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Hypoxia is not the primary mechanism contributing to exercise-induced proteinuria

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ABSTRACT

Introduction Proteinuria increases at altitude and with exercise, potentially as a result of hypoxia. Using urinary alpha-1 acid glycoprotein (α1-AGP) levels as a sensitive marker of proteinuria, we examined the impact of relative hypoxia due to high altitude and blood pressure-lowering medication on post-exercise proteinuria.

Methods Twenty individuals were pair-matched for sex, age and ACE genotype. They completed maximal exercise tests once at sea level and twice at altitude (5035 m). Losartan (100 mg/day; angiotensin-II type 1 (AT1) receptor blocker) and placebo were randomly assigned within each pair 21 days before ascent. The first altitude exercise test was completed within 24–48 hours of arrival (each pair within ~1 hour). Acetazolamide (125 mg two times per day) was administered immediately after this test for 48 hours until the second altitude exercise test.

Results With placebo, post-exercise α1-AGP levels were similar at sea level and altitude. Odds ratio (OR) for increased resting α1-AGP at altitude versus sea level was greater without losartan (2.16 times greater). At altitude, OR for reduced post-exercise α1-AGP (58% lower) was higher with losartan than placebo (2.25 times greater, p=0.059) despite similar pulse oximetry (SpO2) (p=0.95) between groups. Acetazolamide reduced post-exercise proteinuria by approximately threefold (9.3±9.7 vs 3.6±6.0 μg/min; p=0.025) although changes were not correlated (r=−0.10) with significant improvements in SpO2 (69.1%±4.5% vs 75.8%±3.8%; p=0.001).

Discussion Profound systemic hypoxia imposed by altitude does not result in greater post-exercise proteinuria than sea level. Losartan and acetazolamide may attenuate post-exercise proteinuria, however further research is warranted.

INTRODUCTION

Proteinuria typically results from protein leakage from the capillary lumen through the glomerular filter,1 with some removal in the tubes, as shown by studies inhibiting renal tubular reabsorption with lysine infusions.2 Comparisons between albumin, a selectively reabsorbed,3 66 kDa, negatively charged (pI 4.7) protein that passes through the glomerular membrane via the slit diaphragm pores,1 and alpha-1 acid glycoprotein (α1-AGP),5–7 a smaller (41–43 kDa) and more negatively charged protein (pI 2.7),5 7 indicate that urinary α1-AGP is a more sensitive marker of glomerular leakage than albumin.7–9 Urinary α1-AGP excretion has implicated the glomerular origin of the proteinuria exhibited with both, altitude7 and post-exercise.3 Hypothetical mechanisms for such glomerular leak have included changes in renal blood flow,5 10 increases in peritubular pressure and blood pressure2 (BP), hypoxia11 and acid–base disturbances,2 although the mechanisms remain unclear.12 The contributions of BP and hypoxia, especially to post-exercise proteinuria may be uniquely evaluated with altitude exercise.

At altitude, exercise BP is amplified13 14 and such exaggerations are responsive to antihypertensive therapies such as angiotensin II type 1 (AT1) receptor antagonists or blockers (ARBs).13 15 ARBs reduce BP via several mechanisms, including selectively blocking angiotensin II (A-II) from binding to AT1s within the vasculature,16 promoting vasodilation. Specific to the kidney, ARBs
(eg, losartan) ‘block’ vasoconstriction imposed by A-II on afferent arterioles. ARBs also limit both, vasopressin secretion and aldosterone production, further adding to its BP lowering effect. Considering these factors, it would be reasonable to expect altitude post-exercise proteinuria to be greater than at sea level and that AT1 blockers such as losartan could attenuate the post-exercise response. To evaluate this, we planned to compare post-exercise α1-AGP excretion between (1) sea level and altitude exercise and (2) losartan and placebo groups following exercise at altitude.

Exercise oxygen saturation is also altered at altitude and it is profoundly lower compared with sea level but it can be improved by acetazolamide, a carbonic anhydrase inhibitor commonly used to prevent altitude illness. By inhibiting carbonic anhydrase, a catalyst of the reversible CO₂ hydration reaction, acetazolamide promotes carbonic acid formation and dissociation of H⁺ and bicarbonate in the blood which limits hypoxia-induced alkalosis and improves oxygen saturation. Assuming hypoxia as the main mechanism of post-exercise proteinuria, altitude exercise would be expected to produce larger increases in post-exercise proteinuria compared with sea level. In addition, subsequent improvements in arterial oxygenation would be expected to have the ‘reverse’ effect. To evaluate this, we planned to compare post-exercise α1-AGP between (1) sea level and altitude and (2) placebo and placebo +acetazolamide groups at altitude.

