

Acute Mountain Sickness in the Pyrenees: an observational cross-sectional study

Clara Weisweiler

Universitat de Girona

Marc Ayala

Hospital Nostra Senyora de Meritxell

Iñigo Soteras

Hospital de Cerdanya

Enric Subirats

Universitat de Girona

Joan Carles Trullàs (✉ jctv5153@comg.cat)

Hospital d'Olot <https://orcid.org/0000-0002-7380-3475>

Research

Keywords: Altitude, Altitude Sickness, Mountaineering

Posted Date: June 26th, 2020

DOI: <https://doi.org/10.21203/rs.3.rs-37886/v1>

License:  This work is licensed under a Creative Commons Attribution 4.0 International License.

[Read Full License](#)

Abstract

Background

The prevalence of acute mountain sickness (AMS) ranges between 15% and 80% depending on the absolute altitude reached, speed of ascent, and individual susceptibility. However, there is a lack of information regarding AMS at moderate to high altitudes (2,500-3,500 m) and, even less, in the Pyrenees. Our aim is to determine the prevalence and risk factor of AMS in the Pyrenees.

Methods

A cross-sectional study including mountaineers who climbed a mountain with a height greater than 2,500 m in the Pyrenees region during July and August 2019. Sociodemographic data, medical history and activity information were collected using a questionnaire. The diagnosis of AMS was based on the 2018 modified Lake Louise Score. A logistic regression analysis was performed to examine the association of different variables (risk factors) and AMS.

Results

From 437 participants, 117 met diagnostic criteria of AMS, establishing a prevalence of 26.7% (95% confidence interval: 22.6%-30.9%). Individuals affected by AMS had mild (88%) or moderate (12%) affection. The most common symptoms (in addition to headache which is mandatory for AMS diagnosis) were fatigue or weakness, gastrointestinal symptoms and dizziness. In an adjusted multivariate analysis, heavy perceived exertion, bad physical condition, nonsteroidal anti-inflammatory drugs use and previous history of altitude illness were independent risk factors for developing AMS.

Conclusions

One fourth of climbers in the Pyrenees experienced mild or moderate AMS. Previous history of AMS, nonsteroidal anti-inflammatory drugs use and other modifiable risk factors such as physical exertion and physical condition were strong and independent predictors of AMS. These findings suggest that educational/informational programs for individuals planning to climb to moderate-high altitudes in the Pyrenees may contribute to prevent AMS.

Background

High-altitude illness (HAI) is an encompassing term for the range of pathology that the unacclimatised individual can develop at increased altitude. This includes acute mountain sickness (AMS), high-altitude cerebral edema and high-altitude pulmonary edema [1].

AMS is by far the most common HAI, arises after at least 4–6 hours spent at an altitude above 2,000–2,500 m and its reported prevalence ranges from 8–25% at 2,500–3,000 m and from 40–60% at 4,500 m [1]. The prevalence of AMS varies widely according to different ascent profiles. Studies conducted in Nepal, Colorado, Kilimanjaro, and the Alps show a prevalence of AMS ranging from 9–58%, with a higher prevalence at higher altitudes [2].

On the other hand, AMS has been associated with a number of potential risk factors including: age, gender, genetics, residence at an altitude below 800 m, pre-existing medical conditions, physical condition and intensity of exercise, rate of ascent and reached altitude, lack of acclimatization and previous history of HAI [3, 4].

There is a risk of AMS in the Pyrenees, but to our knowledge, no studies have determined the prevalence and risk factors of AMS in this area. For this reason, our aim is to determinate the prevalence and risk factors for AMS in mountaineers exposed to moderate-high altitude in the Pyrenees region.

Methods

Study design

This is an observational cross-sectional study to determine prevalence.

Study setting, study population and data collection

The study population includes mountaineers who climbed a mountain with a height greater than 2,500 m in the Pyrenees (the maximum altitude in this area is 3,404 m at the Aneto summit) during July and August 2019.

