Adrenal, thyroid and gonadal axes are affected at high altitude

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Abstract

Humans cannot live at very high altitude for reasons, which are not completely understood. Since these reasons are not restricted to cardiorespiratory changes alone, changes in the endocrine system might also be involved. Therefore, hormonal changes during prolonged hypobaric hypoxia were comprehensively assessed to determine effects of altitude and hypoxia on stress, thyroid and gonadal hypothalamus–pituitary hormone axes. Twenty-one male and 19 female participants were examined repetitively during a high-altitude expedition. Cortisol, prolactin, thyroid-stimulating hormone (TSH), fT4 and fT3 and in males follicle-stimulating hormone (FSH), luteinizing hormone (LH) and total testosterone were analysed as well as parameters of hypoxemia, such as SaO2 and paO2, at 550 m (baseline) (n = 40), during ascent at 4844 m (n = 38), 6022 m (n = 31) and 7050 m (n = 13), at 4844 m (n = 29) after acclimatization and after the expedition (n = 38). Correlation analysis of hormone concentrations with oxygen parameters and with altitude revealed statistical association in most cases only with altitude. Adrenal, thyroid and gonadal axes were affected by increasing altitude. Adrenal axis and prolactin were first suppressed at 4844 m and then activated with increasing altitude; thyroid and gonadal axes were directly activated or suppressed respectively with increasing altitude. Acclimatisation at 4844 m led to normalization of adrenal and gonadal but not of thyroid axes. In conclusion, acclimatisation partly leads to a normalization of the adrenal, thyroid and gonadal axes at around 5000 m. However, at higher altitude, endocrine dysregulation is pronounced and might contribute to the physical degradation found at high altitude.

Introduction

Humans have populated a wide variety of climatic and geographical regions of the world. However, altitudes of around 5000 m are only permanently colonized in rare cases, for example mining communities in the Andes. This fact can be explained not only by the hostile climate, but also by the inadequate physiological capacity of the
human organism to adapt to this altitude in the long term or to reproduce in this environment.

A longer-term existence at high altitude requires a variety of cardiorespiratory and metabolic changes (Luks, 1985), a compensated metabolism and, in the case of permanent settlement at higher altitudes, adequate fertility.

This raises the question of whether the endocrine system, which plays a central role in the regulation of almost all bodily functions, becomes significantly dysregulated above a certain altitude.

Several studies have already investigated various hormone concentrations at high altitude (1, 2, 3, 4, 5, 6, 7, 8, 9, 10).

However, the study groups were mostly small, limited to men, were only carried out at altitudes of up to 5000 m above sea level or the study design was not ideal. This might be the reason for the often heterogeneous results of these studies, which have not allowed a conclusive statement about the effect of altitude on the endocrine system to be made up to now.

The aim was therefore to perform an observational study in a large collective, with both male and female subjects at various altitudes up to 7000 m under standardized conditions and to investigate the changes in the adrenal, thyroid and gonadal hypothalamus–pituitary end-organ axes.

Materials and methods

Setting

This was a prospective observational cohort study conducted during the Swiss High Altitude Medical Research Expedition Himlung Himal 2013 (11). The aim of this project was to investigate the cerebral, cardiovascular, pulmonary and humoral adaptation to prolonged hypobaric hypoxia during an ascent to Mount Himlung Himal (7126 m) in Nepal.

Baseline and post-expedition testing was performed in two groups in Berne, Switzerland (550 m) 8 and 9 weeks before the start of the expedition (PR=Pre-expedition). For logistical reasons, the journey as well as the expedition testing were planned in two groups with identical ascent profiles. Comprehensive medical assessments were performed in Base Camp (BC1) at 4844 m on day 7, in Camp 2 (C2) (6022 m) on day 13 and in Camp 3 (C3) (7050 m) on day 23/24 respectively. An additional measurement was carried out at Base Camp on day 19/20 (BC2) to study the effect of acclimatization. A final analysis was performed 4–5 weeks after the expedition at 550 m (PO=Post-expedition).

