- ³ Farrow, S. C., Fisher, D. J., and Johnson, D. B., *British Medical Journal*, 1972, 3, 686.
- ⁴ *British Medical Journal*, 1974, 3, 703.
- ⁵ Nelson, S. D., and Tovey, G. H., *British Medical Journal*, 1974, 1, 622.
- ⁶ Martin, D. C., Smith, G. A., and Fareed, D. O., *Surgical Forum*, 1969, 20, 529.
- ⁷ Belzer, F. O., *Transplantation Proceedings*, 1971, 3, 268.
- ⁸ Frost, A. B., *et al.*, *Lancet*, 1970, 1, 620.
- ⁹ Scott, D. F., *et al.*, *Transplantation*, 1971, 11, 90.
- ¹⁰ Festenstein, H., *et al.*, *Lancet*, 1971, 2, 225.
- ¹¹ Ackermann, J. R., and Snell, M. E., *British Journal of Urology*, 1968, 40, 515.
- ¹² Calne, R. Y., *British Medical Journal*, 1969, 2, 565.
- ¹³ Gelin, L. E. *et al.*, *Scandinavian Journal of Urology and Nephrology*, 1968, 2, 1.
- ¹⁴ Collins, G. M., Bravo-Shugerman, M. and Terasaki, P. I., *Lancet*, 1969, 2, 1219.
- ¹⁵ Bell, P. F. R., *et al.*, *British Medical Journal*, 1972, 4, 408.
- ¹⁶ Hall, C. L., *et al.*, *Lancet*, 1974, 1, 532.
- ¹⁷ Belzer, F. O., Ashby, B. S., and Dunphy, J. E., *Lancet*, 1967, 2, 536.
- ¹⁸ Hartley, L. C. J., Collins, G. M., and Clunie, G. J. A., *New England Journal of Medicine*, 1971, 285, 1049.
- ¹⁹ Tanaka, N., Stevens, L. E., and Terasaki, P. I., *Transplantation*, 1971, 12, 348.
- ²⁰ Collaborative Study, *Lancet*, 1972, 2, 887.

Criteria of Fitness for Anaesthesia in Patients with Chronic Obstructive Lung Disease

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British Medical Journal, 1975, 3, 670-673

Summary

Twelve patients with severe chronic obstructive lung disease undergoing 15 operations were assessed with preoperative lung function tests and blood gas estimations. Their operative and postoperative course was followed. There were no deaths or serious complications. Patients fell into three groups: those with low respiratory capacity but normal blood gases, who required no special respiratory treatment apart from physiotherapy and antibiotics; those with hypoxaemia but normal arterial carbon dioxide pressure, who needed more prolonged oxygen treatment after operation; and those with hypoxaemia and hypercapnia, who needed postoperative ventilatory support.

While forced expiratory volume in one second (FEV₁) is a good screening test in preoperative assessment it should be supplemented by arterial blood gas estimations in patients with an FEV₁ of less than 1 litre.

Introduction

Doctors are often confronted with the problem of whether to recommend or accept for surgery a patient with chronic obstructive lung disease. The decision rests partly on surgical considerations but also on the likely effect of anaesthesia and surgery on the respiratory state of the patient. This is difficult to predict, and numerous reviews have described techniques for preoperative assessment of respiratory function but given no guidance on interpretation of findings in relation to the expected course of events.

The commonest routine preoperative test of pulmonary function is the measurement of forced vital capacity, with the results expressed as the forced expiratory volume in one second (FEV₁) and also as a percentage of the slow vital capacity (VC). The theoretical hazards of impaired ventilatory capacity in relation to anaesthesia are widely appreciated, but there is no reference to the actual level of reduction of FEV₁

which in itself constitutes a definite hazard or contraindication to anaesthesia and surgery. Diamant and Palmer,¹ however, claim that a reduction in FEV₁ below 70% indicates that a patient is at risk of developing postoperative pulmonary complications, but this conclusion was not supported by any study of their patients' progress after operation.