Data were collected using a questionnaire at the Renclusa refuge (Bellasque Valley, Huesca, Spain) at 2,138 m, where most of the mountaineers who climb mountains over 3,000 meters high pass through. During the study period all mountaineers passing through the refuge were asked to voluntarily participate in our study by answering the questionnaire. The questionnaire could be answered in two different ways, in person and online. The online form consisted of a link or QR code that was given to the mountaineers once the mountain activity was finished allowing to answer it in a quiet place and once the symptoms (if present) were resolved. The estimated time to answer the questionnaire was about 3 minutes and no personal data was collected to ensure data protection. The privacy and the safety of this online form has been always ensured. Once the questionnaire was completed all participants received a tryptic with more information about high-altitude diseases.

Study variables

The questionnaire included 28 questions with the following variables: sociodemographic (age, gender, height, weight, place of residence and sea level situation), smoking habits, medical conditions (including diabetes, cholesterol, hypertension, chronic obstructive pulmonary disease, obstructive sleep apnea

syndrome (OSAS), cardiovascular diseases, asthma, migraine and anemia), drug consumption (including nonsteroidal anti-inflammatory drugs [NSAIDs] in the previous 48 hours), physical condition (bad / acceptable / good / very good), training habits (hours per week), recent (previous 3 days) activity at more than 2,500 m high, spending the previous night at the refuge (or at height) and previous history of HAI. We also collected the following variables related to the mountain activity: maximum altitude reached, time of ascension and altitude difference, physical exertion (normal / moderate / severe), sport or discipline (mountaineering, alpinism, trail-running) and liquid consumption during the activity. Finally, the questionnaire included five questions about four different AMS-related symptoms (experienced during the activity) to assess the diagnosis of AMS based on the Lake Louise Score (LLS) (see below).

Diagnosis of acute mountain sickness

For the diagnosis of AMS we used the 2018 modified LLS because is a well-accepted standard for AMS diagnosis. The LLS can be self-administrated and includes the following items: headache, nausea/vomiting, fatigue, and dizziness/light-headedness. For a positive AMS definition, it is mandatory to have a headache score of at least one point, and a total score of at least three points (Table 1) [5].

Table 1
Lake Louis Score 2018*

Headache
0. None at all
1. A mild headache
2. Moderate headache
3. Severe headache, incapacitating
Gastrointestinal symptoms
0. Good appetite
1. Poor appetite or nausea
2. Moderate nausea or vomiting
3. Severe nausea and vomiting, incapacitating
Fatigue and/or weakness
0. Not tired or weak
1. Mild fatigue/weakness
2. Moderate fatigue/weakness
3. Severe fatigue/weakness, incapacitating
Dizziness/light-headedness
0. No dizziness/ light-headedness
1. Mild fatigue/weakness
2. Moderate dizziness/light-headedness
3. Severe dizziness/light-headedness, incapacitating
Acute mountain sickness Clinical Functional Score. Overall, if you had acute mountain sickness symptoms, how did they affect your activities?
0. Not at all
1. Symptoms present, but did no force any change in activity or itinerary
2. My symptoms forced me to stop the ascent or to go down on my own power
3. Had to be evacuated to a lower altitude

*The Lake Louise Score for an individual is the sum of the score for the four symptoms (headache, nausea/vomiting, fatigue, and dizziness/light-headedness). For a positive AMS definition, it is mandatory to have a headache score of at least one point, and a total score of at least three points. The severity of AMS can be classified as follows: mild (3–5 points), moderate (6–9 points) and severe (10–12 points).

Altitude ranges definition

We defined altitude ranges according to the terminology recommended by an international expert panel: low altitude (500–2,000 m), moderate altitude (2,000–3,000 m), high altitude (3,000–5,500 m) and extreme altitude (> 5,500 m) [6].

Statistical analysis

Sample size: based on previous studies we estimated that a sample of 400–500 subjects would be enough to estimate with a 95% confidence and a precision of ± 4 per cent units, a population percentage considered to be around 25–30% [7, 8].

