Pulmonary endocrine cells of Aymara Indians from the Bolivian Andes

D Williams, D Heath, J Gosney, J Rios-Dalenz

Abstract

Introduction There is evidence to suggest that life at high altitude causes changes in the population of pulmonary endocrine cells, possibly because of exposure to chronic hypoxia. A study was made of the populations of pulmonary endocrine cells in three Aymara Indians and three Mestizos of La Paz (3600 m), Bolivia, which were compared with those in four white lowlanders.

Methods Pulmonary endocrine cells were immunolabelled for neurone specific enolase and their two major secretory products, gastrin releasing peptide and calcitonin, and their numbers expressed per cm² of tissue section.

Results No differences in morphology, number, content, or distribution of immunoreactive cells were found when the native highlanders were compared with the lowlanders.

Conclusions If chronic hypoxia as such exerts an influence on human pulmonary endocrine cells it was not apparent in this morphological study. There was no increase in gastrin releasing peptide containing pulmonary endocrine cells, such as have previously been seen in patients with pulmonary hypertension characterised by plexogenic pulmonary arteriopathy. This may be due to the fact that in plexogenic pulmonary arteriopathy there is free migration of smooth muscle cells. Although three of the highlanders in this present study showed pulmonary vascular remodelling, this was in contrast only modest.

(Thorax 1993;48:52-56)

Clear cells in the epithelium of many organs were first described in detail by Feyrter, who regarded them as components of a "diffuse endocrine epithelial organ," now known generally as the "diffuse endocrine system." In the lung, in all species so far studied, they occur in the epithelium of bronchi and bronchioles and to a lesser extent in alveolar walls. They are found as single cells or in groups. Some of these groups of cells have been found to be innervated and consequently have been termed "neuroepithelial bodies"; others are probably merely aggregates of solitary cells. Experimental evidence has suggested that, in acute hypoxia at least, neuroepithelial bodies act as chemoreceptors monitoring oxygen tension in the airways. Hence chronic exposure to the hypobaric hypoxia of natural high altitude might be expected to affect the function, morphology, content, or distribution of these cells. The aim of this study was to see whether any such changes were apparent in the lungs of a group of Aymara Indians collected during an expedition to the Bolivian Andes.

Methods

REGION OF STUDY

Obtaining permission and facilities for the performance of necropsies presents difficulties in the Andes and a study such as that presented here has to be carried out in a pathology laboratory at very high altitude adjacent to a hospital providing a service in morbid anatomy. Very few areas meet these requirements. One that does is La Paz, the capital of Bolivia, which is situated on the eastern slopes of the Andean range. There is considerable variation in the elevation of its various districts. The highest point is the airport, El Alto, at an altitude of 4120 m, and the lowest comprises the nearby jungle valleys at about 3200 m; the central plaza is at 3600 m and may be taken as an average for the city. We were privileged to receive lung tissue for study from the Obrero Hospital and the Hospital of the University of San Andres.

SUBJECTS

We studied 10 subjects, in the age range 15–42 years (table 1). Six had been born in La Paz, or were long standing residents, and were free of cardiopulmonary disease at death. Although all had died in hospital in La Paz, it was important to ascertain both the altitude of their birthplace and their normal place of residence, as in a large and busy city such as La Paz not everyone dying at high altitude is native to the area. Consequently, ethnic origin was considered important. Its definition was that used in the 1976 Bolivian census. Individuals were classified as "Indian" if they had an Aymara surname, had emigrated to La Paz from a rural community, and lived in a predominantly Indian neighbourhood. Those termed "Mestizo" had a mixed Spanish and Indian family history and had lived in La Paz for at least one generation. The surname in the Mestizo group could be either Spanish or Aymara. By these criteria three of our highlanders were Aymara Indians and three were Mestizos. The four remaining subjects, who constituted a control group, were white lowlanders (cases 7–10, table 1). These also died without any cardiopulmonary disease.
Pulmonary endocrine cells of Aymara Indians from the Bolivian Andes