Published works gives little or no guidance on the potential hazard in patients with an exceptionally low FEV₁. The lowest value reported is 0.81 litre.² Thornton³ lists several patients with values below 1 litre who underwent surgery but gives no details of progress during or after surgery. We therefore collected details of 15 anaesthetics in 12 patients with grossly reduced FEV₁ (range 0.45-1.04 litres) to determine whether a reduction of FEV₁ to these low levels in itself introduces a hazard to anaesthesia for routine surgery. In most cases we also noted the preoperative blood gas levels, since Stein *et al.*² reported that a raised alveolar carbon dioxide pressure was an important factor discriminating those patients who would develop postoperative pulmonary complications.

Patients and Management

Fifteen operations were performed on 12 patients with chronic obstructive airway disease who presented consecutively for surgery with an FEV₁ of 1 litre or less. The anaesthetic management was entirely at the discretion of the seven different anaesthetists on whose lists the patients came for surgery.

Preoperative Assessment.—Patients were subjected to routine preoperative clinical assessment by the anaesthetist. Tests of ventilatory capacity were also carried out in all patients. FEV₁ and VC were measured with a Vitalograph dry spirometer, usually one or two days before operation. Predicted normal values were taken from the data of Cotes.⁴ Arterial gases were measured preoperatively in 10 patients using a Corning Eel model 165, oxygen pressure (PaO₂) calibration being carried out with 30% glycerol in water equilibrated with air. The normal range of blood gases was taken from the data of Raine and Bishop.⁵

Preoperative Management.—All patients received preoperative pulmonary physiotherapy except one patient (case 12) who was admitted as an emergency and four (cases 2, 7, 9, and 11) for whom it was deemed unnecessary. All patients received antibiotics before and after surgery. Details of premedication are given in table I.

Anaesthesia.—Induction of anaesthesia was with thiopentone in all cases. Spontaneous respiration and anaesthesia with halothane and nitrous oxide without endotracheal intubation was used four times for relatively minor operations not requiring relaxation. For the remaining 11 operations, patients were intubated and ventilated artificially with a Manley Ventilator incorporated in a Blease "Northwick Park" anaesthetic apparatus. Anaesthesia was maintained with nitrous oxide supplemented with halothane, fentanyl, or diamorphine. The relaxant pancuronium was used in nine cases. In six cases

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TABLE 1—Operations Performed and Anaesthetics Used in 12 Patients with Chronic Obstructive Lung Diseases

