Renal Excretion of Bicarbonate in High Altitude Natives and in Natives with Chronic Mountain Sickness

CARLOS MONGE C., RODOLFO LOZANO, AND AMADOR CARCELÉN

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Natives of high altitudes are known to have a low tension of CO₂ in the arterial blood, due to their greater ventilation (1-6). In 1928 Monge M. (7, 8) described a disease in the Peruvian Andes that is characterized by excessive polycythemia (above the physiological level for the corresponding altitude) and congestive symptoms that are relieved dramatically on descent to lower levels. This disease may develop in sea level people after years of residence at high levels, or in natives of the high altitudes, and it has been called chronic mountain sickness or Monge’s disease (9-11). Hurtado (12) has found that the arterial oxygen saturation of the hemoglobin of patients with chronic mountain sickness is lower than that which corresponds to the physiological altitude level, and he has recently postulated hyperventilation, secondary to reduced sensitivity of the respiratory center to CO₂, as an important factor in the pathogenesis of this entity (13). On the other hand, Pitts and Lotspeich (14) and Pitts, Ayer, and Schiess (15) have defined the threshold of bicarbonate both in animals and in humans, and Dorman, Sullivan, and Pitts (16) have shown that the tension of CO₂ in the arterial blood is one of the factors that govern the reabsorption of bicarbonate in the renal tubule.

The present investigation is concerned with the parameters of acid-base equilibrium of the arterial blood in sea level controls, natives living at 4,300 m above sea level, and natives living at the same altitude but suffering from chronic mountain sickness, and attempts to establish a relationship between the pressure of the CO₂ in the arterial blood and the renal reabsorption of bicarbonate in the three groups. The results show that the maximal reabsorption (Tm) of bicarbonate, expressed as mmoles per 100 ml of glomerular filtrate, is the same in the high altitude natives as in the sea level controls, despite the lower arterial Pco₂ of the former. The cases of chronic mountain sickness have a greater maximal bicarbonate reabsorption and a higher arterial Pco₂ compared with the control native group. The results are interpreted as suggesting that the normal high altitude native is in a new state of acid-base equilibrium with low arterial Pco₂ and a normal bicarbonate reabsorption. The possible roles of high arterial Pco₂, hypokalemia, and anoxia in the elevation of bicarbonate Tm of patients with chronic mountain sickness is discussed.

Methods

The studies have been carried out in 17 human male volunteers: six were normal people at sea level, six were normal high altitude natives, and five were high altitude natives with chronic mountain sickness. The first group was studied in Lima (150 m above sea level) and the last two in Cerro de Pasco (4,300 m above sea level). In contrast with other studies carried out by us and others in the Andes of Peru, we have selected for the sea level controls a group of men native to sea level and who in our opinion have a closer nutritional, anthropological, and social resemblance to the altitude natives than former sea level groups selected among the higher classes. This selection may explain some of the biochemical abnormalities found in the blood of the sea level group, such as slight anemia, low serum potassium, and perhaps the lower blood pH.

The criteria for diagnosis of chronic mountain sickness were a hematocrit of 70% or higher, congestive symptoms, and an arterial oxygen saturation significantly lower than the average usually found at the altitude of Cerro de Pasco. All the subjects had a chest X-ray film taken in addition to a physical examination, in order to discard gross pulmonary pathology. The volunteers slept in the laboratory the night before the experiment, which was conducted early in the morning under fasting conditions and at an approximate room
temperature of 20° C, both at sea level and at high altitude. Adequate hydration was started the night before the studies by oral water intake.

The experiment began with the insertion of a Cournand needle into the brachial artery, which remained in place until the end of the study. An indwelling bladder catheter was inserted, and after a few minutes of rest, and with the subject quiet, a sample of arterial blood was obtained and a urine collection completed. These samples were used to calculate the parameters of acid-base equilibrium under “basal” conditions. After the basal samples were obtained, an indwelling plastic catheter was placed in one vein of the arm opposite to the one with the arterial needle, and a constant infusion of 1.5% sodium bicarbonate was started at a rate of 10.1 ml per minute. This solution contained inulin calculated to give an approximate concentration of 2.0 mg per 100 ml in the blood plasma. After 1 hour of equilibration, urine was discarded and four consecutive 20-minute urinary collections were made under anaerobic conditions. No air or saline was used to wash the bladder, and only manual compression assured adequate emptying.