The number of subjects included in this study was restricted due to logistical and safety concerns. Forty healthy subjects (21 male, 19 female), with basic mountaineering experience and aged between 18 and 70 years, were included in the trial. Exclusion criteria consisted of the presence of any neurologic, cardiac or respiratory disease, diabetes mellitus type I or II or the need for any regular medication, particularly thyroid hormones, corticosteroids and other medication, which could possibly affect hormone analysis. In addition, subjects were excluded if a known susceptibility for high altitude sickness was present. Susceptibility was defined as a history of severe acute mountain sickness, high altitude cerebral oedema (HACE) or high altitude pulmonary oedema (HAPE) after a rapid ascent (<3 nights) at altitudes below 3500 m.

The study was approved by the Cantonal Ethics Committee (KEK 226/12) and has been registered on clinicaltrials.gov (NCT01953198). Consent has been obtained from each patient after full explanation of the purpose and nature of all procedures used.

Measurements

Arterial blood gas sampling and oxygen transport parameters: Arterial blood was aspirated using a blood gas sampler (safe PICO Aspirator, Radiometer Medical, Denmark) and immediate analysis was done with a pocket analyser (Epoc, Alere, Wadenswil, Switzerland) to assess arterial oxygen saturation (SaO$_2$), partial pressure for O$_2$ (paO$_2$) and haemoglobin (Hb).

Blood sampling: At each study site, fasting blood samples were taken between 07.15 and 09.00 h. The samples were collected in sterile serum tubes (Sarstedt, Sevelen, Switzerland). All samples were immediately centrifuged for 10 min at 2000g (EBA 20, Hettich AG, Bäch, Switzerland). Aliquots were frozen to −40° to −60°C on-site and stored at −80°C thereafter.

Cortisol, prolactin, thyroid-stimulating hormone (TSH), free tetraiodothyronine (fT4), free triiodothyronine (FT3), follicle-stimulating hormone (FSH), luteinizing hormone (LH) and total testosterone (testosterone) were analysed. FSH, LH and testosterone were only evaluated in males due to the menstrual-dependent fluctuations in females.

Hormone analysis: Cortisol levels were determined using a competitive immunoassay based on a
Results

Of the 40 subjects, one met an exclusion criterion during the baseline testing and was therefore excluded from the following expedition. Number of subjects was lower at high altitudes but the average age of the analysed subjects was not different. SaO₂, paO₂ and Hb varied significantly at different altitudes (Table 1).

Hormone concentrations were assessed for the effect of hypoxia parameters and altitude. In most cases, only altitude resulted in a significant effect (Tables 2, 3, 4, 5, 6, 7, 8, 9 and 10). Prolactin concentrations were also sex dependent (P=0.015). fT3 (P=0.024) and testosterone (P=0.028) were SaO₂ dependent and TSH (P=0.011) was PoO₂ dependent. Accordingly, in the modelling approach, we adjusted for these parameters while assessing the effect of altitude. Average hormone concentrations stayed within the normal ranges at all altitudes; however, in individual cases concentrations reached pathological concentrations as shown in Figs 1 and 2 for cortisol and fT4.

Hormone axes were affected by altitude as follows:

Adrenal axis and prolactin

Cortisol concentrations revealed a U-shaped pattern with increasing altitude (Fig. 1 and Table 2). Concentrations first decreased significantly from 474 ± 163 nmol/L before the expedition (baseline) to 369 ± 106 at 4844 m and increased 1.7-fold thereafter to reach 643 ± 91 nmol/L at 7050 m altitude. Acclimatization at 4844 m led to normalization of cortisol concentration compared to baseline. These changes were neither sex nor SaO₂ or PaO₂ dependent.

Changes in prolactin (Table 3) concentrations were less pronounced. Prolactin also dropped significantly at 4844 m and increased at 6022 m, but the increase was not significant. However, prolactin also normalized at 4844 m after acclimatization as prolactin concentration

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**Table 1 Basic characteristics of subjects.**

<table>
<thead>
<tr>
<th>Subjects total, n</th>
<th>PR</th>
<th>BC1</th>
<th>C2</th>
<th>BC2</th>
<th>C3</th>
<th>PO</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Subjects total, n</td>
<td>40</td>
<td>38</td>
<td>31</td>
<td>29</td>
<td>13</td>
<td>38</td>
<td></td>
</tr>
<tr>
<td>Females, n</td>
<td>19</td>
<td>17</td>
<td>14</td>
<td>13</td>
<td>5</td>
<td>17</td>
<td></td>
</tr>
<tr>
<td>Males, n</td>
<td>21</td>
<td>21</td>
<td>17</td>
<td>16</td>
<td>8</td>
<td>21</td>
<td></td>
</tr>
</tbody>
</table>

Altitudes: PR, Pre expedition = 550 m; BC1, Base Camp 1 = 4.844 m; C2, Camp2 = 6022 m; BC2, Base Camp 2 (after acclimatization) = 4.844 m; C3, Camp3 = 7050 m; PO, Post expedition = 550 m.