Descriptive analysis: Qualitative or categorical variables are expressed as number of patients and percentages and quantitative variables as median and interquartile range [IQR]. The Kolmogorov–Smirnov test was used to determine whether quantitative variables were normally distributed.

Bivariate analyses: The Chi-square test (χ^2) has been used to compare qualitative variables (when the expected number of cases in any of the cells was lower than 5, the Fisher exact test was used) and the Student's T test to compare normally distributed quantitative variables (the Mann-Whitney U test for the non-normally distributed ones). Multivariate analyses: A binary logistic regression analysis has been performed to assess the association between different predictor variables (independent variables) with AMS (dependent and dichotomous variable). This analysis has been adjusted for variables that are statistically significant ($p < 0.05$) in the bivariate analysis and those risk/protective factors previously described in the literature. We applied manual and automatic procedures (including backward and forward stepwise).

Statistical significance was set at 0.05. Analyses were performed with the software Statistical Package for Social Sciences version 20.0 (SPSS, Inc., Chicago, Illinois, USA).

Ethical Aspects

Before answering the questionnaire, all participants read and accepted a study information form and informed consent. To maintain the confidentiality and data security, no personal data were collected (including names, postcodes, addresses or birth dates). Data security was ensured with a locked network only accessible for the principal investigator of the study. According to the national and international laws regarding autonomy, the study is governed by the Organic Law (15-1999 December 13th) for personal data protection. A local review board from Universitat de Girona approved the study.

Results

Four hundred and forty-seven individuals participated in the study (only 9 mountaineers refused to participate because they were not feeling well (two cases) or they did not have time (seven cases)). After reviewing the questionnaires, 10 participants were excluded for the following reason: 3 answered the

questionnaire inappropriately, 4 had climbed to heights greater than 3,404 m (it is not possible to climb these heights in the Pyrenees as the maximum altitude is 3,404 m at the Aneto summit) and 3 had climbed to heights less than 2,500 m. The final sample size included 437 individuals (124 filled out the questionnaire in person and 313 on-line).

Baseline characteristics are shown in Table 2 and variables related with the mountain activity in Table 3. Most participants were young males with normal body-mass index, with a good health status (low proportion of smokers and comorbidities) and good physical condition. Almost 90% lived below 800 meters from sea level and 24% had previous history of HAI. More than one fourth of participants had done a recent activity (previous 3 days) or slept above 2,500 meters. All the participants reached moderate-high altitudes in a median time of 5 hours, describing a moderate physical exertion in most cases.

Table 2
Baseline characteristics and univariate analysis risk for acute mountain sickness

Variable	Total	No AMS	AMS	P value
Number	437	320	117	-
Male gender	314 (71.9%)	236 (73.8%)	76 (65.0%)	0.07
Age (years)	34 [18]	34 [17]	33 [19]	0.70
BMI (kg/m^2)	22.8 [3.2]	22.8 [3.1]	22.9 [3.4]	0.92
BMI category	13 (3%)	-	-	-
Low weight	348 (79.6%)			
Normal	69 (15.8%)			
Overweight	7 (1.6%)			
Obesity				
Smoking habits	307 (70.3)	-	-	-
No Smoker	85 (19.4)	-	-	-
Former smoker	45 (10.3)	32 (10.0%)	13 (11.1%)	0.94
Smoker				
Comorbidities	1 (0.2%)	1 (0.3%)	0	0.55
Diabetes	16 (3.7%)	12 (3.8%)	4 (3.4%)	0.87
Cholesterol	11 (2.5%)	7 (2.2%)	4 (3.4%)	0.47
Hypertension	3 (0.7%)	1 (0.3%)	2 (1.7%)	0.12
Cardiovascular	12 (2.7%)	9 (2.8%)	3 (2.6%)	0.89
Asthma	6 (1.4%)	2 (0.6%)	4 (3.4%)	0.03
OSAS	0	0	0	0.91
COPD	7 (1.6%)	5 (1.6%)	2 (1.7%)	0.49
Migraine	8 (1.8%)	5 (1.6%)	3 (2.6%)	0.40
Anemia				