Table 1  Age, sex, ethnic background, and diagnosis at necropsy

<table>
<thead>
<tr>
<th>Case no</th>
<th>Age (y)</th>
<th>Sex</th>
<th>Ethnic background</th>
<th>Diagnosis at necropsy</th>
</tr>
</thead>
<tbody>
<tr>
<td>HIGH ALTITUDE</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>1</td>
<td>28</td>
<td>M</td>
<td>Aymara</td>
<td>Cerebellar tumour</td>
</tr>
<tr>
<td>2</td>
<td>37</td>
<td>M</td>
<td>Aymara</td>
<td>Peritonitis: ruptured appendix</td>
</tr>
<tr>
<td>3</td>
<td>42</td>
<td>M</td>
<td>Aymara</td>
<td>Gastric lymphoma</td>
</tr>
<tr>
<td>4</td>
<td>15</td>
<td>F</td>
<td>Mestizo</td>
<td>Lupus erythematosus</td>
</tr>
<tr>
<td>5</td>
<td>27</td>
<td>F</td>
<td>Mestizo</td>
<td>Pyelonephritis</td>
</tr>
<tr>
<td>6</td>
<td>32</td>
<td>M</td>
<td>Mestizo</td>
<td>Pyelonephritis</td>
</tr>
<tr>
<td>SEA LEVEL</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>7</td>
<td>21</td>
<td>M</td>
<td>Caucasian</td>
<td>Road accident</td>
</tr>
<tr>
<td>8</td>
<td>29</td>
<td>F</td>
<td>Caucasian</td>
<td>Friedreich's ataxia</td>
</tr>
<tr>
<td>9</td>
<td>23</td>
<td>M</td>
<td>Caucasian</td>
<td>Road accident</td>
</tr>
<tr>
<td>10</td>
<td>24</td>
<td>M</td>
<td>Caucasian</td>
<td>Road accident</td>
</tr>
</tbody>
</table>

LUNG TISSUE

Lungs were distended with 10% neutral buffered formalin until the pleural surfaces were smooth. Because only a limited amount of tissue could be brought back from La Paz care, it was taken to ensure, so far as possible, that pulmonary sampling was consistent in its site in all subjects studied from both altitudes. Thus tissue was taken from a point midway between the hilum and the pleural surface such that intrapulmonary bronchi and terminal bronchioles, the sites of greatest concentration of human pulmonary endocrine cells, were well represented; but more distal airways and pulmonary parenchyma were also included. In most cases, eight blocks of tissue were taken from each of the three lobes of the right lung; but in case 5 difficulty in distending the right lung made it necessary to use the left, and eight blocks of tissue were taken from both lobes. These blocks were processed, embedded in paraaffin wax and sections cut at a thickness of 4 \( \mu \)m.

IMMUNOLABELLING

These sections were labelled for neurone specific enolase (NSE), gastrin releasing peptide (GRP), and calcitonin by the avidin biotin complex technique. Antisera were obtained from Dako Ltd (NSE and calcitonin) and Cambridge Research Biochemicals (GRP). After being dewaxed and taken to absolute alcohol the sections were pretreated with 1% hydrogen peroxide in methanol for 20 minutes and with normal swine serum for 20 minutes. This prevented false positive staining due to endogenous peroxide and non-specific binding of immunoglobulin respectively. The primary antisera were applied at optimum dilution (neurone specific enolase at 1:2000, gastrin releasing peptide at 1:4000, calcitonin at 1:2000) and incubated for one hour, after which sections were washed in buffered saline. A biotinylated anti-rabbit secondary antibody was applied at a dilution of (neurone specific enolase at 1:2000, gastrin releasing peptide at 1:4000, calcitonin at 1:2000) and incubated for one hour, after which sections were washed in buffered saline. A chromogen used was diaminobenzidine, which after the addition of hydrogen peroxide produces a black-brown reaction product at antigenic sites. Positive