SaO₂, saturation arterial oxygen (%); paO₂, partial pressure of oxygen (mmHg); Hb, haemoglobin (g/L).
Thyroid axis

TSH (Table 4) concentrations were not different compared to baseline apart from 7050 m altitude at which TSH concentration were 1.7-fold higher. In contrast, fT4 (Fig. 2 and Table 5) increased constantly with each altitude from 12.7 ± 1.5 pmol/L before the expedition to reach 16.1 pmol/L at 7050 m. During acclimatization at 4844 m, fT4 concentration decreased but stayed higher than baseline. These changes were less pronounced for fT3 (Table 6). At 4844 m fT3 concentrations also increased significantly, but at 6022 m and 7050 m altitude fT3 concentrations were similar to baseline. The thyroid axis was partly hypoxia dependent. TSH was paO2 dependent.

Gonadal axes

FSH (Table 7) decreased from 5.8 ± 5.9 mU/L before the expedition constantly with increasing altitude to reach 3.3 ± 2.1 mU/L at 7050 m, which corresponds with a 40% decrease. After acclimatization at 4844 m, FSH concentrations normalized compared to baseline. LH concentrations (Table 8) also decreased significantly by around 50% between baseline and 7050 m altitude. However, the decrease could not be observed at all altitudes. At 6022 m, there was an even smaller increase compared to 4844 m. The normalizing effect of acclimatization could also be seen for LH. At 4844 m, LH concentrations were significantly lower compared to baseline, an effect that was not recognizable anymore after acclimatization.

Testosterone concentrations were similar at all altitudes apart from 7050 m altitude. At this altitude, testosterone concentration was – in parallel to reduced concentrations of LH at that altitude – 50% lower compared to baseline. A hypoxia dependency of the
Table 6  fT3 concentrations (pmol/L) at different altitude levels in the whole group of subjects and differences relative to PR.

<table>
<thead>
<tr>
<th>Mean ± s.d.</th>
<th>n</th>
<th>Differences relative to Pr (adjusted for SaO2)</th>
<th>Coefficient ± s.e.</th>
<th>P value</th>
<th>95% CI</th>
</tr>
</thead>
<tbody>
<tr>
<td>PR</td>
<td>4.57 ± 0.60</td>
<td>39</td>
<td></td>
<td>0.69 ± 0.10</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>BC1</td>
<td>5.26 ± 0.73</td>
<td>38</td>
<td></td>
<td>0.02 ± 0.10</td>
<td>0.855</td>
</tr>
<tr>
<td>C2</td>
<td>4.56 ± 0.74</td>
<td>31</td>
<td></td>
<td>0.21 ± 0.14</td>
<td>0.143</td>
</tr>
<tr>
<td>C3</td>
<td>4.76 ± 0.76</td>
<td>13</td>
<td></td>
<td>0.15 ± 0.11</td>
<td>0.169</td>
</tr>
<tr>
<td>BC2</td>
<td>4.74 ± 0.56</td>
<td>29</td>
<td></td>
<td>0.28 ± 0.10</td>
<td>0.004</td>
</tr>
<tr>
<td>PO</td>
<td>4.89 ± 0.61</td>
<td>38</td>
<td></td>
<td>0.02 ± 0.10</td>
<td>-0.585</td>
</tr>
</tbody>
</table>

Variation of fT3 was not sex but SaO2 (P=0.018) and paO2 (P=0.024) dependent.

Table 8  LH concentrations (mU/L) at different altitude levels in the whole group of subjects and differences relative to PR.