Quantitative variables are expressed as median [interquartile range]. Categorical variables are expressed as number and percentage. Abbreviations: AMS: acute mountain sickness; BMI: body mass index; NSAIDs: nonsteroidal anti-inflammatory drugs; OSAS: obstructive sleep apnea syndrome; COPD: chronic obstructive pulmonary disease; HAL: high altitude illness

Variable	Total	No AMS	AMS	P value
Medication	47 (10.8%)	32 (10.0%)	15 (12.8%)	0.40
Drugs	11 (2.5%)	7 (2.2%)	4 (3.4%)	0.48
Herbal medicine	60 (13.7%)	36 (11.2%)	24 (20.5%)	0.01
NSAIDs < 48 hours				
Residence below 800 m	393 (89.9%)	283 (88.4%)	110 (94.0%)	0.09
Previous HAI	105 (24%)	65 (20.3%)	40 (34.2%)	0.003
Physical activity (h/week)	7 [5]	7.5 [5.0]	6.0 [4.0]	< 0.001
Physical condition	6 (1.4)	4 (1.2%)	2 (1.7%)	0.02
Bad	82 (18.8)	49 (15.3%)	33 (28.2%)	
Acceptable	268 (61.3)	203 (63.4%)	65 (55.6%)	
Good	81 (18.5)	64 (20.0%)	17 (14.5%)	
Very good				

Quantitative variables are expressed as median [interquartile range]. Categorical variables are expressed as number and percentage. **Abbreviations:** AMS: acute mountain sickness; BMI: body mass index; NSAIDs: nonsteroidal anti-inflammatory drugs; OSAS: obstructive sleep apnea syndrome; COPD: chronic obstructive pulmonary disease; HAI: high altitude illness

Table 3
Mountain activity and univariate analysis risk for acute mountain sickness

Variable	Total	No AMS	AMS	P value
Number	437	320	117	-
Recent (previous 3 days) activity > 2,500 m	148 (33.9%)	116 (36.2)	32 (27.4)	0.08
Slept > 2,500 m	116 (26.5%)	83 (25.9)	33 (28.2)	0.64
Maximum reached altitude (meters)	3,404 [263]	3,404 [294]	3,404 [260]	0.10
	Mean 3,275	Mean 3,264	Mean 3,294	
Was altitude > 3000 meters reached? (yes answer)	387 (88.6%)	281 (87.8%)	106 (90.6%)	0.42
Ascension time (hours)	5 [9]	5.0 [1.0]	5.0 [2.0]	0.03
	Mean 4.7	Mean 4.64	Mean 4.87	
Altitude difference (meters) ^a	1,500 [300]	1,500 [313]	1,500 [290]	0.08
	Mean 1,516	Mean 1,503	Mean 1,553	
Ascension rate (meters/hour)	300 [110]	320 [95]	316 [108]	0.54
	Mean 335	Mean 337	Mean 329	
Physical exertion	99 (22.7%)	84 (26.2%)	15 (12.8%)	< 0.001
Normal	237 (54.2%)	183 (57.2%)	54 (46.2%)	
Moderate			48 (41.0%)	
Intense	101 (23.1%)	53 (16.6%)		
Discipline ^b	284 (65%)	206 (64.4%)	78 (66.7%)	0.74
Mountaineering	48 (11%)	36 (11.2%)	12 (10.3%)	
Running	102 (23.3%)	75 (23.4%)	27 (23.1%)	
Alpinism				
Fluid intake (liters)	2 [1]	2 [1]	2 [1]	0.57

Quantitative variables are expressed as median [interquartile range] and mean when specified.

Categorical variables are expressed as number and percentage. ^aAltitude difference means difference in altitude from the starting point to the highest altitude. ^bRunners had a significantly lower ascension time and higher ascension rate in comparison with other disciplines, but no significant differences regarding maximum altitude were found.

