Periodic breathing in healthy young adults in normobaric hypoxia equivalent to 3500 m, 4500 m, and 5500 m altitude

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Abstract
Purpose The occurrence of periodic breathing (PB) at high altitude during sleep and the quality of sleep are individually different and influenced by multiple factors including sex. Although poor sleep quality at high altitude might not be directly linked to oxygen desaturations, the PB upsurge at high altitude leads to significant oscillations in oxygen saturation.

Methods Thirty-three students were recruited. Participants were randomly assigned to three groups (A, B, C) sleeping one full night in a dormitory with normobaric hypoxia at a FIO2 of 14.29% (A), a FIO2 of 12.47% (B), or a FIO2 of 10.82% (C). Full polysomnography was performed in each participant.

Results Mean total sleeping time decreased significantly with increasing hypoxia ($p < 0.001$). Respiratory events changed from central hypopneas to central apneas (CA) with increasing hypoxia: CA = 17.8%, 50.0%, 92.2% of AHI (37.96 events per hour (n/h), 68.55 n/h, 93.44 n/h). AHI ($p = 0.014$) and time duration of respiratory events ($p = 0.003$) were significantly different between sexes, both greater in men. REM sleep was reduced.

Conclusions Men tend to be more prone to PB in normobaric hypoxia. Further research should implicate a longer acclimatization period around simulated 4500 m in order to find out if the exponential increase in PB between 4500 m and 5500 m could be shifted to lower hypoxic levels, i.e., higher altitudes.

Keywords Hypoxia · Periodic breathing · Sleep · Polysomnography · Altitude
instability in men seems to be driving increased PB at altitude [11–13]. This sex difference has also been reported to be based on hormonal mechanisms directly and indirectly contributing to ventilatory control and central breathing stimulation in subjects at altitude, however, these data are still controversially discussed [8–14]. The understanding of PB in normobaric hypoxia in different simulated altitudes could add to a better understanding of the underlying mechanisms and sex-related differences to hypoxia. Therefore, our aim was to study differences in PB in normobaric hypoxia equivalent to 3500 m, 4500 m, and 5500 m (FIO2 = 14.29%, 12.47%, 10.82%). We hypothesized that there might be an inspired oxygen fraction (FIO2)-dependent upsurge for PB and that there are sex-related differences in the occurrence and degree of PB.

Materials and methods

Subjects

Thirty-three healthy students have been recruited from the University of Ulm and the University of Innsbruck and gave written informed consent. Inclusion criteria were the absence of pre-diagnosed sleeping disorders and an overall good health status as non-smoker. All students were questioned a priori by a physician to assure health status and exclude pregnancy. According to the physician, none of the students was overweight indicating a BMI between 18.5 and 24.9 kg/m2. All students were of German or Austrian ethnicity. Preexisting sleep disorders have been excluded via BERLIN questionnaire. We evaluated 19 male and 14 female subjects with a mean age of 23.36 ± 2.52 years (Table 1). All participants were randomly assigned into three groups. Group A contained 5 male and 6 female subjects (age 24.45 ± 1.75 years), group B 8 male and 3 female subjects (age 24.64 ± 1.91 years), and group C 6 male and 5 female subjects (age 21.00 ± 2.10 years) (Table 1). They had comparable daytime routines since they followed the same course program containing lectures and physical activity for the past semester as well as hiking at altitudes < 2000 m the days before the measurements. Previous exposition to moderate and high altitudes during the last 2 weeks prior to study start could be excluded for all subjects. Group characteristics are displayed in Table 1.

Procedure

The three groups were assigned to three different normobaric altitude simulations. Group A at a FIO2 of 14.29%, Group B at a simulated FIO2 of 12.47% and Group C at a FIO2 of 10.82%. According to the assigned group, the corresponding altitudes were Group A at 3500 m, Group B at 4500 m, and Group C at 5500 m. Group B and C underwent an acclimatization night at a FIO2 of 14.29% (equivalent to 3500 m) the night before the actual testing night in order to prevent symptoms of acute mountain sickness (AMS). The trial took place in the normobaric altitude sleeping room of the Hermann Buhl Institute for Hypoxia and Sleep Medicine Research. Normobaric hypoxia was provoked by an oxygen expulsion System (normobaric hypoxia, low oxygen systems; Berlin-Buch, Germany). This allows reducing oxygen in the whole chamber down to a minimum of 9.3%. The participants’ medical history was assessed by an experienced physician. Two students at a time were connected to a 12-channel PSG each study day (Sidas, Stimotron Inc., Roth, Germany). Polysomnography was carried out and scored by a sleep physician according to the American Academy of Sleep Medicine Standard of 2017 [15]. Monitoring time was 11:00 pm until 06:30 am. During this period, following data was collected continuously: heart rate (HR), EMG, EEG, EKG, EOG, peripheral oxygen saturation (SpO2), nasal air flow (NAF), position, and abdominal and thoracic movement. The assessment of AMS symptoms was self-administered by each participant upon awakening using the Lake Louise Score. If subjects experienced symptoms of AMS higher than 6 on the Lake Louise Score, they were allowed to interrupt measurements and leave the hypoxic room.