<table>
<thead>
<tr>
<th>Mean ± s.d.</th>
<th>n</th>
<th>Differences relative to Pr</th>
<th>Coefficient ± s.e.</th>
<th>P value</th>
<th>95% CI</th>
</tr>
</thead>
<tbody>
<tr>
<td>PR</td>
<td>3.81 ± 1.98</td>
<td>18</td>
<td></td>
<td>-1.57 ± 0.36</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>BC1</td>
<td>2.39 ± 1.55</td>
<td>19</td>
<td></td>
<td>-1.45 ± 0.49</td>
<td>0.003</td>
</tr>
<tr>
<td>C2</td>
<td>3.00 ± 1.52</td>
<td>18</td>
<td></td>
<td>-1.85 ± 0.54</td>
<td>0.001</td>
</tr>
<tr>
<td>C3</td>
<td>1.88 ± 1.25</td>
<td>6</td>
<td></td>
<td>0.79 ± 0.39</td>
<td>0.044</td>
</tr>
<tr>
<td>BC2</td>
<td>3.26 ± 1.87</td>
<td>14</td>
<td></td>
<td>-0.34 ± 0.37</td>
<td>0.346</td>
</tr>
<tr>
<td>PO</td>
<td>3.60 ± 1.59</td>
<td>19</td>
<td></td>
<td>-0.02 ± 0.10</td>
<td>0.66</td>
</tr>
</tbody>
</table>

Variation of LH was not sex, SaO2 and paO2 dependent.

Table 9  Testosterone concentrations (nmol/L) at different altitude levels in the whole group of subjects and differences relative to PR.

<table>
<thead>
<tr>
<th>Mean ± s.d.</th>
<th>n</th>
<th>Differences relative to Pr (adjusted for SaO2)</th>
<th>Coefficient ± s.e.</th>
<th>P value</th>
<th>95% CI</th>
</tr>
</thead>
<tbody>
<tr>
<td>PR</td>
<td>21.5 ± 10.1</td>
<td>19</td>
<td></td>
<td>0.10 ± 0.44</td>
<td>0.011</td>
</tr>
<tr>
<td>BC1</td>
<td>21.1 ± 13.6</td>
<td>19</td>
<td></td>
<td>-1.66 ± 0.62</td>
<td>0.007</td>
</tr>
<tr>
<td>C2</td>
<td>23.9 ± 13.5</td>
<td>19</td>
<td></td>
<td>-0.66 ± 0.48</td>
<td>0.167</td>
</tr>
<tr>
<td>C3</td>
<td>10.7 ± 3.2</td>
<td>6</td>
<td></td>
<td>0.16 ± 0.37</td>
<td>0.712</td>
</tr>
</tbody>
</table>

Variation of testosterone was not sex or paO2 but SaO2 (P=0.28) dependent.

In summary, adrenal, thyroid and gonadal axes as well as prolactin secretion were affected by increasing altitude. Adrenal axis and prolactin were first suppressed at 4844 m and then activated with increasing altitude; thyroid axis and gonadal axis were both directly activated respectively suppressed with increasing altitude. Acclimatisation at 4844 m led to normalization of adrenal and gonadal but not of thyroid axes.

Discussion

Our study revealed that adrenal and thyroid axes as well as prolactin secretion are activated, and male gonadal axis is suppressed at high altitude. The study also revealed that acclimatisation at 4844 m led to normalization of adrenal and gonadal but not of thyroid axes.

Variation of fT3 was not sex but SaO2 (P=0.018) and PaO2 (P=0.024) dependent.

Previous studies have already analysed endocrine parameters at different altitudes but revealed heterogeneous or even controversial results, which might be due to the small study sizes or suboptimal study designs.

As we performed a study with a large study group with both male and female participants, with an analysis at various altitude levels up to 7050 m and under study conditions, which were standardized as much as possible, we assume that our study results are quite representative. However, our study design could not exclude all confounders, which a field study is confronted with. We therefore tried to include several hormone parameters for each analysed hormone axis to get a more representative and more comprehensive picture of the endocrine system.

The reason for our study was not only to better understand what kind of metabolic changes and adaptions mountaineers face at high altitude expeditions, we also hoped to understand if the endocrine systems contribute to the incapability of humans to live above around 5000 m altitude.
It is well known that hypobaric hypoxia leads to several cardiorespiratory and metabolic changes (12), which preserve sufficient performance at higher altitude. This is impressively demonstrated by the ability of humans to ascend to over 7000 m above sea level without additional oxygen. However, staying at this altitude is only possible for a brief period of time. The critical altitude for living permanently is usually about 5000 m, although this limit may vary depending on individual disposition and population group. The reasons for the limited ability of humans to live at a high altitude are not clear. It is known that a life-threatening form of altitude sickness can develop, which leads to physical degeneration and deterioration and is fatal. However, it is unclear what controls the metabolic changes at high altitude. This raises the question of whether these processes are also under endocrinological control.