The prevalence of AMS and the severity of symptoms are detailed in Table 4. According to the 2018 LLS, 117 participants met diagnostic criteria of AMS, establishing a prevalence of 26.7% (95% confidence interval: 22.7–30.9%). The prevalence of AMS in online and in person responders was 26.6% and 29.9%, respectively (this difference did not reach statistical significance). All cases were mild (88%) or moderate (12%) and the most frequent (and severe) symptom (apart from headache, which is mandatory for diagnosis) was fatigue (98.3%) followed by gastrointestinal symptoms (59%) and dizziness (47%).

Table 4
Prevalence and severity of symptoms of acute mountain sickness^a

AMS	Number		Percentage		
	Total	Mild	2	3	Total
	Not at all	Mild	Moderate	Severe	
Total	117		26.7%		
Mild	103		88%		
Moderate	14		12%		
Severe	0		0%		
Symptoms	0	1	2	3	Total
Headache	0	88 (75.2%)	25 (21.4%)	4 (3.4%)	117 (100%)
Gastrointestinal	48 (41%)	65 (94.2%)	3 (4.3%)	1 (1.4%)	69 (59%)
Fatigue	2 (1.7%)	57 (49.6%)	48 (41.7%)	10 (8.7%)	115 (98.3%)
Dizziness	62 (53%)	45 (81.8%)	9 (16.4%)	1 (1.8%)	55 (47%)

^aAccording to the 2018 Lake Louis score. Results are expressed as number and percentage.
Abbreviations: AMS: acute mountain sickness.

In the univariate analysis the following variables were associated with an increased risk of AMS (Tables 2 and 3): history of OSAS, the use of NSAIDs in the previous 48 hours, previous history of HAI, less physical activity per week, worse physical condition, slower ascension time and intense physical exertion during the activity. We also observed a greater risk of AMS in women, residents below 800 meters from sea level and not performing a recent activity above 2,500 meters, but these differences didn't reach statistical significance.

We found no differences in AMS with respect to the maximum altitude reached, but those who climbed above 3000 meters suffered more intense symptoms (mild AMS in 90.6%) compared to those who climbed below 3000 meters (mild AMS in 63.6%), being these differences statistically significant ($p = 0.009$)

The individuals with AMS who took NSAIDs in the previous 48 hours had higher scores in the LLS compared to those who did not take them (median [IQR]: 4.5 [2.5] and 4.0 [1.0], respectively [p value

0.03]). In addition, the intensity of AMS was different according to NSAIDs consumption; AMS was moderate in 25% of those who took NSAIDs and in 8.6% of those who did not take them (*p* value 0.02). There were no differences in the proportion of patients who took NSAIDs based on whether they had previous history of AMS (14% vs 14%, *p* value 0.85). Thus, NSAIDs use was not associated with the history of AMS.

Table 5 summarizes the univariate and multivariate analysis risk factors for AMS. In the multivariate adjusted analysis the following variables remained as risk factors for developing AMS: physical exertion during ascent (OR 2.24; 95% CI 1.22–4.12), use of NSAIDs in the previous 48 hours (OR 1.95; 95% CI 1.09–3.50) and previous history of HAI (OR 1.98; 95% CI 1.22–3.21). On the other hand, having a good physical condition was a protective factor against suffering AMS (OR 0.48; 95% CI 0.29–0.79).

Table 5
Risk factors for acute mountain sickness. Univariate and multivariate analysis