Statistical analysis

Data are presented as means ± standard deviation (SD). Data analyses were performed with the SPSS statistical software package (PASW Statistics for Windows version 21.0, SPSS Inc., Chicago, IL, USA). Normal distribution of data has been tested via the Shapiro-Wilk test and has been visually checked. A multifactor ANOVA was applied to identify differences between altitudes and sex. Significance level was set at p < 0.05. Post-hoc power calculation via G-power for sex differences in the main parameter AHI gave a power of 0.97.

Table 1 Group characteristics. Values are presented as means ± SD

| FIO2 (%) | Simulated altitude (m) | n   | Sex (m/f) | Age (years) |
|---------|------------------------|-----|-----------|-------------|
| Group A | 14.29 | 3500 | 11 5/6 | 24.45 (± 1.75) |
| Group B | 12.47 | 4500 | 11 8/3 | 24.64 (± 1.91) |
| Group C | 10.82 | 5500 | 11 6/5 | 21.00 (± 2.10) |

FIO2, inspired oxygen fraction
Results

The measurements and altitude simulations at a FIO2 of 14.29% (equivalent to 3500 m) and 12.47% (equivalent to 4500 m) were well tolerated. None of the participants had to leave the hypoxic room due to moderate AMS symptoms (Lake Louise Score > 6). At the measurements at a FIO2 of 10.82% (equivalent to 5500 m) almost all participants were suffering from moderate AMS (Lake Louise Score > 6) symptoms and left the hypoxic room after 4 to 7 h. We observed a highly significant decrease in total sleeping time (p < 0.001) and peripheral oxygen saturation (p < 0.001) with decreasing FIO2 considering all subjects (Table 2). The heart rate was slightly increased with greater simulated altitudes but showed no significance considering all subjects (p = 0.100) (Fig. 1). We could measure a significant shift from mostly hypopneas towards apneas at a FIO2 of 12.47% (4500 m) and at a FIO2 of 10.82% (5500 m) (p = 0.001, p = 0.032) with an increase of total respiratory events, but found no changes in duration of the events (p = 0.527). There was a significant increase in mean AHI (p = 0.017). (Fig. 2) The arousal frequency did not change from a FIO2 of 14.29% (3500 m) up to a FIO2 of 12.47% (4500 m) in all participants and at a FIO2 of 10.82% (5500 m) only one subject expressed arousals (87/h) by the classic definition of short (few seconds) alpha rhythm in the EEG (Fig. 1). All other subjects had no arousals during sleep phases but only full awakenings from respiratory disturbances. REM sleep was low, compared to normal, at all hypoxia levels (7.45%, 10.8%, 5.8% of TST) with no difference between sexes [16]. Significant sex differences were seen in AHI and event duration parameters. Female subjects showed a lower AHI (p = 0.014) as well as a shorter event duration (p = 0.003) in all hypoxic conditions. No significant sex differences could be seen in SpO2 or TST during all hypoxic conditions (Fig. 3).

Discussion

To our knowledge, this is the first study to assess PB in normobaric hypoxia at different simulated altitudes. The low impact of disruptive factors using normobaric hypoxia compared to hypobaric chambers like in the Operation Everest II in Loma Linda (California) or real altitude seems to be unique [17]. Due to our findings, the occurrence of PB increases with decreasing FIO2 first in a more linear matter and then with an exponential upsurge between a FIO2 of 12.47% (4500 m) and a FIO2 of 10.82% (5500 m). Due to the signal chain of the carotidal chemoreceptors and the delay of the feedback response, PB patterns are more pronounced in hypoxic environments in an hypoxic environment of a FIO2 of 10.82% (5500 m). This applies more to non-acclimatized subjects. PB has been thought by some colleagues to have a stabilizing effect on oxygen saturation and to be an acute adaption to the demanding situation of hypoxia [5, 18–20]. Our data supports this theory since PB is increased reaching critical altitudes. However, in our sample, we could not establish a correlation between AHI and SpO2. Considering the prevalence of males in the hypoxic condition with an FIO2 of 12.47% (4500 m) we cannot exclude an effect on the mean AHI, given the small sample size. However, we report a homogenous subject group concerning age, as the age difference was not significant between groups. Therefore, the anthropometric age data should have not influenced measured parameters.