We hypothesised that exercise at altitude would increase post-exercise α1-AGP levels compared with sea level, primarily due to greater systemic hypoxia, and that losartan and acetazolamide would attenuate the observed increases by improving blood and peritubular pressures and alleviating renal hypoxia, respectively.

As part of an expedition to Quito, Ecuador, we measured post-exercise urinary α1-AGP excretion in 20 pair-matched individuals once at sea level (before placebo or losartan administration) and twice at altitude as outlined in the following sections.

METHODS

Design and participants

Twenty participants (14 men, 6 women) free of any pre-existing conditions were included in the study. ACE genotyping (II, ID or DD) was performed to limit potential differences between groups that could be attributed to ACE genotype (eg, response to losartan, response to altitude, etc.). Participants were pair-matched participants for ACE genotype, age, sex, previous altitude exposure and glomerular filtration rate (GFR, obtained 4 weeks prior to ascent) (figure 1).

Following matching, a double-blind, randomised, placebo-controlled trial design was adopted. Individuals within each pair were randomly assigned to either placebo or losartan group (figure 1). Losartan administration (100 mg/day or placebo) began in the UK 21 days prior to departure for Quito, Ecuador (2850 m). On arrival at Quito, participants ascended over 10 days to the Whymper Hut on the flank of Chimborazo volcano (5035 m, figure 1).

Baseline and daily measures

Baseline measures of height, body mass, GFR and creatinine (μmol/L) were recorded 4 weeks prior to ascent (figure 1A). Estimated GFR (eGFR) was calculated for each individual using the Modification of Diet in Renal Disease study equation:

\[
eGFR(\text{mL/min/1.73m}^2) = \frac{32788 \times (\text{serum creatinine} \ \mu\text{mol/litre})^{1.154}}{(\text{Age})^{-0.203}} (\text{then 0.742 for females})
\]

During ascent (figure 1B), resting measures of systolic BP (SBP) and diastolic BP (DBP) were collected each morning using a manual sphygmomanometer.

Exercise protocols and measures

Sea-level exercise

Baseline sea-level graded maximal exercise tests were performed 4 weeks prior to ascent on a cycle ergometer (Alticycle) designed for altitude studies (figure 1A). Volitional fatigue was used as the primary end point criterion for the test. Maximal power output (Watt max) was measured by the Alticycle and heart rate (HR) recordings were facilitated by telemetry (Polar Electro, UK).

Altitude exercise

The first altitude exercise tests were commenced on the Whymper Hut for five pairs (day 7) and completed on the following day (day 8) for remaining pairs (figure 1). These initial tests were immediately followed by acetazolamide administration (125 mg orally, two times per day) which occurred 48 hours prior to the second round of altitude exercise tests in all participants. Similar to the first tests, the second round of altitude exercise tests were performed across 2 days (on days 9 and 10, figure 1).

At altitude, pre-exercise measurements of oxygen uptake (VO₂), carbon dioxide production (VCO₂), ventilation (V₆), HR, SBP and DBP were collected. Participants undertook a 5 min self-paced warm-up followed by a modified graded exercise test on the Alticycle which was commenced at 30% of sea-level Watt max. Intensity was increased every 3 min by 10% until the participant reached 80% of sea-level Watt max. From this point, intensity was increased by 10% each minute until volitional fatigue. Expired respiratory gases were analysed breath-by-breath using a Cosmed K4b² (Metabolic Company, Rome, Italy) portable metabolic system alongside continuous measurements of HR (via three-lead ECG), pulse oximetry (SpO₂, Datex Ohmeda 3900, GE Healthcare, USA) and beat-to-beat measurements of SBP and DBP by photoplethysmography (Portapres, Finapres Medical Systems BV, The Netherlands). Change in SBP and DBP was calculated as the difference between pre-exercise and the value obtained at Watt max.
Figure 1  (A) Visual representation of the study design. Step 1: Participants were matched for: ACE genotype (II, ID or DD; see table 3), age, sex previous altitude exposure, and GFR. Step 2: Within each pair, participants were randomly assigned to placebo or losartan groups. Step 3: Baseline characteristics were recorded and baseline exercise tests were conducted (4 weeks before ascent) and were followed by the initiation of losartan administration (21 days before ascent). Step 4: Ascent is initiated with both groups ascending together in accordance with (B). Step 5: The first round of altitude exercise tests were conducted for members of both groups (5035 m). Step 6: Immediately following the first altitude exercise tests, acetazolamide was administered (125 mg orally, two times per day) and continued for 48 hours until next exercise test. Step 7: Repeat altitude exercise tests were conducted for all individuals (only placebo group data reported). (B) Expedition ascent profile. Day 0: Birmingham, UK (130 m), day 1: Quito, Ecuador (2800 m), day 2: Quito, Ecuador (2800 m), day 3: bus to Chunquiragua in Chaupi (3400 m), day 4: bus to Estrella del Chimborazo MARCO cruz (3950 m), day 5: Estrella del Chimborazo MARCO cruz (3950 m) with day hike to 5000 m and back, day 6: bus to Carrel hut (4800 m), days 7–10: Whymper hut (5035 m). GFR, glomerular filtration rate.