Arterial blood samples were obtained under anaerobic conditions in the middle of the urine collection periods with heparinized syringes. All the chemical analyses were carried out on arterial blood, with the exception of hemoglobin and hematocrit, which were determined on venous blood the day before the experiment. With the exception of sodium, potassium, and chloride in plasma and urine, which were determined in the laboratory at sea level, the different analyses of the high altitude groups were carried out at the Cerro de Pasco laboratory.

Immediately after obtaining blood and urine, their pH was determined in a radiometer M4 pH meter at 38° C. The machine was standardized frequently by a Beckman buffer that had previously been checked with a disposable high-precision buffer of pH 7.381 provided by the manufacturing company. The Pco2 was determined in anaerobically separated plasma by the Astrup equilibration technique (17) with a gas cylinder of an approximate CO2 pressure of 40 mm Hg and a temperature of 38° C. At sea level, the CO2 concentration was checked by a Sholander gas analysis apparatus and by equilibrating a solution of 25 mmoles per L of potassium bicarbonate as recommended by Astrup (17). This technique gave good checks with the direct gas analyses, and it was used in the high altitude laboratory for a daily control of the CO2 concentration. With the actual blood pH and the plasma pH after equilibration known, the Pco2

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* Hgb. = hemoglobin; hct. = hematocrit. The subscript s corresponds to determinations carried out in plasma.
was calculated by a nomogram. Plasma bicarbonate was obtained by using the Henderson-Hasselbalch equation with a pK' of 6.1 and a CO₂ solubility factor of 0.0300. Urine CO₂ was determined in the Van Slyke manometric apparatus immediately after collection. Urine bicarbonate was calculated by using the Henderson-Hasselbalch equation with a pK' of 6.1 and a CO₂ solubility factor of 0.0300. Inulin was determined in the plasma and urine according to Schreiner (18), plasma and urine chloride according to Keys (19). Plasma and urine sodium and potassium were measured by internal standard flame photometry. Donnan factors of 0.95 for anions and of 1.05 for cations were used in the calculation of filtered loads of ions in all subjects.

Results

Table I shows that the high altitude natives had a lower body weight and a smaller surface area than the sea level controls. Detailed anthropological analyses of them have been made by Hurtado (20). The hemoglobin concentrations and the hematocrits of the Cerro de Pasco normal natives were in the expected range (21), but they were found to be much higher in the natives with chronic mountain sickness whose arterial hemoglobin oxygen saturations were 59.6, 74.2, 75.9, 78.8, and 80.0%, in contrast with 81%, which is the average value corresponding to that in Morococha (4,540 m).

The mean blood pH of the normal Cerro de Pasco natives was 7.431, and in cases of chronic mountain sickness, it was 7.429. Both are higher than the sea level control group, which was 7.362. There is no significant difference between the means of the high altitude groups. The mean Pco₂ of the normal natives was 32.5 mm Hg, lower than 39.6, which was the average at sea level. The mean plasma bicarbonate concentration in the normal native group was 20.9 mmol/L and does not differ significantly from 21.7,
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**Cerro de Pasco (4,300 m)**

**Normal natives**

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**Cerro de Pasco (4,300 m)**

**Natives with chronic mountain sickness**

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<td>SD</td>
<td>2.89</td>
<td>22.5</td>
<td>0.2 2.2 0.024 1.7 0.5 0.3</td>
<td>43.2 14.6 0.182 18.9 28.4</td>
<td>0.08 0.24 0.4</td>
</tr>
</tbody>
</table>

* Each datum is the mean of four consecutive clearance periods.

TABLE II

Tubular reabsorption of bicarbonate, chloride, and sodium during the infusion of sodium bicarbonate*
which was the corresponding sea level figure. In the natives with chronic mountain sickness the mean was 25.1, significantly higher than the other two groups.