Case No.	Premedication (mg)	Operation	Induction (mg)	Maintenance	Airway	Respiration	Length of Operation (min)	Postoperative Regional Analgesia
1	Morph. (2.5), Atrop. (0.6), Drop. (2.5), Hydc. (100)	Cystoscopy, bilateral hernia	Thiop. (300)	N ₂ O/Hal	Mask	Spontaneous	90	Epidural
2 a	Papa. (10), Hyos. (0.2), Hydc. (100)	Sigmoid colectomy	Thiop. (100)	N ₂ O/Hal, Panc (6 mg)	E.T. tube	Artificial	185	
2 b	Papa. (10), Hyosc. (0.2)	Closure colostomy	Thiop. (100)	N ₂ O/Hal, Panc (4 mg)	E.T. tube	Artificial	45	
3 a	Morph. (5), Drop. (5), Atrop. (0.6)	Cystoscopy	Thiop. (150)	N ₂ O/Hal	Mask	Spontaneous	25	
3 b	Papa. (10), Hyosc. (0.2)	Cystectomy	Thiop. (250)	N ₂ O/Hal, Panc (8 mg)	E.T. tube	Artificial	180	
4	Hydc. (100)	Transvesical prostatectomy	Thiop. (50)	N ₂ O/Hal	E.T. tube	Artificial	80	Epidural
5	Papa. (15), Hyosc. (0.3)	Cryodiatheirmy to eye	Thiop. (50)	N ₂ O, fentanyl, Panc (11 mg)	E.T. tube	Artificial	180	
6	Papa. (10), Hyosc. (0.2)	Transurethral prostatectomy	Thiop. (250)	N ₂ O, Panc (8 mg)	E.T. tube	Artificial	45	Epidural
7	Papa. (10), Hyosc. (0.2)	Repair of incisional hernia	Thiop. (100)	N ₂ O/Hal, Panc (4 mg)	E.T. tube	Artificial	150	Epidural
8	Morph. (7.5), Drop. (5), Atrop. (0.6)	Cholecystectomy	Thiop. (250)	N ₂ O, diamorph. (4 mg), Panc (6 mg)	E.T. tube	Artificial	130	Intercostal block (T 6-12, bupivacaine)
9	Atrop. (0.6)	Cystoscopy, prostatectomy, repair hernia	Thiop. (225)	N ₂ O/Hal, fentanyl	E.T. tube	Artificial	75	Epidural
10 a	None	Removal of pacemaker	Thiop. (100)	N ₂ O/Hal	Mask	Spontaneous	60	
10 b	None	Insertion of pacemaker	Thiop. (100)	N ₂ O/Hal	Mask	Spontaneous	60	
11	Papa. (10), Atrop. (0.6)	Abdominoperineal resection of rectum	Thiop. (250)	N ₂ O/Hal, Panc (6 mg)	E.T. tube	Artificial	135	
12	None	Suture of bleeding ulcer and vagotomy	Thiop. (250)	N ₂ O, Panc (6 mg)	E.T. tube	Artificial	120	

Morph. = Morphine. Atrop. = Atropine. Drop. = Droperidol. Hydc. = Hydrocortisone. Papa. = Papaveretum. Hyosc. = Hyoscine. Thio. = Thiopentone. Hal. = Halothane. Panc. = Pancuronium. E.T. = Endotracheal.

regional analgesia was instituted for control of postoperative pain in accord with current practice at this hospital (table I).

Operations.—The operations were predominantly abdominal and urogenital (table I). The series included no thoractomies or neurosurgical procedures but there was one operation for detachment of the retina.

Results

Preoperative Assessment.—All patients were dyspnoeic on exertion. One patient (case 1) gave a history of late onset asthma and two (cases 7 and 11) had unremitting dyspnoea without cough or sputum. The remaining nine patients had a history of chronic cough and expectoration. The FEV₁ ranged from 0.45-1.04 litres (body temperature and pressure, saturated with water) with a mean value of 0.78, or 40% of predicted (table II). The mean FEV₁:VC ratio was 43%. Arterial carbon dioxide pressure (Paco₂) was 6.0 kPa (45 mm Hg) or above in four patients, though two of these had FEV₁ values above 0.85 litres. Pao₂ was below the normal range for patients' ages in all four patients with raised Paco₂ and in another three patients with normal or reduced Paco₂.

Anaesthesia.—No difficulties arose in the course of any anaesthetic. On the four occasions when patients breathed spontaneously they seemed to maintain satisfactory levels of ventilation as judged by

observation of the reservoir bag and did not become cyanosed when breathing 35% oxygen. The other patients were ventilated artificially without difficulty and air trapping was not clinically apparent.

POSTOPERATIVE COURSE

Table III summarizes the postoperative course and blood gas results. The time spent in the recovery room or intensive therapy unit was a measure of the opinion of the anaesthetist on the patient's recovery from the operation and anaesthetic. In nine cases the patient returned to the ward within four hours, after a stay which was within normal limits for the type of surgery undertaken. None of these patients gave any indication of difficulties referable to the respiratory system though two patients (cases 6 and 9) had a Paco₂ of 6.0 and 6.6 kPa (45.0 and 49.2 mm Hg) respectively, and two (cases 9 and 10) had FEV₁ values of 0.54 litres and 0.45 litres respectively before operation. Early postoperative blood gas values in case 9 indicated a slight increase in Paco₂ from 6.6 to 7.2 kPa (49.2 to 54 mm Hg). Postoperative blood gases showed only usual changes in cases 2 and 7.