Urine collection and storage
Twenty-four-hour urine samples were collected over 1 day at sea level and on each day of the expedition (10 days) with four aliquots (2 mL each) taken from each daily collection and frozen on dry ice.

For pre-exercise collections, participants were instructed to drink 500 mL of water between 90 and 30 min pre-exercise and to provide a timed (60 min) urine collection surrounding exercise as outlined in the following section.
specimen immediately prior to exercise. Post-exercise urine specimens were collected at 60, 120 and 180 min. Four aliquots (2 mL each) were taken from exercise specimens and frozen on dry ice, with the residual volumes returned to each individual’s 24 hours collection bottle. Urine samples were transported back to Birmingham, UK and stored at (−80°C) until analysis.

Urine analysis
All samples were thawed at room temperature for 1 hour before analysis. The 24-hour urine samples were first analysed for α1-AGP using radial immunodiffusion (RID; Talks et al., 2018) and the results were converted to excretion rates (mg/24 hours or μg/min). A subset of samples (119 out of 201) were then analysed using a latex-enhanced immunoassay on the Optilite turbidimetric analyser (The Binding Site, Birmingham, UK; online supplementary appendix 1). This automated method was used for all exercise samples, with the analysed concentrations (mg/L) converted to excretion rates (μg/min) based on sample volumes and collection durations.

Data analysis
Statistical analyses were performed using SPSS (IBM SPSS Statistics, V.25). Normality of distribution was determined by the Shapiro–Wilk test, with data (24-hour urinary α1-AGP excretion) log-transformed where possible. Continuous variables are presented as mean±SD or median ±IQR where appropriate. All tests for significance were two-tailed with statistical significance set at p≤0.05 unless otherwise indicated.

We used independent t-tests to compare group means (placebo vs losartan) where data were normally distributed and when measures were not repeated (eg, age and baseline HR). We used repeated-measures ANOVA with pairwise comparisons (Bonferroni corrected) to determine group (placebo vs losartan) and interaction effects (where appropriate) across days for normally distributed data (eg, transformed 24-hour α1-AGP, DBP and SBP). We used Friedman tests to determine the main effect of time on α1-AGP excretion surrounding exercise (ie, pre-60, post-60, post-120 and post-180 min) where data were not normally distributed. These results are presented as χ²(df), p value. When appropriate, the post-hoc Wilcoxon signed rank test with Bonferroni correction for repeated measures was used to distinguish significance between time points (significance set at p≤0.0125; ie, corrected for three comparisons). We then used Mann–Whitney U tests to compare significant time points between groups (ie, placebo vs losartan at post-60 min).

To avoid any superimposed effects of the two drugs on results, we limited comparisons between the two altitude tests (before and after acetazolamide administration, placebo vs placebo +acetazolamide) to individuals from the placebo group completing both tests (n=9, tables 1 and 2). For these comparisons, we applied the Friedman test and Wilcoxon signed rank test as previously described with additional use of the Wilcoxon signed rank test to compare α1-AGP excretion rates (within-individuals) between placebo and placebo +acetazolamide at post-60 min. We used Spearman’s correlation to evaluate relationships between daily measures (eg, DBP and 24 hours α1-AGP excretion) and exercise measures (eg, SpO₂ and post-exercise α1-AGP excretion).

We calculated ORs for urinary α1-AGP excretion in placebo and losartan groups at rest (pre-exercise and post-exercise at 120 and 180 min) and with exercise (at post-60 min) based on the relative changes (relative increase (+) or decrease (−)) from baseline sea-level measures within each individual.

Patient and public involvement
This study was supported by the Birmingham Medical Research Expeditionary Society which provided input for the conduct of the research. Patients were not included. Public involvement was limited to recruitment. Notification was given to participants at the time of consent that acquisition of personal data was permitted on request. Permission was also obtained at this time for the dissemination of de-identified data within the research team and only externally when a reasonable request was submitted directly to the corresponding author(s) of the present study within 6 months of its publication. A portion of the cohort was invited to review the research methods for accuracy and readability.