TST was significantly reduced with greater simulated altitudes. The disturbance due to the lower FIO2 seems to impact sleep severely and does not allow longer REM periods [21, 22]. If this is due to the low oxygen levels alone or co-affected by the mechanical disturbance due to PB remains to be investigated [21]. Surprisingly in our experiment and in opposition to previous studies like Operation Everest II, arousals did not play a predominant role [17]. Although there could be observed a slight increase in arousal frequency from a FIO2 of 14.29% (3500 m) up to a FIO2 of 12.47% (4500 m), subjects either awoke or slept without arousals at a FIO2 of 10.82% (5500 m) with few arousals in their short sleep periods. This opposing finding could be due to less external disruptive factors in the normobaric hypoxia room, where sleep was not disturbed by compressors. More studies on this matter are needed.

Consistent with other studies, the shift from mostly hypopneas to apneas was quite significant. The lower oxygen levels seem to amplify respiratory responses and lead to a

| FIO2 (%) | TST (min) | nREM(%) | REM(%) | SpO2(%) minimum | SpO2 (below 90%) | SpO2 (below 80%) |
|---------|----------|---------|--------|-----------------|-----------------|-----------------|
| 14.29   | 430 (± 51.56) | 92.91 (± 5.61) | 7.36 (± 6.07) | 66.0 (± 10.76) | 79.91 (± 28.70) | 6.64 (± 13.77)   |
| 12.47   | 340 (± 83.28) | 90.36 (± 9.30) | 10.82 (± 9.24) | 56.82 (± 9.81) | 98.27 (± 2.24)  | 58.0 (± 30.90)   |
| 10.82   | 188 (± 118.20) | 98.41 (± 13.88) | 5.96 (± 5.64) | 55.60 (± 4.03) | 99.73 (± 0.85)  | 97.76 (± 4.07)   |

FIO2, inspired oxygen fraction in %; TST, total sleep time in minutes; nREM/REM, (non) rapid eye movement sleep in % of TST; SpO2, oxygen saturation values in % of TST.
Fig. 1 General polysomnographic data recorded at 3 different levels of normobaric hypoxia at an F IO2 of 14.29% (3500 m), 12.47% (4500 m), and 10.82% (5500 m). Values are presented as means ± SD. Legend: TST, total sleeping time, HF, heart frequency, SpO2, peripheral oxygen saturation; arousals: events per hour (n/h) * = level of significance, \( p \leq 0.001 \) (in regard to different hypoxic conditions)

Fig. 2 Respiratory parameters from polysomnographic data recorded at 3 different levels of normobaric hypoxia at an F IO2 of 14.29% (3500 m), 12.47% (4500 m), and 10.82% (5500 m). Values are presented as means ± SD. CA, central apneas; CH, central hypopneas; AHI, apnea hypopnea index; n/h = events per hour; levels of significance, * = \( p \leq 0.05 \); *** = \( p \leq 0.001 \)(in regard to different hypoxic conditions)
higher oscillation. The duration of respiratory events did not seem to be affected by lower F IO2 levels. According to Orr et al. 2017, the duration of high altitude provoked PB events remains quite stable at approximately 10 s [23]. PB was more pronounced in male subjects which could indicate higher instability of the carotidal chemoreceptors in men [24]. Furthermore, male subjects seem to show a longer event duration regardless to the dose of hypoxia compared to women. This could be due to higher lung volumes and the slower breathing frequency male subjects show in general and therefore a slower responsiveness to carotidal signals [25]. The sex difference could also be related to the effect of sex hormones directly and indirectly affecting respiration and ventilation mechanisms as well as cerebral blood flow regulations [14]. Hormones such as estrogens and androgens influence cerebral blood circulation, which in turn affects central chemoreflex activity [26]. During normal menstrual cycle, estrogens, androgens, and testosterone take action in the central neural control of breathing, which affects cyclic fluctuations in ventilation. The effect of cerebral blood flow exerted by female hormones might contribute to improve the stability of ventilator control [27]. Additional information on the menstrual cycle phase of our female subjects as well as application of contraceptives could have supported this theory and should be assessed in further investigations. Usually, very few obstructive events are registered in healthy subjects at sea level, however, we could not detect any obstructive events in our subjects at altitude. We assume that at the studied altitudes, the frequent periodic breathing might have masked few obstructive events.

