Effect of chronic airflow limitation on resting oxygen consumption

C Lanigan, J Moxham, J Ponte

Abstract
The work of breathing in patients with severe chronic airflow limitation is increased even at rest but little is known about the magnitude of this increase. Resting oxygen consumption (\(\dot{V}O_2\)), carbon dioxide production (\(\dot{V}CO_2\)), and respiratory quotient (RQ) were measured in 13 patients with severe chronic airflow limitation (mean FEV\(_1\) 0.78 l, vital capacity 2.1 l) and compared with those of 13 age, weight, and height matched control subjects. Whereas mean RQ was the same in the two groups (0.82), mean \(\dot{V}O_2\) and \(\dot{V}CO_2\) were higher in the patients (+18 ml min\(^{-1}\) and +15 ml min\(^{-1}\) respectively). When \(\dot{V}O_2\) was standardized for body surface area it was 10.9% higher in the patients (\(p < 0.05\)). If the increased resting \(\dot{V}O_2\) in these patients were solely due to increased activity of the respiratory muscles, it would represent a fourfold increase in the oxygen cost of breathing.

The diaphragm accounts for most of the work of ventilation during quiet breathing.\(^1\) In patients with chronic airflow limitation additional energy is expended throughout the breathing cycle, most of which is borne by the inspiratory muscles.\(^2\) These patients may also have raised energy requirements because of a hyperkinetic circulation, increased blood viscosity, shifts in the haemoglobin dissociation curve, hypoxia, and increased shunt and dead space.\(^3\) Some of the increased work results from the need to generate larger subatmospheric pressures to expand the thoracic cage and from the loss of mechanical efficiency of the diaphragm.\(^3\)

Chronic airflow limitation causes a disproportionate increase in total oxygen consumption (\(\dot{V}O_2\)) for a given minute ventilation during exercise.\(^4,5\) Evidence for an increased energy expenditure at rest is less convincing. Consequently we undertook a prospective controlled study of resting \(\dot{V}O_2\), carbon dioxide production (\(\dot{V}CO_2\)), and respiratory quotient (RQ) in chronic airflow limitation.

Methods

PATIENTS AND CONTROL SUBJECTS

Thirteen patients (nine of them male)—mean (SD) age 64 (6.8) years, weight 61 (15.6) kg, height 1.67 (0.1) m, body surface area 1.67 (0.19) m\(^2\)—were recruited, after informed consent had been obtained and with the approval of the hospital ethics committee, from the chest clinics at King's College and Dulwich Hospitals. They had moderate to severe airflow limitation with minimal reversibility and large lung volumes with reduced carbon monoxide transfer (table). Control subjects consisted of 13 healthy, non-smoking volunteers (six male), mean age 53 (16) years, weight 65 (12.5) kg, height 1.67 (0.12) m, and body surface area 1.72 (0.12) m\(^2\).

EXPERIMENTAL PROCEDURE

Resting \(\dot{V}O_2\), \(\dot{V}CO_2\), and RQ were measured by an open canopy technique with a mass spectrometer (Airspec Ltd).\(^8\) The subject's head and chest were enclosed in a clear chamber, from which gases were exhausted at a set rate of 80 to 90 l min\(^{-1}\) through a mixing box to which argon (Ar, 250 ml min\(^{-1}\)) was added as a tracer for measurement of flow. The fractional differences between oxygen and carbon dioxide in air and in the outflow from the chamber, measured at six second intervals, were multiplied by flow through the system to produce \(\dot{V}O_2\), \(\dot{V}CO_2\), and RQ (\(\dot{V}CO_2/\dot{V}O_2\)). The resolution, tested by methanol combustion, was 8 ml min\(^{-1}\) for \(\dot{V}O_2\) and 6 ml min\(^{-1}\) for \(\dot{V}CO_2\).

Patients' medication was unaltered; all subjects had at least 30 minutes' rest within the chamber before the study started and listened to music throughout the 20 minutes' continuous gas exchange measurement (200 data points) that followed.

DATA ANALYSIS

Gas analysis was corrected for instrument drift and methanol combustion tests and converted to STPD. The mean and SD for two consecutive 10 minute periods were calculated, and the lower of the two subsequently included in the analysis. \(\dot{V}O_2\) values were standardised for body surface area (BSA) and calculated as follows:\(^9\)

\[
\log BSA = \log \text{weight} \times 0.425 + \log \text{height} \times 0.725 + 1.8564
\]

Resting energy expenditure (kJ/day) was compared with predicted values\(^10\) on the basis of an oxygen equivalent of 4.83 kJ/ml. Differences between groups were analysed by means of the Wilcoxon signed rank test (\(p < 0.05\)).