Ethics approval
This study did not aim to investigate any safety or efficacy of the already Food and Drug Administration (FDA)-approved drugs included, thus no clinical trial approval was obtained. There were no active FDA recalls for either drug for the duration of the study.

RESULTS
Baseline measures
Baseline data are presented in table 3. Placebo and losartan groups were not significantly different at baseline for measures of age, body mass, height, GFR, eGFR or creatinine. ACE genotype was exactly matched in 9 participants achieved exhaustion during baseline sea-level exercise tests. Baseline measures of HR_max, absolute
Maximal exercise test results compared between groups at baseline and twice at altitude

<table>
<thead>
<tr>
<th>Placebo</th>
<th>Losartan</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>HR\text{\text{max}} (bpm)</td>
<td>171.7±16.5</td>
<td>173.7±22.9</td>
</tr>
<tr>
<td>Absolute Watt\text{\text{max}} (W)</td>
<td>243±66</td>
<td>238±70</td>
</tr>
<tr>
<td>Relative Watt\text{\text{max}} (W)</td>
<td>3.3±0.5</td>
<td>3.1±0.5</td>
</tr>
</tbody>
</table>

Altitude exercise measures of absolute Watt\text{\text{max}}, relative Watt\text{\text{max}}, SBP at Watt\text{\text{max}}, DBP at Watt\text{\text{max}} and HR\text{\text{max}} were similar before and after acetazolamide, as were the changes in SBP. In contrast, changes in DBP were significantly greater with acetazolamide (p=0.04). SpO\text{\text{2}} at Watt\text{\text{max}} was significantly increased with acetazolamide (p<0.01).

Urine studies

Twenty-four-hour excretion

Twenty-four-hour urinary α1-AGP excretion (μg/min) significantly increased with ascent (p<0.01) although no differences were observed between groups for log-transformed excretion rates (p=0.97, figure 2).

Baseline sea level

Baseline urine results are presented in table 2 and figure 3A. Urinary α1-AGP excretion rates were no different between groups before exercise. Exercise increased urinary α1-AGP excretion in both placebo
groups with excretion rates peaking at post-60 min (figure 3A). Urinary α1-AGP excretion at post-60 min was similar between groups (p=0.63) and resolved after 120 min in both groups (figure 3A).

**Altitude (placebo vs losartan)**

Altitude exercise results are presented in table 2 and figure 3B. Collectively, pre-exercise α1-AGP excretion was elevated at altitude compared with sea level, although excretion rates were similar between groups (p=0.62). The odds ratio (OR) for a relative increase in α1-AGP excretion at rest (pre-120, post-120, post-180 min) was 2.16 times greater without losartan at altitude.

Altitude exercise significantly increased urinary α1-AGP excretion in both placebo ($\chi^2(3)=10.73$, p=0.013) and losartan ($\chi^2(3)=15.86$, p<0.01) groups (figure 3B). Post-60 min α1-AGP excretion was lower with losartan compared with placebo, although the difference was not statistically significant (p=0.28, figure 3B and table 2). In the losartan group only, the change in α1-AGP excretion from pre-60 min to post-60 min was lower compared

<table>
<thead>
<tr>
<th>Placebo</th>
<th>P value (Z-score)</th>
<th>Losartan</th>
<th>P value (Z-score)</th>
<th>Placebo +acetazolamide</th>
<th>P value (Z-score)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pre-exercise</td>
<td>3.1±5.3</td>
<td>0.022 (−2.29)</td>
<td>2.4±3.4</td>
<td>0.009 (−2.60)*</td>
<td>4.4±5.7</td>
</tr>
<tr>
<td>ΔPre- to post-60 min</td>
<td>7.9±14.3</td>
<td>–</td>
<td>4.8±9.9</td>
<td>–</td>
<td>1.2±11.03†</td>
</tr>
<tr>
<td>Post-60 min</td>
<td>11.5±19.2</td>
<td>–</td>
<td>8.2±5.9</td>
<td>–</td>
<td>7.7±12.1</td>
</tr>
<tr>
<td>Post-120 min</td>
<td>3.2±2.4</td>
<td>0.007 (−2.70)*</td>
<td>2.8±4.4</td>
<td>0.012 (−2.52)*</td>
<td>3.4±5.0</td>
</tr>
<tr>
<td>Post-180 min</td>
<td>1.3±4.3</td>
<td>0.011 (−2.55)*</td>
<td>1.5±1.5</td>
<td>0.018 (−2.4)</td>
<td>2.7±4.2</td>
</tr>
</tbody>
</table>

Urinary α1-AGP excretion rates (μg/min) are presented as median ±IQR before (pre-exercise, 0 min) and after (post-60, post-120 and post180 min) exercise initiation. Results are presented for placebo versus losartan (baseline), placebo versus losartan (first altitude exercise) and pre-acetazolamide versus acetazolamide (first compared with second altitude exercise).