A summary of the acid-base equilibrium as expressed by the Henderson-Hasselbalch equation is given in Figure 1. This corresponds to the triaxial nomogram of Shock and Hastings, reconstituted in such a way that the three axes intersect at the values corresponding to the mean of the sea level group. The areas numbered 1, 2, and 3 represent the basal figures obtained in the sea level controls, high altitude natives, and the natives with chronic mountain sickness, respectively. There is no overlapping of figures. When compared with the sea level group, the normal natives of the altitude are in the area of high pH and low Pco2, and the natives with chronic mountain sickness belong to an area of high pH with high bicarbonate and normal Pco2.

Table I shows the plasma concentrations of sodium, potassium, and chloride, which did not differ significantly in the three groups.

Table II gives the experimental details on each subject. In spite of changes in the glomerular filtration rate, the reabsorption of bicarbonate, when expressed as mmoles per 100 ml glomerular filtrate, was a constant. The constancy of the sodium and chloride reabsorption, similarly expressed, was also evident.

Each individual figure is the average of four periods. The data on potassium reabsorption were extremely variable in the three groups and have not been included. The only significant differences between the means in the bicarbonate reabsorption were in the natives with chronic mountain sickness, which were higher than in the other two groups (p < 0.001). The plasma bicarbonate concentration of the same group was also higher than in the other two groups (p < 0.001).

Figure 1 includes, in addition to the areas numbered 1, 2, and 3, already described, the areas 1', 2', 3'. These areas have been constructed with the Pco2 and pH values corresponding to the highest plasma bicarbonate concentration reached during the infusion of bicarbonate. Both high altitude groups reached a higher pH than the sea level group.

Both in the sea level group and in the natives with chronic mountain sickness, the Pco2 response to the induced metabolic alkalosis was negligible. In the normal natives, there was a small but significant rise of the Pco2 after the bicarbonate infusion.

**Discussion**

The healthy native of high altitudes has a high ventilatory rate and a low arterial Pco2. His blood pH was found to be within normal limits but slightly lower than the sea level controls by Dill, Talbott, and Consolazio (3) and Hurtado, Aste-Salazar, Velasquez, and Reynafarge (4−5), using an equilibration technique and a calculated pH. Monge, Encinas, Heral, and Hurtado (2) found a higher venous pH in the natives with a colorimetric technique. Recently, Severinghaus and Carcelén (22), employing the glass electrode, found a pH of 7.431, 7.424, and 7.426, corresponding to altitudes of 3,720, 4,545, and 4,820 m in Peruvian high altitude natives. Our results show a mean arterial blood pH of 7.431 in our native group living at 4,300-m altitude, in close agreement with those obtained by Severinghaus and Carcelén. As these authors, after completion of their work, had to recalculate their values using frozen samples of a buffer that had deteriorated while in transit to Peru, their absolute pH values should be considered with caution. Nevertheless, our figure for arterial Pco2 in the normal natives of 32.5 mm Hg is identical to their value found at Morococha (4,540 m). These authors were particularly careful in their determination of this value, using the Pco2 electrode, the alveolar air technique, and the pH−CO2 technique. Our patients with chronic mountain sickness had a pH that is no different from the one in the normal high altitude native group, but the Pco2 was definitely higher.

Figure 1, areas 1, 2, and 3, shows that when the three variables of the Henderson-Hasselbalch equation are taken into consideration, the three groups studied occupy a different position in the triaxial nomogram of Shock and Hastings. This different location may be an important characteristic of chronic mountain sickness and indicates a difference from the normal native population.

As our sea level group pH average value of 7.362 is somewhat lower than the average reported for
sea level groups in general, a second group of
seven sea level volunteers of similar anthropologi-
cal and social characteristics was studied after
the completion of this work, and a mean of 7.370 was
found. This figure does not differ significantly
from the one reported in this study.