One main limitation of this study is the lack of baseline polysomnography at sea level which would have provided us with individual starting points regarding respiratory parameters. We tried to address this shortcoming with a detailed assessment of the subjects’ medical history excluding any cardiorespiratory conditions, which might influence respiration at sea level and non-hypoxia induced PB-related events at altitude. Although the number of incidences of preexisting sleep disorders in young subjects is rising, we assumed that the prevalence of preexisting sleep disorders in adults of age is still higher [28]. Therefore, we would anticipate a rather low incidence of preexisting sleep disorders in our study population. Due to the fact, that we aimed at the assessment of acute hypoxia exposure, we did not have a familiarization period or acclimatization nights at conditions of a F IO2 of 14.29% (3500 m) and a F IO2 of 12.47% (4500 m). Furthermore, some subjects interrupted sleep at a F IO2 of 10.82% (5500 m) because of AMS symptoms. This might have led to shorter TST.

In conclusion, our findings indicate that from a F IO2 of 12.47% (4500 m) up to a F IO2 of 10.82% (5500 m) PB increases exponentially. This is of importance to know for mountaineers and other persons reaching very high altitudes for recreational or professional purposes because the exponential
increase of periodic breathing might have significant impact on health and wellbeing. It is very likely that PB has a SpO2 stabilizing effect and is a necessary adaptive response to hypoxia. Considering the fact, that men tend to be more prone to PB than women in normobaric hypoxia, we assume that female sex hormones regulating the menstrual cycle also contribute to improve nightly ventilator control stability in hypoxia. Further investigations examining the hormonal threshold during sleep while exposed to hypoxia concerning PB are suggested. Overall, possible differences of normobaric (NH) and hypobaric (HH) hypoxia regarding ventilator response during sleep have to be taken into account [29]. HH could induce lower nocturnal oxygen saturation values and more AHI compared to NH [19]. The main difference could lie in NO metabolism altering pulmonary capillary vasodilation or an increase of physiological dead space [30, 31]. These hypotheses will need to be confirmed in further studies. However, the occurrence of PB appears to show similar metrics in both, NH and HH, which still leaves normobaric hypoxia a valuable tool for further investigations [19].

Further research should implicate a longer acclimatization period around a FIO2 of 12.47% (4500 m) in order to find out if the exponential increase in PB between a FIO2 of 12.47% (4500 m) and a FIO2 of 10.82% (5500 m) could be shifted to lower hypoxic levels, i.e., higher altitudes.

Author contributions S. P.: conception or design of the work; or acquisition, analysis, or interpretation of data for the work; drafting the work or revising it critically for important intellectual content. R. S.: acquisition, analysis, or interpretation of data for the work; drafting the work or revising it critically for important intellectual content. A. P.: acquisition, analysis, or interpretation of data for the work; drafting the work or revising it critically for important intellectual content. L. R.: acquisition, analysis, or interpretation of data for the work; drafting the work or revising it critically for important intellectual content. N. C. N.: conception or design of the work; acquisition, analysis, or interpretation of data for the work; drafting the work or revising it critically for important intellectual content.

All authors: approved the final version of the manuscript and agree to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved and all persons designated as authors qualify for authorship, and all those who qualify for authorship are listed.

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Compliance with ethical standards

Conflict of interest Stephan Pramsohler declares that he has no conflict of interest. Robert Schilz declares that he has no conflict of interest. Andreas Patzak declares that he has no conflict of interest. Linda Rausch declares that she has no conflict of interest. Nicolaus C. Netzer declares that he has no conflict of interest.

Ethical approval All procedures performed in studies involving human participants were in accordance with the ethical standards of the institutional and/or national research committee and with the 1964 Helsinki declaration and its later amendments or comparable ethical standards. The study protocol has been reviewed and approved by the ethical review board of the Bavarian Physician Chamber on the 22.05.2014. This article does not contain any studies with animals performed by any of the authors.

Informed consent Informed consent was obtained from all individual participants included in the study.

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