in recurrent pleural effusions: experience with 51 cases. *G Chir* 1995;16:437-41. (In Italian.)
- 128.** Gharagozloo FR, Wang KP. Thoracoscopy in pleural effusion. *J Bronchol* 1996;3:209-16.
- 129.** Aelony Y, King RR, Boutin C. Thoracoscopic talc poudrage in malignant pleural effusions: effective pleurodesis despite low pleural pH. *Chest* 1998;113:1007-12.

Copyright © 2001 Massachusetts Medical Society.