

SAFETY DURING BRONCHOSCOPY

Pages with reference to book, From 146 To 148

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Bronchoscopy can only be learnt through observing and performing and not by reading manuals. As in surgery, experience gained is directly related to the number of procedures performed. The procedure should not be performed unless there are valid indications and results or benefits outweigh the potential risks. Consideration should always be given to alternate methods for obtaining same information¹. However, if bronchoscopy appears to be the best way of getting same information, this is always indicated. Complications can arise with increased frequency when the procedure is earned out by the staff under training or gaining experience. In Pakistan, practice of pulmonology is at a primitive stage, facilities for fiberoptic bronchoscopy are available only at selected centres, even in teaching institutions and only limited experience has been gained. The procedure is likely to be introduced at more centres in near future. Present review is intended to provide information about the potential difficulties which may arise during the procedure.

HAZARDS OF BRONCHOSCOPY

In the long list of new departments added to pulmonology, bronchoscopy has been introduced at almost every teaching centre in developed countries. While rigid bronchoscope remained in the domain of surgeons, physicians developed intense desire for early learning and routine application of fiberoptic bronchoscope², since its introduction in late sixties. At present, in pulmonary practice, bronchoscopy is the 3rd most common invasive procedure after arterial blood gases and thoracentesis³. Whether bronchoscopy is considered a procedure⁴ or a science⁵, it needs to be performed with an art. Availability of fibroscope to many countries around the world has resulted in an epidemic. Based on experience and sound clinical judgement, the procedure should be carried out with firm indications, whether useful information will emerge. The procedure may be employed in a variety of situations for both diagnostic as well as therapeutic purposes⁶. These include evaluation of focal and diffuse lung diseases, therapeutically for atelectasia⁷ and foreign bodies for diagnosis of unexplained pleural effusion⁸ and hemoptysis⁹, diagnosis and staging of neoplasm¹⁰, assessment of localised infection, bronchiectasis and slow resolving pneumonia¹¹, bronchial lavage in immunocompromised or sputum negative patients¹², laser therapy¹³, diagnostic biopsy procedures¹⁴, evaluation of thoracic trauma¹⁵, bronchography and in case of difficult intubation.

COMPLICATIONS

In experienced hands it is a safe procedure, however, complications¹⁶ do occur, attributable to the procedure, lack of skill and failure to adequately monitor the patients. A large proportion of patients undergoing procedure are highly susceptible and carry risk factors for the development of complications like smoking¹⁷. Complications can arise during premedication and preparation of the patient, technical problems may occur during insertion of the scope and often complications are associated with the procedure itself, biopsy, bronchial lavage and inadequate post-procedure monitoring. A major complication is one which is considered to endanger life or require urgent therapeutic intervention.

PREMEDICATION

Sedation with diazepam or morphine can lead to respiratory depression and acute respiratory failure, hypotension, syncope or hyperexitement. Among patients with COPD, fall in FEV₁ by 10% and rise in airway resistance by 30% was observed 15 minutes after the procedure¹⁸. Sedation may have an additive adverse effect. Almost 50% of all major complications during the procedure are related to sedatives and local anaesthetics. With adequate anaesthesia there may be no need of premedication¹⁹, particularly in the presence of obstructive airway disease. Asthma and COPD patients may be given hydrocortisones, atropine and nebulised salbutamol as premedication and lavage fluid should be limited to 200-300 ml at body temperature.

LOCAL ANAESTHESIA

Lignocain is used in form of 2-4% spray and 2% gel for local anaesthesia. Inhaled lignocain can anaesthetise upto midtrachea. With maximum recommended dose of 300-400 mg²⁰ a blood level of 30-50% may be achieved through mucosal absorption²¹. In a normal person complications with lignocain are rare as much of this is aspirated out. Patients with bronchial hyper-reactivity may get severe bronchospasm. This response is not related to airway; histamine-responsiveness or to the preservatives in the lignocain preparation²². Other adverse effects include hypersensitivity reaction and cardiorespiratory arrest. Potentially lignocain can interfere with the growth of aerobes anaerobes, fungi, mycobacteria²³ and impair the function of pulmonary alveolar macrophages and other immune competent cells, however, the concentration of aspirated lignocain has been found to be too low²⁴. Rapid absorption of lignocain from inflamed mucosa²⁵ can result in toxic level (>6ug/ml). Signs of toxicity are directly related to the free plasma fraction of lignocain, this in turn is inversely proportional to the level of plasma alpha₁-acid glycoprotein (AAG) which can rise in case of malignancy or thoracic trauma. Higher level of lignocain is tolerable if no rise in free form is achieved.