Three patients (cases 7, 10, and 11) were admitted to the intensive therapy unit overnight for observation. All three recovered normally from anaesthesia and received no ventilatory support.

Postoperative difficulties were encountered in three patients. One (case 8) had arterial hypoxaemia Pao₂ 7.2 kPa (53.9 mm Hg) before operation, which was accentuated after the operation. Two hours after operation his Pao₂ was 12 kPa (90 mm Hg) while breathing 50% oxygen and he was gradually weaned from oxygen-enriched gas mixtures over the next five days, by which time he had a Pao₂ of 5.9 kPa (44 mm Hg) when breathing air. Paco₂ rose to 6.7 kPa (50 mm Hg) during the first few hours after operation but was within the normal range the next day. Ventilatory support was not required and he received no respiratory treatment other than oxygen and physiotherapy. There were no significant changes in four postoperative chest x-ray films other than slight opacity at the left costophrenic angle. The intercostal block provided 12 hours of analgesia and no postoperative opiates were required. He left hospital 13 days after his cholecystectomy.

One patient (case 5) who had preoperative hypercapnia was unable to breathe satisfactorily after operation and was ventilated artificially for 24 hours. On return of spontaneous respiration Paco₂ ranged from 7.5 to 8.4 kPa (56-63 mm Hg) during the first four hours, but thereafter returned to its preoperative level. Oxygenation of arterial blood was impaired for 48 hours (Pao₂ 5.7 kPa, 43 mm Hg) while breathing air

TABLE II—Pulmonary Function Test Results. Oxygen Pressure Values that Fell below 2 S.D. of Predicted are Shown in *Italic*

Case No.	Age and Sex	VC (Litres)	FEV ₁ in Litres (% of Predicted)	FEV ₁ as % of VC	Blood Gases		
					Pao ₂ (kPa)	Paco ₂ (kPa)	PH
1	56 M.	3.19	0.88 (37)	28	10.7	5.6	7.380
2	57 M.	1.75	0.82 (30)	47	10.6	5.1	7.371
3	76 M.	2.03	1.04 (45)	51			
4	77 M.	2.30	0.86 (41)	37	10.5	5.6	7.347
5	70 M.	2.09	1.00 (34)	48	6.4	6.8	7.368
6	69 M.	1.85	0.88 (34)	48	8.8	6.0	7.380
7	73 F.	1.21	0.77 (49)	64	7.8	4.4	7.428
8	69 M.	2.10	0.80 (27)	38	7.2	5.0	7.470
9	64 M.	1.74	0.54 (48)	31	8.5	6.6	7.310
10	65 F.	1.32	0.45 (24)	34	7.6	5.8	7.390
11	88 F.	1.30	0.76 (77)	59			
12	70 M.	1.43	0.50 (35)	35	5.9	7.6	7.320

Conversion: SI to Traditional Units—Blood gases: 1 kPa ≈ 7.5 mm Hg.

TABLE III—Postoperative Course in 12 Patients. Blood Gases were Estimated on Day of Operation, usually Two to Four Hours after Operation

Case No.	Time in Recovery Ward or I.T.U. (h)	Ventilatory Support	Oxygen Enrichment (% O ₂)	Blood Gases		Length of Physiotherapy (d)	Length of Stay in Hospital (d)
				Pao ₂ (kPa)	Paco ₂ (kPa)		
1	1.5	None	28	11.7	6.4	10	10
2 a	0.75	None				None	18
2 b	3.0	None				None	6
3 a	0.2	None				4	5
3 b	2.25	None	29	6.3	5.6*	11	19
4	1.3	None				14	16
5	4 d	24 h				11	>11†
6	2	None				6	9
7	18	None	40	22.0	4.9	10	11
8	5 d	None	28	7.5	6.3	9	13
9	3.75	None	28	11.1	7.2	None	6
10 a	2 d	None	21 35	5.9 9.6	7.6 6.0*	2	15 49
10 b	0.5	None				None	
11	18	Nonc				4	
12	10 d	5 d					

*On intermittent positive pressure ventilation.