	Unadjusted	<i>p</i>	Adjusted	<i>p</i>
	OR (CI 95%)	value	OR (CI 95%)	value
Age	0.99 (0.98–1.01)	0.62	0.98 (0.96–1.0)	0.091
Male gender	0.65 (0.42–1.04)	0.072	0.62 (0.36–1.06)	0.081
Residence < 800 meters	2.06 (0.89–4.75)	0.086	1.75 (0.72–4.26)	0.220
BMI	1.01 (0.95–1.09)	0.74	1.04 (0.93–1.16)	0.477
OSAS	5.63 (1.02–31.4)	0.026	3.54 (0.52–24.2)	0.197
NSAIDs < 48 hours	2.04 (1.16–3.56)	0.013	1.95 (1.09–3.50)	0.026
Smoke	1.07 (0.67–1.69)	0.78	1.03 (0.62–1.73)	0.889
History of HAI	2.04 (1.23–3.26)	0.003	1.98 (1.22–3.21)	0.006
Physical activity (hours/week)	0.92 (0.82–0.98)	0.004	0.94 (0.89–1.00)	0.062
Good physical condition	0.47 (0.28–0.76)	0.002	0.48 (0.29–0.79)	0.004
Maximum altitude (meters)	1.0 (1.0–1.0)	0.25	1.0 (0.99–1.0)	0.327
Ascension time (hours)	1.19 (0.99–1.44)	0.063	1.06 (0.82–1.36)	0.672
Altitude difference (meters) ^a	1.0 (1.0–1.0)	0.15	1.0 (0.99–1.0)	0.603
Physical exertion	2.42 (1.33–4.40)	0.003	2.24 (1.22–4.12)	0.009
Recent (3 days) activity > 2,500 m	0.66 (0.42–1.06)	0.082	0.69 (0.42–1.14)	0.150

Results are expressed as OR (odds ratio) and 95% CI (confidence interval). ^aAltitude difference means difference in altitude from the starting point to the highest altitude. Abbreviations: BMI: body mass index; NSAIDs: nonsteroidal anti-inflammatory drugs; OSAS: obstructive sleep apnea syndrome; HAI: high altitude illness

Discussion

Using the modified LLS criteria, we found a prevalence of AMS of 26.7% in mountaineers ascending to moderate-high altitudes in the Pyrenees region. To our knowledge, this is the first study that analyses the prevalence and risk factors of AMS in this region.

There is less information regarding the prevalence of AMS at moderate to high altitudes (around 3,000 m) in comparison with high to extreme altitudes. Our impression is that the prevalence found in our study is higher than the expected but it is difficult to compare our results with previous studies performed at similar heights because of differences in geographic locations and studied populations. AMS occurred in 25% of visitors to moderate altitudes (1,920-2,956 meters) in the Rocky Mountains of Colorado [7]. The prevalence of AMS in other studies performed at Mount Fuji [9] and Western and Eastern Alps [10] was 29.5% (3,776 meters), 34.9% (3,817 meters) and 38.0% (3,454 meters), respectively. It is also important to highlight that these studies were conducted before the LLS was modified in 2018 and, to our knowledge, no studies regarding AMS prevalence have been published using this modified score. Other factors could explain the differences in AMS prevalence at similar altitudes (different hut locations, different weather conditions, different levels of mountaineer experience, etc.) but the explanation to them is beyond the scope of this study.

We have also found different risk factors for AMS, many of them already described in the literature that deserve discussion.

Men and women appear to be equally at risk for AMS although some observational studies suggest a slightly higher risk for women [7]. We also found a higher risk for AMS in women but the statistical significance of this difference was borderline (p value = 0.07).

Regarding pre-existing medical conditions we only found that OSAS was associated with a greater risk for AMS in the univariate analysis. Several studies have shown that obesity and nocturnal hypoxemia are risk factors for the development of AMS. Patients with OSAS and significant arterial desaturation at sea level would be expected to have more profound arterial desaturation during apneic periods at high altitude, but there are no data on this issue [11].

The most relevant risk factor (with a higher odds ratio) for developing AMS in our study was the intensity of exertion perceived by mountaineers. Those individuals who reported heavy exertion had more than 2 fold risk of developing AMS. This finding has also been found in other studies and could be explained due to the stress caused in the autonomic nervous system by additional hypoxia that generates an intense exercise [3, 10]. The role of exercise in this study must be discussed. We found a high proportion of subjects being fatigued (half of them moderately to severe) and it is difficult to assess whether these symptoms are secondary to AMS or indicate the effect of exercise. Furthermore, Moore et al have recently drawn attention to the inclusion of fatigue in the LLS. These authors suggest that fatigue may contribute to increase false positive AMS diagnoses and propose to remove this symptom from the score [12]. For

this reason, we believe the reported AMS scores would have been lower without the inclusion of fatigue and the prevalence of AMS would have been lower.