Results

There were more men in the patient group, but there were no significant differences in terms of...
Mean pulmonary function data for patients*

<table>
<thead>
<tr>
<th>Variable</th>
<th>Initial</th>
<th>After bronchodilator</th>
<th>% predicted</th>
</tr>
</thead>
<tbody>
<tr>
<td>PEF (l min⁻¹)</td>
<td>170 (92-5)</td>
<td>180 (95-7)</td>
<td>41 (21-2)</td>
</tr>
<tr>
<td>FEV₁ (l)</td>
<td>0.78 (0-36)</td>
<td>0.83 (0-36)</td>
<td>31 (11-3)</td>
</tr>
<tr>
<td>VC (l)</td>
<td>2.1 (0-6)</td>
<td>2.4 (0-5)</td>
<td>62 (15-6)</td>
</tr>
<tr>
<td>FEV₁/FVC</td>
<td>0.78 (0-72)</td>
<td>0.75 (0-72)</td>
<td>55 (21-5)</td>
</tr>
<tr>
<td>TLC (mmol min⁻¹ l⁻¹)</td>
<td>4.5 (1-39)</td>
<td>5.7 (2-22)</td>
<td>139 (37-6)</td>
</tr>
<tr>
<td>KCO</td>
<td>1.3 (0-42)</td>
<td>8.8 (2-53)</td>
<td>267 (90-1)</td>
</tr>
<tr>
<td>RV (l)</td>
<td>5.1 (2-20)</td>
<td>7.3 (2-23)</td>
<td>139 (37-6)</td>
</tr>
<tr>
<td>TLCo (l)</td>
<td>7.3 (2-23)</td>
<td>139 (37-6)</td>
<td>267 (90-1)</td>
</tr>
</tbody>
</table>

*

Discussion

Although numbers were small the difference in VO₂ and VCO₂/BSA between the groups is likely to be real and of considerable clinical interest, as it may largely reflect the difference in the metabolic cost of breathing between the two groups. If we assume normal cost of breathing is about 21 W/kg, then the difference in 300 gr of total VO₂ (that is, < 8 ml min⁻¹ kg⁻¹), the cost of breathing in our patients was about 24 ml min⁻¹—four times the normal value, representing 10⁻⁸⁰, of total VO₂. At least three other factors in addition to those listed in the introduction might have influenced the result.

Somatometric differences

Large scale studies of resting energy expenditure have shown that VO₂ declines with age, increases with weight, height, and body surface area, and is lower in women than men. The group differences were too small to account for the differences in VO₂ and VCO₂ after normalisation for body surface area. Although there were more men in the patient group, separate analysis of the data for men still showed a higher mean VO₂/BSA, the difference being 23 ml min⁻¹ (p < 0.05).

Drug treatment

Patients were in a stable clinical condition but their drug treatment included beta agonists, methyl xanthenes and corticosteroids, all potentially capable of increasing VO₂. The precise influence of these drugs on VO₂ was not measured because halting treatment might have exacerbated the condition, but bronchodilator treatment itself might have reduced the work of breathing. The values of VO₂ and VCO₂ obtained probably reflected the "best case" ventilatory conditions.

Acute hyperventilation

Placing dyspnoeic patients inside a chamber could have provoked hyperventilation and would have caused the RQ to rise. Acclimatisation minimised this effect as no individual RQ exceeded 0.88, and the average RQ was identical and normal at 0.82 for both groups.

The reliability of our method of measuring VO₂ and VCO₂ was confirmed by the close agreement found between measured and predicted resting energy expenditure in the control subjects. The lack of correlations between VO₂/BSA and indices of airways obstruction is not surprising given the complex relationship between lung mechanics and the work of breathing, the small numbers studied, and the relatively uniform severity of the airflow obstruction. We decided not to monitor minute or tidal ventilation because of the inevitable artefact introduced by the recording method, which could have produced different weighting in patients and in control subjects. Previous authors have reported a 30% increase in minute volume when a facemask is worn by normal subjects and a 14% increase when a mouthpiece and noseclips are used. Indirect methods of measuring tidal ventilation (for example, magnetometry and inductive plethysmography), though adequate for detecting relative changes within subjects, are not suitable for measuring absolute differences between subjects.

One other group has found an increased metabolic rate in patients with chronic airflow obstruction. Although they used a similar
open canopy method and control subjects, the data obtained by these authors are not comparable to ours as they studied only malnourished patients. Another group specifically studied resting VO₂ in 43 patients with chronic airflow limitation 16 but normal control subjects were not studied. Several other studies have obtained VO₂ data from smaller groups of patients (up to 10) before exercise or rebreathing tests. 18-21 All these studies made use of mouthpieces, noseclips or face masks, which might have affected the results. Nevertheless, all authors have reported increases in metabolic rate, which was 8.5-20% above predicted values.

In conclusion, our results indicate that the resting VO₂ in patients with chronic airflow limitation, normally nourished and undisturbed by measuring appliances, is about 10% higher than in normal subjects. If this increase in VO₂ were solely due to greater respiratory muscle work, then the patients had a fourfold increase in the oxygen cost of breathing at rest.

Effect of chronic airflow limitation on resting oxygen consumption.

C Lanigan, J Moxham and J Ponte

Thorax 1990 45: 388-390
doi: 10.1136/thx.45.5.388

Updated information and services can be found at:
http://thorax.bmj.com/content/45/5/388

These include:

Email alerting service

Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/