Hurtado (13) has recently postulated hypo-
ventilation, secondary to diminished sensitivity of
the respiratory center to CO₂ as an important
factor in the pathogenesis of chronic mountain
sickness. The high arterial Pco₂ found in the
patients with altitude sickness, in comparison
with their normal native control group and in the
absence of gross pulmonary pathology, is in ac-
cordance with Hurtado’s views.

When the bicarbonate reabsorption is ex-
pressed as mmoles per 100 ml glomerular filtrate, the
difference between the means of the sea level and
high altitude normal native groups is not sig-
ificant. The corresponding figure is higher in the
natives with chronic mountain sickness. Hy-
perventilation with a low arterial Pco₂ has been
shown to depress the bicarbonate threshold in
acute experiments in dogs (23). Our results
suggest that the native of high altitudes is in a new
steady state of acid-base equilibrium with low
Pco₂ and normal bicarbonate Tm.

The higher bicarbonate Tm found in patients
with chronic mountain sickness could be ex-
plained on the basis of their higher Pco₂, which
would increase their tubular reabsorption of bi-
carbonate as has been shown to occur in acute
experiments in dogs (23). The renal adjustment
would keep the arterial pH at the same level as
the native control group. Robin has recently
pointed out that total renal compensation in
respiratory acidosis is a frequent finding (24).
But other factors should also be considered, such
as the possibility of hypokalemia, which is known
to increase the renal bicarbonate reabsorption.
Table I shows that low serum potassium values
are found in the three groups studied. When the
postbicarbonate infusion figures are averaged, the
sea level group presents a mean drop from 3.7
mmoles per L to 3.5 mmoles per L (5.4%), the
normal native group from 3.9 to 3.2 (17.9%),
and the chronic mountain sickness group from
3.5 to 2.9 (17.1%); both altitude groups show a
drop that is of higher magnitude than the sea
level group. Although these figures do not explain
the differences between the bicarbonate Tm of
normal and sick natives, nevertheless they should
be considered as a possible factor in the genesis of
the high Tm found in chronic mountain sickness
patients. Another possibility that should be con-
sidered is the role of anoxia as responsible for the
high Tm. If the kidney responds to severe
anoxia with a primary elevation of Tm, then we
might speculate that the resultant metabolic al-
kalosis could result in compensatory hypoventila-
tion and an elevated arterial Pco₂. This in turn
might aggravate the anoxemia and a vicious circle
would be established. There is no experimental
evidence to support this possibility. Table II and
Figure I show that the normal native usually re-
sponds to the infusion of bicarbonate with an ele-
vation of the arterial Pco₂.

The areas 1', 2', and 3' of Figure I show that
the pH displacement after bicarbonate loading
has been larger in both altitude groups than in
the sea level controls. This does not necessarily
mean that the high altitude groups have a smaller
alkali buffer capacity, since the natives have less
body weight than the sea level men and have a
lower filtration rate, as has been shown by Becker,
Schilling, and Harvey (25). Although we do
not have prebicarbonate-infusion figures, the av-
average postinfusion values for glomerular filtration
rates give figures of 130 ml per m for the sea
level group, 119 ml per m for the high altitude
native group, and 103 ml per m for the natives
with chronic mountain sickness. These factors
might contribute to a greater accumulation of
alkali, since the amount of bicarbonate given was
the same in all the experiments.

Summary

When compared with sea level controls, natives
from high altitudes (4,300 m above sea level) have
lower arterial Pco₂ and the same renal
maximal reabsorption (Tm) of bicarbonate.
Natives with chronic mountain sickness have a
higher arterial Pco₂ than their own native control
group and a higher bicarbonate Tm. The results
are interpreted as indicating that the normal high
altitude native is in a new state of acid-base
equilibrium. The possible roles of high arterial
Pco₂, hypokalemia, and anoxia in the elevation
of bicarbonate Tm of patients with chronic moun-
tain sickness are discussed.
Acknowledgments

We express our gratitude to Professor Robert Pitts and to Dr. George Graham for their reviews of the manuscript and to Mr. José Whitembury, electronic engineer, for his help and advice in the instrumentation at sea level and at high altitudes.

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