Moreover, self-assessed bad physical condition was found to be another significant risk factor for AMS. This can be explained, in part, by the fact that physical condition is related to the level of exertion during the ascent, suggesting that low fitness climbers do not tolerate the unusual exertion of mountaineering, or they appear to workout excessively. The results regarding this issue in the literature are inconclusive, with some studies finding that physical condition is a risk factor for AMS [7, 10] while others not [4, 13]. This variability between studies may be explained because self-reported physical condition is difficult to objectively evaluate with considerable variability between individuals.

We also found that history of HAI on previous exposures was also a risk factor for suffering AMS (almost more than two fold risk). Based on our findings and previous studies, a strong relationship seems to exist between a self-report of previous HAI and the risk of subsequent development of AMS. There are different theories regarding individual susceptibility to HAI, but it is likely to be derived from both genetic and environmental variables. The genetic influence of AMS remains an active area of investigation with no identified specific genetic predisposition [2, 4, 7, 13–15].

Finally, we also found that NSAIDs consumption in the previous 48 hours was associated with an increased risk for AMS. However, caution should be applied when interpreting this apparently confusing result. We do not believe that taking an analgesic places one at a greater risk for subsequent AMS. It is likely that many of our participants who used NSAIDs did so after they started experiencing symptoms associated with AMS in an effort to alleviate those symptoms. The findings that LLS was higher and AMS was more severe in those taking NSAIDs supports this hypothesis. The same finding and interpretation has also been reported in previous studies [13]. On the other hand, the use of NSAIDs was not associated with a previous history of AMS, so it seems unlikely that their use was preventive to avoid symptoms.

It is commonly accepted that residing at an altitude above 800 meters from sea level offers protection against AMS [7, 16]. In our study, we found a higher proportion of participants living below 800 meters from sea level in the ones who suffered AMS compared with those who not, but these differences did not reach statistical significance ($p = 0.09$). This may be explained, in part, because only 10% of participants were living above 800 meters from the sea level.

This study has some limitations that must be taken into account. First, data were collected in person and on-line and this could contribute to a participant selection bias. Despite this, we did not find differences in the prevalence of AMS between the two types of data collection. Second, the individuals who participated in the study did so voluntarily (non-probabilistic method), allowing perhaps to select individuals with “different” characteristics to those who decide not to participate (more motivation, less severity of symptoms, etc.). Third, the results presented here have performed in a specific area of the Pyrenees and probably cannot be generalized.

Conclusions

The prevalence of AMS (using the modified 2018 LLS) in mountaineers at moderate-high altitudes (lower than 3,404 meters) at the Pyrenees region is not negligible (26.7%) but probably slightly overestimated. Most cases of AMS were mild (88%) or moderate (12%) and we did not find any severe case. The independent risk factors for developing AMS were previous history of HAI, NSAIDs use, and other modifiable risk factors such as physical exertion and bad physical condition. In light of these results, we believe that more educational/informational programs for individuals planning to climb to moderate-high altitudes in the Pyrenees would contribute to prevent AMS. We also encourage researchers to conduct new studies to confirm these results.

Abbreviations

AMS
Acute mountain sickness
HAI
High-altitude illness
IQR
Interquartile range
LLD
Lake Louise Score
NSAIDs
nonsteroidal anti-inflammatory drugs
OSAS
Obstructive sleep apnea syndrome

Declarations

Ethics approval and consent to participate:

A local review board from Universitat de Girona approved the study and all individuals read and accepted a study information form and informed consent before participation.

Availability of data and materials:

All data generated or analysed during this study are included in this published article. In addition, the datasets used and/or analysed during the current study are available from the corresponding author on reasonable request.

Competing interests:

The authors declare that they have no competing interests.