Table 2  Numbers of pulmonary endocrine cells per cm²

<table>
<thead>
<tr>
<th>Case no</th>
<th>NSE</th>
<th>GRP</th>
<th>Calcitonin</th>
</tr>
</thead>
<tbody>
<tr>
<td>HIGH ALTITUDE</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>1</td>
<td>50</td>
<td>40</td>
<td>0.3</td>
</tr>
<tr>
<td>2</td>
<td>42</td>
<td>37</td>
<td>0.2</td>
</tr>
<tr>
<td>3</td>
<td>40</td>
<td>33</td>
<td>0.1</td>
</tr>
<tr>
<td>4</td>
<td>3.5</td>
<td>3.4</td>
<td>0.2</td>
</tr>
<tr>
<td>5</td>
<td>3.8</td>
<td>3.5</td>
<td>0.2</td>
</tr>
<tr>
<td>6</td>
<td>4.7</td>
<td>4.4</td>
<td>0.2</td>
</tr>
<tr>
<td>Mean (SD)</td>
<td>4.2 (0.5)</td>
<td>3.7 (0.4)</td>
<td>0.2 (0.06)</td>
</tr>
<tr>
<td>SEA LEVEL</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>7</td>
<td>6.8</td>
<td>6.5</td>
<td>0.2</td>
</tr>
<tr>
<td>8</td>
<td>20.3</td>
<td>15.6</td>
<td>4.5</td>
</tr>
<tr>
<td>9</td>
<td>7.2</td>
<td>7.0</td>
<td>0.3</td>
</tr>
<tr>
<td>10</td>
<td>10.4</td>
<td>10.0</td>
<td>0.2</td>
</tr>
<tr>
<td>Mean (SD)</td>
<td>11.2 (5.5)</td>
<td>9.8 (3.6)</td>
<td>1.3 (1.9)</td>
</tr>
</tbody>
</table>

NSE—neurone specific enolase; GRP—gastrin releasing peptide.
tissue controls included immunolabelling of human fetal lung (gastrin releasing peptide), human medullary carcinoma of the thyroid (calcitonin), and human pancreas (neurone specific enolase). In the negative control procedures labelling was carried out after replacement of the primary antiserum with non-immune serum and after omission of each stage of the procedure in turn. Sections adjacent to those used for immunohistochemistry were stained by the elastic van Gieson method and with haematoxylin and eosin.

CELL COUNTING
The numbers of immunoreactive cells in each section were counted by using a microscope fitted with an eyepiece graticule of 100 squares. After calibration by means of a stage micrometer this enabled both the number of cells to be counted and the area of a section to be measured. The whole of a section was examined and the total area measured was in the range 32.5–50.5 cm² per case, 1 cm² yielding 100 fields, a total of 3250–5050 fields. A search was made for the remodelling of the pulmonary arterial tree that is seen in hypobaric hypoxia.8,7 Sections stained with haematoxylin and eosin served to exclude any undisclosed pathological changes.

Results
There was no difference in numbers, distribution, morphology, or content of endocrine cells when the two groups were compared. The numbers of immunoreactive cells per cm² of section for the two peptides studied are shown in table 2. Although there appeared to be a tendency for more pulmonary endocrine cells to occur in the lungs of the lowlanders, there was a range of less than 2 standard deviations on either side of the means and so both sets of data came under the same distribution curve. In the lungs of both groups nearly all cells were solitary (fig 1) and usually in the epithelium of bronchi and bronchioles; only one cluster of cells was seen and it was in an alveolar duct (fig 2). Solitary cells had the shape and situation typical of pulmonary endocrine cells in normal mammalian lungs. They were usually elongated in shape with a round, basal nucleus and located on the basement membrane (fig 1). Most contained gastrin releasing peptide, calcitonin containing cells comprising a much smaller population (table 2, fig 3). All cells that were labelled for one or other of these secretory products (there was no evidence of co-storage) were neurone specific enolase positive, but occasional neurone specific enolase immunoreactive cells contained neither.

All three Aymaras (cases 1–3) showed muscularisation of their pulmonary arterioles, and in addition case 1 showed intimal longitudinal muscle and the formation of inner muscular tubes.6,7 There were no abnormalities of the pulmonary arterial tree in cases 4–6 or in the lungs of the lowlanders.