INSERTION OF BRONCHOSCOPE

Amount of pressure required to insert a bronchoscope is a matter of experience and learning. As asymmetry of nasal passages is not uncommon, gentle attempt should be made to find more patent side. Saline lavage of nasal passage, topical vasoconstriction and rotatory movements of scope may help facilitate effective and safe passage. In case of inadequate anaesthesia laryngial and broncho-spasm can occur, resulting in hypoxia which may lead to seizures or cardiac arrhythmias. On an average there is fall in P_{O₂} by 20 mmHg, particularly ml at body temperature. during suction and rise in PCO₂; Fall P_{O₂} may persist for many hours and supplemental O₂ is recommended as a routine¹⁶ during the procedure, particularly for high risk patients²⁷.

MONITORING DURING PROCEDURE

During the procedure monitoring of the patient is required by an assistant, as bronchoscopist is often so engrossed in the endoscopic findings that physiological condition of the patient is forgotten. As patient is not under general anaesthesia, minimum conversation about the findings during the procedure will lessen patient's anxiety with minimal complications.

BIOPSY PROCEDURES

Incidence of complications is increased 20 times when biopsy procedures are carried out. Biopsies are only helpful if proper indications are sought for and proper information is collected through clinical and physical examination²⁸. Transbronchial needle aspiration is safe in experienced hands and only minor complications are seen²⁹. Complications of bronchial biopsy include pneumothorax³⁰ and haemorrhage. Intubation is needed in 40-50% of pneumothoraces. Small leak can occur after a delay, this does not need a routine post-procedure chest radiograph³¹. Risk of haemorrhage is minimised by having coagulation profile checked. Incidence of serious bleeding (>50 ml) is increased among

immuno-compromised and uraemic patients. Other complications include, vasovagal episodes, serious arrhythmias, myocardial infarction and pulmonary oedema. Fever and pulmonary infiltrate has also been noted following the procedure. During bronchoalveolar lavage if tip of the bronchoscope is not wedged into an airway much of the instilled saline may be distributed to the adjacent or proximal airways. This may disseminate infection and even result in respiratory compromise.

FATALITY

Mortality rate as low as 0.01-0.04% has been reported. Higher mortality and morbidity has been related to the procedure being carried out by the staff under training. In two series by Suratt et al³² (48,000 procedures) and Simpson et al³³ (40,000 procedures reviewed) causes of death have included reaction to topical anaesthesia, massive haemorrhage from bleeding tumour, myocardial infarction, chronic respiratory failure, severe pneumonia, bronchospasm and general anaesthesia. During another series by Herf et al³⁴, in 2,628 patients mortality rate of 0.5% was observed. The procedure may be best avoided among high risk patients, having severe hypoxia, serious arrhythmias, unstable angina, recent myocardial infarction and poor cooperation. Biopsy procedures are contraindicated in bleeding diathesis, uraemia, pulmonary hypertension, severe anaemia and in patients on mechanical ventilation or positive pressure breathing.

CONCLUSION

Fibreoptic bronchoscopy is a safe procedure even for a beginner³⁵, provided a little extra care is applied and procedure is carried out properly and rapidly in well selected patients, however, safety is no indication for routine use³⁶. It is an aggressive bronchoscopist who is more likely to encounter complications than a conservatist.

REFERENCES

1. Grant, I.W. Hazards of bronchoacopy (letter). *Br. Med.J. (Clin. Res.)*, 1986; 293:955.
2. Kvale, P.A. Training in laserbronchoscopy and proposal for credentialing. *Chest*, 1990; 97:983-89.
3. Parakash, U.B. Bronchoacopy. *Chest*, 1989; 95:259.
4. Stangel, P. Bronchoacopy, enough or not enough. *Chest*, 1990;9&774.
5. Sen, R.P. and Walab,T.E. Bronchoscopy, enough or too much. *Cheat*, 1989;96:710-11.
6. George, La, Light, LW. and Mathay, LA. ed. *Chest medicine*. New York, Churchill Livingatone, 1983,pp.219-21.
7. Jawarski, A., Goldberg. S.K., Walkenstein, M.D.. Wilson, E and Lippavann, M.L Use of immediate postlobectomy tibreoptic bronchoscopy in preventing atelactsa. *Cheat*, 1988;94:38-43.
8. Heaton, LW. and Roberta, C.M. The role of fibreoptic bronchoscopy in the inveatigation of pleural effusion. *Poatgrad. Med.)*. 1988;64:581-82
9. Lederle, F.A., Nichol, KU and Parenti, CM. Bronchoscopy to evaluate hemoptysis in older menwith nonapacificchest roentgenogram. *Chest*, 1989;95:1043-47.
10. Mod, K, Yanase, N., Kaneco, M. et al Diagnoais of peripheral lung cancer in cases of tumour 2cm orless in size. *Chest*, 1989;95:304-8.
11. Peinsilver, S.H., Fein, A.M., Neiderman, M.S.,Shultz, DR and Faegenburg, D.H. Utility oftibreopticbronchoscopy in nonreaolving pneumonia. *Cheat*, 1990;98:1322-26.
12. Jaiawal, A.K., Kalopati, D.D., Jam, N.K. and Singh, M.M. Role of bronchoscopy in the early diagnosis of suspected smear negative caaes of pulmonary tuberculosis, *Indian.J. Tuberc.*, 198936:231.
13. Geracin, V.A., Levashov, Y.N., Shaflovsey, B.B. et al. Bronchoscopic laser photocoagulation of superficial carcinomaof bronchi. *Cheat*, 1990;98:238
14. Shure, D. Transbronchial biopsyand needle aspiration. *Chest*, 1989;95:1130-38.