Our study provides evidence for this hypothesis, as the adrenal axis is activated at high altitude, but can at least partially normalize at an altitude of about 5000 m. The strong activation of the adrenal axis as well as the thyroid axis, and inhibition of the gonadal axes at very high altitude suggest that adaptation might not be longer possible at this altitude, and thus, might contribute to the physical degradation. However, it needs to be stressed that the endocrine changes might only reflect the adaptation of the body without significantly affecting the physical status at high altitude. The dysregulation of the endocrine system at high altitude is only mild to moderate and the endocrine system might to some extent adapt to very high altitude.

Interestingly, the effects are not significantly different in men and women. We found similar effects in the adrenal and thyroid axes in both women and men. The gonadal axis could only be studied in men due to the

Table 10 Pairwise comparisons of hormones shown in Tables 2, 3, 4, 5, 6, 7, 8 and 9 (P values, bold <0.05).

<table>
<thead>
<tr>
<th>Overall effect</th>
<th>Cortisol female and male</th>
<th>Prolactin female and male</th>
<th>TSH female and male</th>
<th>fT3 female and male</th>
<th>fT3 female and male</th>
<th>FSH male</th>
<th>LH male</th>
<th>Testosterone male</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ascent</td>
<td>&lt;0.001</td>
<td>0.054</td>
<td>&lt;0.001</td>
<td>&lt;0.001</td>
<td>&lt;0.001</td>
<td>&lt;0.001</td>
<td>&lt;0.001</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>BC1 vs PR</td>
<td>0.001</td>
<td>0.016</td>
<td>0.240</td>
<td>&lt;0.001</td>
<td>&lt;0.001</td>
<td>0.159</td>
<td>&lt;0.001</td>
<td>0.751</td>
</tr>
<tr>
<td>C2 vs PR</td>
<td>0.971</td>
<td>0.953</td>
<td>0.594</td>
<td>&lt;0.001</td>
<td>0.096</td>
<td>&lt;0.001</td>
<td>0.046</td>
<td>0.271</td>
</tr>
<tr>
<td>C3 vs PR</td>
<td>0.015</td>
<td>0.464</td>
<td>0.001</td>
<td>&lt;0.001</td>
<td>&lt;0.001</td>
<td>0.100</td>
<td>0.009</td>
<td>1.000</td>
</tr>
<tr>
<td>C2 vs BC</td>
<td>0.001</td>
<td>0.994</td>
<td>1.000</td>
<td>1.000</td>
<td>0.967</td>
<td>0.393</td>
<td>1.000</td>
<td>0.387</td>
</tr>
<tr>
<td>C3 vs BC1</td>
<td>&lt;0.001</td>
<td>1.000</td>
<td>&lt;0.001</td>
<td>0.009</td>
<td>1.000</td>
<td>0.999</td>
<td>1.000</td>
<td>0.997</td>
</tr>
<tr>
<td>C3 vs C2</td>
<td>0.484</td>
<td>0.999</td>
<td>&lt;0.001</td>
<td>0.076</td>
<td>0.864</td>
<td>1.000</td>
<td>1.000</td>
<td>0.018</td>
</tr>
<tr>
<td>Descent</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>PO vs C3</td>
<td>0.008</td>
<td>0.823</td>
<td>&lt;0.001</td>
<td>&lt;0.001</td>
<td>0.122</td>
<td>0.046</td>
<td>0.077</td>
<td>1.000</td>
</tr>
<tr>
<td>PO vs C2</td>
<td>0.887</td>
<td>1.000</td>
<td>0.008</td>
<td>&lt;0.001</td>
<td>0.652</td>
<td>&lt;0.001</td>
<td>0.275</td>
<td>0.104</td>
</tr>
<tr>
<td>PO vs BC2</td>
<td>0.536</td>
<td>0.940</td>
<td>0.980</td>
<td>&lt;0.001</td>
<td>1.000</td>
<td>0.746</td>
<td>0.985</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Acclimatization</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>BC1 vs PR</td>
<td>0.001</td>
<td>0.016</td>
<td>0.240</td>
<td>&lt;0.001</td>
<td>&lt;0.001</td>
<td>0.159</td>
<td>&lt;0.001</td>
<td>0.751</td>
</tr>
<tr>
<td>BC2 vs PR</td>
<td>0.079</td>
<td>0.451</td>
<td>0.074</td>
<td>&lt;0.001</td>
<td>0.161</td>
<td>0.936</td>
<td>0.492</td>
<td>0.006</td>
</tr>
<tr>
<td>BC2 vs BC1</td>
<td>&lt;0.001</td>
<td>0.999</td>
<td>0.017</td>
<td>&lt;0.009</td>
<td>&lt;0.001</td>
<td>0.709</td>
<td>0.550</td>
<td>0.839</td>
</tr>
<tr>
<td>Long term effect</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>PO vs PR</td>
<td>1.000</td>
<td>1.000</td>
<td>0.012</td>
<td>0.945</td>
<td>0.059</td>
<td>1.000</td>
<td>0.998</td>
<td>0.991</td>
</tr>
</tbody>
</table>