Discussion
Although acute exposure to hypoxia undoubtedly induces activity in neuroepithelial bodies in rabbits,4,8 the effects of chronic hypoxia are uncertain. Studies on animals in the laboratory, where hypoxia has been induced by using hypobaric chambers, have produced contradictory results.8–15 This may be due to a combination of factors, including variability in experimental conditions, differences in species or strain susceptibility,13 and the fact that the changes induced by hypoxia may be subtle, requiring careful study for their detection.16,17

Studies at natural high altitude have produced similarly variable results.16–21 Rabbits native to the Peruvian Andes and thus exposed all their lives to hypobaric hypoxia were reported to show increased clusters of argyrophilic cells in the lung,18 but we were unable to find such an effect in various animal species in the Himalaya of Ladakh.19 We studied the pulmonary endocrine cells of sheep, goats, and the yak, together with the yak’s interbreeds with cattle (the dzos and stol) at Sakti (4500 m) and compared them with those of sheep and goats at Srinagar (1590 m). There were no differences in their number or distribution, as shown by immunolabelling with neurone specific enolase, that could be ascribed to the effects of hypobaric hypoxia, though species differences were substantial. More recently guinea pigs from the Peruvian Andes have been shown to have significantly more clusters of pulmonary endocrine cells than controls from sea level, whereas there was no difference in the number of solitary cells.20

In an abstract Memoli et al21 reported that in six of 20 samples of lung from native highlanders of Bolivia, who had spent their entire lives at altitudes ranging from 3500 to 4300 m, there were significantly more neuroepithelial bodies, as identified by neurone specific enolase, than in a group from sea level. The present study, however, is the first in which individual peptides have been investigated in such subjects. In the normal adult lung endocrine
cells contain predominantly gastrin releasing peptide and calcitonin, especially the former. The absence of discernible differences between the two groups of subjects in the present study may simply be because numbers were too small, or because any changes that had occurred in the highlanders were too subtle to be detected by the methods we used. Furthermore, we cannot entirely exclude the possibility that differences in endocrine cell numbers were present in those parts of the lung that we did not sample—namely, the extrapulmonary and large intrapulmonary bronchi and subpleural parenchyma. Any such difference, however, would be likely to be most apparent in that part of the lung most richly populated with endocrine cells, the intrapulmonary bronchi and terminal bronchioles, which we were careful to include in the tissue we took.

We are confident that there was no increase in endocrine cell clusters in the subjects from La Paz, as has been reported in previous studies in man and animals from high altitude, when hypoxia has been postulated as its cause.

The lack of any apparent proliferation of gastrin releasing peptide containing cells in the lungs of the highlanders contrasts with the situation in subjects with plexogenic pulmonary arteriopathy due to primary pulmonary hypertension or secondary to congenital cardiac shunts. In these conditions, particularly when associated with cellular plexiform lesions and especially in the plexiform phase, when migration of vascular smooth muscle cells from the media to the intima is at its height, gastrin releasing peptide containing cells are abundant. A similar association has been described in subacute infantile mountain sickness in Tibet. This condition is a manifestation of a failure of initial aclimatisation to high altitude in Han infants taken up to reside in Lhasa, and is also characterised by migration of smooth muscle cells from the media to the intima and a prominence of gastrin releasing peptide containing endocrine cells.

Possibly these findings are related to the role of gastrin releasing peptide as a trophic agent. It has been shown to stimulate the growth of normal human bronchial epithelial cells in vitro as well as murine fibroblasts. It increases the incorporation of tritiated thymidine into developing bronchial epithelium of fetal mouse lungs in utero and in vitro and of human fetal lung in organ culture.

Aymara Indians have studied previously and in the present study we have found remodelling of the pulmonary arterial tree but with only limited migration of medial vascular smooth muscle cells into the intima. In the present study we could not find an association between the migration of smooth muscle cells and prominence of gastrin releasing peptide containing cells. This may be because the remodelling of the pulmonary arterial tree that had occurred in the Aymara Indians is characterised by only limited migration of smooth muscle cells.

Increased numbers of calcitonin containing endocrine cells seem to be most strongly associated with inflammation, which was not a feature of the lungs examined in the present study. Proliferation of such cells does occur in its absence and may in some way be associated with a more generalised disturbance in the structure and function of the pulmonary endocrine system.


Pulmonary endocrine cells of Aymara Indians from the Bolivian Andes.

D Williams, D Heath, J Gosney and J Rios-Dalenz

Thorax 1993 48: 52-56
doi: 10.1136/thx.48.1.52