†Transferred to another hospital.

24 hours after operation), and oxygen enrichment (24-33%) was used for 48 hours. Physiotherapy was continued for 11 days, after which he was transferred to an ophthalmic unit. Postoperative chest x-ray examination showed a band shadow at the left base which could have been due to a pulmonary infarct, but there was no other evidence for this and the shadow resolved.

The third patient requiring postoperative respiratory treatment (case 12) was admitted as an emergency with upper gastrointestinal bleeding. Postoperative artificial ventilation was clearly indicated on clinical grounds and continued uneventfully for five days. On return to spontaneous breathing he maintained a Paco₂ in the range 7.2-7.9 kPa (54-59 mm Hg) and required oxygen-enrichment of inspired gas (Pao₂ 5.3 kPa (40 mm Hg) breathing air). He left the intensive therapy unit after 10 days with a Paco₂ of 8.1 kPa (61.0 mm Hg) and a Pao₂ of 7.7 kPa (57.7 mm Hg) breathing air. Physiotherapy was continued for 21 days.

Discussion

From this survey we may distinguish three groups of patients who present for surgery with chronic obstructive airway disease. Firstly, those patients with low FEV₁ but normal Paco₂ and without overt arterial hypoxaemia (Pao₂ > 7.3 kPa (55 mm Hg)). This group included cases 1, 2, 4, 7, 10, and probably 3 and 11, in whom arterial blood gases were not measured because this did not seem necessary. No patient in this group gave the least cause for anxiety even though preoperative values for FEV₁ were grossly reduced. Routine anaesthetic techniques and normal postoperative attention gave entirely satisfactory results.

The second group consists of patients with low FEV₁ and low Pao₂ (< 7.3 kPa) but normal Paco₂. Only one of our patients (case 8) fell into this group. The only special treatment was the prolonged use of oxygen during the postoperative period, when the inspired oxygen concentration was titrated against his Pao₂ to give a safe value. He was treated in the intensive therapy unit more for convenience than for medical necessity.

The third group consists of patients with hypercapnia (Paco₂ > 5.9 kPa (> 44 mm Hg)), greatly reduced FEV₁, and Pao₂ more than 2 S.D. below the normal range. This group included four of our patients (cases 5, 6, 9, and 12), overt hypoxaemia (Pao₂ < 7.3 kPa) was present in cases 5 and 12. Two patients (cases 6 and 9) had only moderate hypercapnia (Paco₂ 6.0-6.6 kPa (45.0-49.2 mm Hg)) and made uneventful recoveries from prostatectomy with no special respiratory treatment. The two patients with hypoxaemia and Paco₂ in the range 6.8-7.6 kPa (50.9-57.3 mm Hg) (cases 5 and 12), however, needed full ventilatory support in the postoperative period.

We attach considerable importance to the fact that none of our patients had appreciable secretions at the time of operation. This was due partly to preoperative physiotherapy and antibiotics and partly to the changing pattern of chronic bronchitis which over the past 10 years has shown far less of the copious purulent secretions that presented such difficulties to the anaesthetist in former times. Howard⁶ has attributed the less rapid

progression of the disease and the diminished role of infection to the clean air policy and, certainly, we are now seeing a different type of patient from those seen and anaesthetized immediately after the war.