Data are adjusted for sex (Prolactin), SaO2 (fT3, Testosterone) and paO2 (TSH). Sidak adjustment for multiple comparisons was considered.

Figure 1 Cortisol concentration (nmol/L), representing a stress axis. Effect of altitude at 550 m before the expedition (PR), during ascent at 4.844 m (BC1), 6.022 m (C2) and 7.050 m altitude (C3), during descent at 4.822 m after acclimatization (BC2) and after the expedition at 550 m (PO). Boxes represent the 25th and 75th percentiles, and the bottom and top whiskers are defined by the 25th percentile minus 1.5 times the interquartile range (IQR) and the 75th percentile plus 1.5 times the IQR, respectively. Data points outside this range are plotted as individual points. Horizontal lines represent normal values. Statistical comparisons are shown in Tables 2 and 10.
cycle-related fluctuations in female gonadotrophins, which is a limitation of the study. It can be assumed, however, that suppression of the gonadal axis also occurs in women at high altitude, since the female gonadal axis is very sensitive to stressors and reacts with a dysregulated hypothalamic–pituitary pulsatility (13). We had also analysed the pulsatility of the hormone secretion to identify very subtle changes of the endocrine regulation, which will be published in a different paper.

Previous studies have already investigated the effect of hypobaric hypoxia on the endocrine system.

Cortisol concentrations appear to increase with increasing altitude, according to previous studies. Woods et al. showed a significant increase in resting cortisol concentrations at approximately 5000 m above sea level (8). Our study confirmed cortisol increase at high altitudes in both women and men. However, our study at 4844 m initially showed a cortisol decrease and a cortisol increase only at higher altitudes. Woods et al. also described lower cortisol levels below 5000 m (8). They found non-significantly lower concentrations at 4270 m compared to 5150 m. Significantly lower concentrations were noted when exercise (6-h trek) took place before the measurement.

Prolactin concentrations were previously found to first decrease in men at 5380 m, but then increase significantly if they stay at this altitude for several months (10). A significant increase in prolactin was also found in nine mountaineers in whom baseline prolactin levels were measured directly after climbing Mount Everest (5). Our study confirmed lower prolactin concentrations at 4844 m and elevated concentrations at 6022 m altitude. However, the increase was not significant.

Thyroid function dysregulations are not clear according to previous studies. Nepal et al. compared thyroid function in indigenous people at 2800 and 3750 m (9). fT4 was significantly elevated at 3750 m, but not fT3 and TSH. Basu et al. found significantly higher fT3 and fT4 concentrations in soldiers after acclimatization at an altitude of about 6300 m (3). Hackney et al. found significantly lower TSH and fT3 concentrations in 15 mountaineers who had climbed Denali in Alaska (6194 m), whereas fT4 concentrations remained unchanged (2). In our study, the most striking differences were found for fT4, which increased with altitude. Results were not clear for fT3 and TSH, but overall, the thyroid axis was activated. As the thyroid gland releases mainly T4, which is metabolized to T3, and as T4 has a much longer half-life than T3, it can be speculated that the altitude adaptations of thyroid function are easier to be detected using T4 levels. TSH might also not be an ideal parameter for the detection of altitude-dependent functional changes as an increase in TSH may barely be detectable due to the negative feedback loop of T3 and T4 (14). Although a comparison of the studies is only possible to a limited extent and the data are somewhat heterogeneous, there appears to be an overall activation of the thyroid with increasing altitude, which is not detected at a pituitary level but in the thyroid gland.