15. Hara, K.S. and Parakaah, U.B. Fibreoptic bronchoscopy in the evaluation of acute chest and upper airway trauma. *Chest*, 1989;96:627-30.
16. Sen, R.P. and Walsh, T.E., Serious complications of fibreoptic bronchoacopy. *Chest*, 1988;94:222.
17. Metzger, U.P. and Altoae, M.D. Effect of fibreoptic bronchoscopy on respiratory performance in patients with chronic airway obstruction. *Thorax*, 1975;30:441.
18. Nolan, J.J. Fibreoptic bronchoacopy. *N. Engl. J. Med.*, 1984;311:1443.
19. Colt, H.G. and Moris, J.F. Fibreoptic bronchoacopy without premedication; a retrospective study. *Chest*, 1990;98:1327-30.
20. Berger, It, McConnel, J.W., Philips, B. and Overman, T.L. Safety and efficacy of using high dose topical and nebulised anaesthesia to obtain endotracheal culture. *Chest*, 1989;95:299-303.
21. Adriani, Land Campbell, D. Fatalities following topical application of local anaesthetic to mucous membrane. *JAMA.*, 1956;162:1527-30.
22. McAlpine, L.G. and Thomson, N.C. Lidocain induced bronchoconstriction in asthmatic patients. *Chest*, 1989;96: 1012- 15.
23. Keto, H., Goto, H., Yuaaa, K, Leki, It and Schimada, K Lidocain concentration in endotracheal aspirated during fibreoptic bronchoacopy. *Chest*, 1989;96:700.
24. Kirkpatrick, M.B. Lidocain topical anaesthesia for flexible bronchoscopy. *Chest*, 1989;96:965-66.
25. Ameer, B., Burlingame, MB. and Harmane, EM. Rapid mucosal absorption of topical lidocain during bronchoacopy in the presence of oral candidiasis. *Chest*, 1989;96:1438-39.
26. Trovillet, J.L., Guigvet, M., Gilbert, C. et al. Fibreoptic bronchoacopy in ventilated patients. *Chest*, 1990;97:927-33.
27. Albertini, R.E., Harrell, J.H, Kurihara, N. and Moser, KM. Arterial hypoxemia induced by fibreoptic bronchoscopy. *JAMA.*, 1974;230:1666-67.
28. Ruhmedder, J.J. Enticement for fruitless bronchoacopy. *Chest*, 1989;96:708-10.
29. Harrow, EM., Oldenberg. F.A., Lingerfeltera, MS. and Smith, AM. Tranabronchial needle aspiration in clinical practice. *Chest*, 1989;96:1270.
30. Pepeira, W., Kovnat, D.M. and Snider, G.L. A prospective cooperative study of complications following fibreoptic bronchoacopy. *Chest*, 1978;73:813-16.
31. Milman, M.G., Evina, A.E. and Sahn, SA. Immediate chest roentgenography following fibreoptic bronchoacopy. *Chest*, 1989;96:477-79.
32. Suratt, P.M., Smiddy, J.F. and Gruber, B. Deaths and complications associated with fibreoptic bronchoscopy. *Chest*, 1976;69:746-51.
33. Simpson, KG., Arnold, A.G., Pruvia, A. et al. Postal survey of bronchoacopic practice by physicians in the United Kingdom. *Thorax*, 1986;41:311.
34. Hen, S.M. and Suratt, P.M. Complications of transbronchial lung biopsies. *Chest*, 1978;73:759.
35. Ticklu, B. Flexible fibreoptic bronchoscopy in Addis Ababa: a 4 years experience. *Br.J.Dis. Chest*, 1986;80:283.
36. Olopade, C.O. and Parskash, U.B. Bronchoscopy in the critical care unit. *Mayo Clin. Proc.*, 1989;64:1255-63.