Funding:

None

Authors' contributions:

study concept and design (all authors); acquisition of the data (CW); analysis of the data (JCT and CW); drafting of the manuscript (JCT and CW); critical revision of the manuscript; and approval of the final manuscript (all authors)

Acknowledgements:

We greatly appreciate all those who have contributed to this study being carried out, especially Montaña Segura and Aneto Seguro, la Renclusa refuge, Amigos de los Pirineos and 3.000 dels Pirineus.

Authors' information:

The results of this study were presented on February 2020 by the first author (CW) as her Final Degree Project at the School of Medicine, University of Girona, Girona, Spain. The final qualification was excellent (9.7/10).

References

1. - Smedley T, Grocott MP. Acute high-altitude illness: a clinically orientated review. *Br J Pain*. 2013;7:85–94.
2. - Taylor AT. High-altitude illnesses: physiology, risk factors, prevention, and treatment. *Rambam Maimonides Med J*. 2011;2:e0022.
3. - Burtscher M, Mairer K, Wille M, Broessner G. Risk factors for high-altitude headache in mountaineers. *Cephalgia*. 2011;31:706–11.
4. - Schneider M, Bernasch D, Weymann J, Holle R, Bartsch P. Acute mountain sickness: influence of susceptibility, preexposure, and ascent rate. *Med Sci Sports Exerc*. 2002;34:1886–91.
5. - Roach RC, Hackett PH, Oelz O, Bärtsch P, Luks AM, MacInnis MJ, et al. The 2018 Lake Louise Acute Mountain Sickness Score. *High Alt Med Biol*. 2018;19:4–6.
6. - Bärtsch P, Saltin B. General introduction to altitude adaptation and mountain sickness. *Scand J Med Sci Sports*. 2008;18(Suppl 1):1–10.

7. - Honigman B, Theis MK, Koziol-McLain J, Roach R, Yip R, Houston C, et al. Acute mountain sickness in a general tourist population at moderate altitudes. *Ann Intern Med.* 1993;118:587–92.
8. - Sánchez-Mascuñano A, Masuet-Aumatell C, Morchón-Ramos S, Ramon JM. Relationship of altitude mountain sickness and smoking: a Catalan traveller's cohort study. *BMJ Open.* 2017;7:e017058.
9. - Horiuchi M, Endo J, Akatsuka S, Uno T, Jones TE. Prevalence of acute mountain sickness on Mount Fuji: A pilot study. *J Travel Med.* 2016;23:taw024.
10. - Mairer K, Wille M, Burtscher M. The prevalence of and risk factors for acute mountain sickness in the Eastern and Western Alps. *High Alt Med Biol.* 2010;11:343–8.
11. - Luks AM, Swenson ER. Travel to high altitude with pre-existing lung disease. *Eur Respir J.* 2007;29:770–92.
12. - Moore J, MacInnis MJ, Dallimore J, Wilkes M. The Lake Louise Score: A Critical Assessment of Its Specificity. *High Alt Med Biol.* 2020. 10.1089/ham.2019.0117. doi:10.1089/ham.2019.0117.
13. - Wagner DR, Fargo JD, Parker D, Tatsugawa K, Young TA. Variables contributing to acute mountain sickness on the summit of Mt Whitney. *Wilderness Environ Med.* 2006;17:221–8.
14. - Rupert J. Will blood tell? Three recent articles demonstrate genetic selection in Tibetans. *High Alt Med Biol.* 2010;11:307–8.
15. - Van Roo JD, Lazio MP, Pesce C, Malik S, Courtney DM. Visual analog scale (VAS) for assessment of acute mountain sickness (AMS) on Aconcagua. *Wilderness Environ Med.* 2011;22:7–14.
16. - Grocott M, Montgomery H, Vercueil A. High-altitude physiology and pathophysiology: implications and relevance for intensive care medicine. *Crit Care.* 2007;11:203.

Supplementary Files

This is a list of supplementary files associated with this preprint. Click to download.

- [Coverletter.docx](#)