*Represents the significance of post-60 min α1-AGP excretion (p≤ 0.0125) compared with excretion at other time points.
†Represents the significant (p≤0.05) difference between groups (ie, placebo vs losartan or placebo vs placebo +acetazolamide) at the respective time point. Z-scores are presented in parenthesis where appropriate.

α1-AGP, alpha-1 acid glycoprotein.
with those changes observed at baseline, although this difference was not significant (p=0.059, figure 3D). The OR for reduced urinary α1-AGP excretion at post-60 min (sea level vs first altitude test) was 2.25 times greater with losartan.

Altitude (placebo vs placebo + acetazolamide)
Results for comparisons of post-exercise α1-AGP between placebo and placebo + acetazolamide are presented in tables 1 and 2 and figure 3C,E. As with placebo, altitude exercise tests, placebo + acetazolamide exercise resulted in an increase in urinary α1-AGP excretion (χ²(3)=10.73, p=0.01, figure 3C). Despite similar exercise peak power outputs (table 1) between placebo and placebo + acetazolamide tests, post-exercise α1-AGP excretion (at 60 min post-exercise) was significantly lower following placebo + acetazolamide exercise (p=0.025, figure 3C). Exercise-induced increases in α1-AGP excretion (change from pre-exercise to post-60 min) were significantly reduced (p=0.036) with acetazolamide by nearly threefold (figure 3E). These changes were not correlated (r=-0.10, p=0.82) with the significant improvements in pre-exercise SpO₂ or SpO₂ at Watt max with acetazolamide administration (table 2). No difference was observed between losartan and losartan + acetazolamide tests (data not shown).

**DISCUSSION**

Compared with sea level and despite substantial systemic hypoxia at Watt max, post-exercise α1-AGP excretion was not greater at altitude suggesting that hypoxia is not the primary mechanism. Altitude-related reductions in exercise intensity could, in part, explain this result, but would fail to explain the increased likelihood for post-exercise α1-AGP to be lower with losartan compared with placebo when exercise intensities were similar. We had expected that losartan would mitigate BP amplification and thus lower post-exercise α1-AGP, however we observed no difference in the BP response to exercise between groups. Therefore, we have no evidence to attribute post-exercise α1-AGP responses to alterations in BP or peritubular pressures.

The direct action of ARBs within the glomerular filtration barrier on ATIs of podocytes could provide an alternative explanation. Activation of ATIs on podocytes induces heparanase expression, which promotes the cleavage of heparan sulfate and inhibits the production and secretion of heparan sulfate proteoglycans (HSPGs). The net result is neutralisation of the negatively charged HSPGs which limits the charge-selective function of the glomerular basement membrane (GBM). Blocking ATIs with losartan thus promotes the localised retention of negatively charged proteins at the GBM and limits the charge-selectivity impairment which, in the present study, manifests as a reduction in the extent of post-exercise α1-AGP. This hypothesis is consistent with findings related to the intensity-dependent increases in angiotensin-II and...
exercise-related increases in α1-AGP, although further investigations are required.

Reductions in post-exercise α1-AGP by acetazolamide were unrelated to the significant improvements in exercise SpO₂ at similar intensity (these effects of acetazolamide on exercise performance were previously published and support previous findings). On its own, this would provide further support indicating that hypoxia is not the primary mechanism of post-exercise proteinuria. Unfortunately, the post-exercise effects of acetazolamide cannot be separated from the acclimatisation effect. Thus, no definitive conclusion regarding acetazolamide’s effects on post-exercise proteinuria can be provided. Future research is required to evaluate acetazolamide-related changes in post-exercise proteinuria. Nonetheless, acetazolamide results support recent evidence demonstrating performance limiting effects of acetazolamide despite of its ability to elicit improvements in oxygen saturation.

Limitations and future directions
The inability to control for extraneous variables (eg activity, sleep, environmental conditions) as well as low subject numbers were weaknesses of the present study.
CONCLUSION
Our findings suggest that post-exercise α1-AGP is (1) more related to exercise intensity than degree of hypoxia or BP and (2) possibly influenced by activation or inhibition of AT1 receptors. Losartan may limit post-exercise proteinuria by helping to maintain the charge-selectivity function in the glomerular filter, although further investigations are required for evaluation.

REFERENCES