There are few studies of the predictive value of pulmonary function tests in patients with obstructive airways disease undergoing anaesthesia and surgery. Stein *et al.*² reported a series of 63 randomly selected patients, 30 of whom were classified as abnormal on lung function testing. Respiratory complications were almost exclusively confined to this group. The best predictive test was found to be maximum expiratory flow rate, which correlates well with FEV₁. These authors also found that a raised Paco₂ was associated with serious respiratory complications in every case. Appleberg *et al.*,¹ in a similar study, found that simple lung function tests increased the ability to predict those patients likely to have postoperative respiratory problems. Lockwood⁸ followed up 81 patients after cardiothoracic operations and found that both respiratory and cardiac complications were more common among patients with abnormal function test results. His results suggested that measurements of lung volumes, especially residual volume, might provide a sensitive predictive test of postoperative respiratory complications. Anderson and Ghia⁹ also found a good correlation between tests of airways resistance and postoperative course in cardiothoracic patients, including open heart cases.

In none of these studies are there any recommendations on the lower limit of values for acceptance for surgery. We do not believe that a reduced FEV₁ without hypercapnia or hypoxia is a barrier to routine surgery. Values below 1 litre or 50% of predicted clearly demand that the patient should be carefully monitored after operation but are compatible with normal recovery from surgery. Admission to the intensive therapy unit may be advisable overnight after surgery.

In patients with low FEV₁ and arterial hypoxaemia but no hypercapnia there should also be no barrier to routine surgery but Pao₂ must be carefully monitored and oxygen treatment provided to correct any tendency to deterioration of Pao₂ after the operation. This may be more convenient in an intensive therapy unit if that is where blood-gas measurement is carried out. The most important aspect of preoperative respiratory assessment is a raised Paco₂. A slight increase may offer no appreciable difficulty but an increase above 6.7 kPa (50 mm Hg) often indicates the need for postoperative artificial ventilation requiring prolonged endotracheal intubation or, rarely, tracheostomy.

Hence, the most important preoperative test is measurement of the arterial blood gases, because this shows the most significant abnormalities. Measurement of FEV₁ (or peak expiratory flow) is convenient and is likely to show an abnormality in any patient with chronic obstructive airway disease but it fails to distinguish those patients who are likely to require ventilatory support after operation. It therefore has a role as a preliminary screen but should be supplemented by measurement of arterial blood gases in patients with low values. We regard 1 litre or 50% of predicted

as the lowest values for FEV₁ that should be accepted for surgery without measurement of arterial blood gases.

Patients thus selected will benefit from preoperative physiotherapy and antibiotics,¹⁰ but we do not advocate any one anaesthetic technique as being superior to others for these patients. The important points are preoperative assessment and an awareness of the relevant pathophysiology. Spontaneous respiration seems acceptable for minor procedures not requiring relaxation; controlled ventilation is probably indicated in more prolonged anaesthesia. Regional analgesia for postoperative pain avoids the need for using morphine in patients who may have reduced respiratory sensitivity to carbon dioxide.

We thank Miss Anne Barrett who carried out most of the lung function tests and blood gas estimations and our surgical and anaesthetic colleagues who were involved in the care of these patients.

References

- ¹ Diamant, M. L., and Palmer, K. N. V., *Lancet*, 1967, 1, 1251.
- ² Stein, M., et al., *Journal of the American Medical Association*, 1962, 181, 765.
- ³ Thornton, J. A., *Problem of Anaesthesia and Surgery in Relation to Chronic Respiratory Disease*, M.D. Thesis, University of London, 1969.
- ⁴ Cotes, J. E., *Lung Function, Assessment and Application in Medicine*. Oxford, Blackwell, 1965.
- ⁵ Raine, J. M., and Bishop, J. M., *Journal of Applied Physiology*, 1963, 18, 284.
- ⁶ Howard, P., *British Medical Journal*, 1972, 2, 89.
- ⁷ Appleberg, M., Gordon, L., and Fatti, L. P., *British Journal of Surgery*, 1974, 61, 57.
- ⁸ Lockwood, P., *Respiration*, 1973, 30, 105.
- ⁹ Andersen, N. B., and Ghia, J., *Journal of Thoracic and Cardiovascular Surgery*, 1970, 59, 474.
- ¹⁰ Stein, M., and Cassara, E. L., *Journal of the American Medical Association*, 1970, 211, 787.