Gonadal axes have previously poorly been studied. In the study by Benso et al., significantly reduced testosterone concentrations were found in eight mountaineers in the Base Camp at 5200 m directly after climbing Mount Everest (5); however, the interpretation of these results is limited due to the small number of test persons and excessive exertion. We found suppression of FSH and partly in LH with increasing altitude. Testosterone did not change very much but interestingly both LH and testosterone were reduced at 4800 m. As LH stimulation regulates the production of testosterone (15), it can be assumed that low testosterone concentrations are a result of suppressed LH.

Acclimatization and its effects have not yet been investigated in other studies with a suitable study design. Only the effect of several months’ stay at 5380 m (10) has been analysed so far. LH and total testosterone concentrations decreased significantly in 52 soldiers after 6 months at this altitude. The hormone levels increased again within the next 6 months. Sperm concentration, motility and vitality were also significantly reduced after 6 months at this altitude. Sperm concentration improved over the following 6 months, but not motility. After returning to lower altitudes, semen analysis normalized.
again. The decrease of semen parameters at high altitude was potentially relevant for fertility. There was a moderate asthenozoospermia at altitude, corresponding with andrological subfertility. Six months after starting the altitude exposure, the overall motility was on average about 35% and after 12 months about 28%. In our study, normalization of the stress axes in both sexes as well as the male gonadal axis after acclimatization was also demonstrated. We were able to show that normalization begins after just a few days.

All these study results raise the question of what causes these changes. Interestingly, the hormonal changes hardly correlated with parameters of oxygen transport such as O₂ saturation and pO₂ values, but only with altitude. Consequently, high-altitude hypoxia may not exclusively cause these changes, and other factors must also have an influence. The cold that occurs at high altitudes is probably not relevant or only of limited relevance because, due to good clothing, any relevant exposure to cold is mainly achieved via respiratory air. An effect of the increased physical burden at altitude due to hypobaric hypoxia is conceivable. 1, 8 demonstrated significantly higher cortisol levels or prolactin levels after physical exercise at an altitude of approximately 5000m. The effect of disturbed sleep due to periodic breathing at high altitudes would also be conceivable in terms of the change in stress axes (16). Insomnia and shortened sleep leads to significantly higher cortisol levels, even at low altitudes (17, 18). However, sleep does not improve with increasing acclimatization (16), and disturbed sleep at high altitude cannot fully explain the endocrine changes, as these, in contrast to sleep, partially normalize at 4800m. Because of this, it can be assumed that several height-dependent factors such as hypoxia, cold, physical stress, disturbed sleep, etc. have an influence and accumulate.

The question also arises if these endocrine changes might have a substantial effect on the health of humans living at high altitude for a long time. Insomnia leads to increased cortisol concentrations and to increased blood pressure and the risk for cardiovascular diseases (19). In line with this, patients with adverse cardiovascular events had higher median levels of cortisol (609.4 vs 594.4 pmol/mL, P<0.001) that those without (20). Clinical and subclinical hyperthyroidism is associated with different kinds of cardiovascular diseases (21). These effects might clinically be relevant as even a mild increase of blood pressure has an impact on mortality (22). The clinical consequences of high altitude on the gonadal axis has already been revealed by the above described studies on soldiers (10) which showed a clinically relevant reduction in sperm quality above about 5000m.

Therefore, in our view, the hypothesis is acceptable that the endocrine system might play a role in the fact that human beings cannot live long term at very high altitude.

In summary, the investigations show an activation of the adrenal and thyroid axes and inhibition of the male reproductive endocrine axis, which partially normalize at least at altitude of about 5000m. The endocrine changes found at higher altitude might contribute to the physical degradation and thereby to the incapability of humans to live permanently at very high altitude.

Declaration of interest
The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

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Author contribution statement
M v W and J P H designed the study, participated in the expedition, analysed the data and prepared the manuscript. M T prepared the samples and performed the analysis. C T N performed the statistical analysis. J D V provided support in the design of the study. T M M provided the vote of the cantonal ethics committee, organized and participated in the expedition. A R H performed the hormone analysis. All authors revised the final manuscript.

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