Assessment of Regional Ventilation by Continuous Inhalation of Radioactive Krypton-81m

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British Medical Journal, 1975, 3, 673-676

Summary

A simple technique is described for producing high-quality functional images of regional ventilation during physiological tidal breathing of the inert gas ^{81m}Kr. These images are quickly obtained on a gamma-camera without the need of computerized systems for data acquisition and display and are directly comparable with those of perfusion obtained with ^{99m}Tc-labelled macroaggregates. The short time required for the procedure, its simplicity, and the extremely low absorbed-radiation dose enable serial images of ventilation to be obtained in multiple views.

Introduction

There is a need for a technique to obtain functional images of ventilation that are comparable to the routinely obtained macroaggregate perfusion scan.¹ Previous methods have involved wash-in, wash-out, or a single breath of radioactive gases or aerosols of labelled particles. Because these methods tend to be either complex or inaccurate and unphysiological they have had limited clinical application.

We present here a new simple technique to obtain functional images of the regional ventilation using a gamma-camera during tidal breathing of the inert gas ^{81m}Kr. This method overcomes many of the limitations of previous techniques and when used in conjunction with perfusion data provides unique information on regional mismatching of ventilation and perfusion.

Theory

Continuous inhalation of an inert gas results in equilibration of the gas in the lungs. If a radioactive gas with a long half life is used the count rate at equilibrium, as measured externally over the chest, will be proportional to lung volume rather than to ventilation. Regional ventilation can be assessed only from the rate of equilibration (wash-in) or elimination (wash-out) of the gas.

Consider now the continuous inhalation of the inert gas ^{81m}Kr, which has a radioactive half life of 13 seconds. The rapid radioactive decay of this gas relative to the ventilation turnover rate per unit volume will result in an alveolar concentration at equilibrium much less than that in the inspired air. Hence the contribution of the ^{81m}Kr wash-out to this dynamic process is small. Equilibration of the isotope in the lung therefore depends on the balance between arrival of ^{81m}Kr and radioactive decay. ^{81m}Kr lung counts are theoretically proportional to

$$\frac{\dot{V}}{\dot{V}/\text{vol} + \lambda}$$

where $\dot{V}$ is total ventilation, $\dot{V}/\text{vol}$ the ventilation per unit of lung volume, and λ the radioactive decay constant of ^{81m}Kr, which is 3.2/min. The denominator is dominated by the high value for λ , hence the lung signal is more dependent on ventilation ($\dot{V}$) than on ventilatory wash-out ($\dot{V}/\text{vol}$). This results in an almost linear relation between radioactivity and regional ventilation for both normal (about 1.0 l min⁻¹ l⁻¹) and reduced ventilation rates in adults (fig. 1).

Therefore, the equilibrium images recorded with the gamma-camera during the continuous inhalation of ^{81m}Kr reflect the regional arrival of the gas—that is, ventilation. In other words, continuous tidal breathing of ^{81m}Kr effectively produces the same information as the sum of a series of separate tidal breaths of a radioactive gas with a long half life.

Methods

The ^{81m}Kr, which emits 190 keV gamma-rays, can be produced continuously by passing air through a cation exchange column to which its parent isotope ⁸¹Rb is bound. ⁸¹Rb has a half life of four and a half hours and is produced for us on the M.R.C. cyclotron.² The half life of the parent enables ^{81m}Kr to be continuously available all day for ventilation measurements after a morning production of ⁸¹Rb.

If both ventilation and perfusion scans are required 1 mCi of ^{99m}Tc-labelled macroaggregates or microspheres (^{99m}Tc-HAM) is first injected intravenously with the patient in the seated position. The patient is then positioned in front of the gamma-camera (Toshiba

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