Incidence and severity of acute mountain sickness and associated symptoms in children trekking on Xue Mountain, Taiwan

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Abstract

Background

Acute mountain sickness (AMS) occurs in non-acclimatized people after an acute ascent to an altitude of 2,500 m or higher. The aim of this study was to examine the incidence and severity of AMS and associated symptoms in children.

Methods

The prospective observational study included 197 healthy, non-acclimatized 11 and 12-year-old children trekking the round-trip from the trailhead to the summit of Xue Mountain, Taiwan (2,179 m to 3,886 m) over 3 days. AMS was evaluated at Qika Hut (2,460 m) on Day 1, at Sanliujiu Hut on Day 2 (3,100 m), and at the same altitude (3,100 m) after reaching the summit on Day 3. We used the Lake Louise Score (LLS) to diagnose AMS and record daily AMS-associated symptoms. We gave acetazolamide to children with mild to moderate AMS. Dexamethasone was reserved for individuals suffering from severe AMS. Acetaminophen was administrated to children with headache, and metoclopramide for those with nausea or vomiting.

Results

There were 197 subjects eligible for analysis. The overall incidence of AMS was 40.6%, which was higher in males and in subjects with a higher body mass index (BMI) (p < 0.05).
The prevalence of AMS on Day 1 was 5.6%, which was significantly lower than that on Day 2 (29.4%) and Day 3 (23.4%). The mean LLS of all subjects was 1.77 ± 2.08. The overall incidence of severe AMS (LLS ≥ 5) was 12.5%. The mean LLS of the AMS group (3.02 ± 2.46) was significantly higher than that of the non-AMS group (0.92 ± 1.16, p < 0.001). Among the AMS group, the mean LLS was 1.00 ± 1.55 on Day 1, 4.09 ± 1.97 on Day 2, and 3.98 ± 2.42 on Day 3. The most common symptom was sleep disturbance followed by dizziness, and headache. The prevalence of headache was 46.2% on Day 2 at 3,100 m, and 31.3% on Day 3 at the same altitude after climbing the summit (3,886 m). Males experienced significantly more headache and fatigue than females (p < 0.05). The LLS and prevalence of all AMS symptoms were significantly higher in the AMS than the non-AMS group (p < 0.05).

**Conclusions**

The AMS incidence among children trekking to Xue Mountain was 40.6%. AMS is common and mostly manifests as mild symptoms. Gender (male) and a higher BMI could be considered two independent risk factors of higher AMS incidence. Sleep disturbance is the most common symptom, and the lower prevalence of headache on Day 3 may be due to the effects of medication and/or acclimatization.

**Introduction**

Acute mountain sickness (AMS) often occurs in non-acclimatized people after an acute ascent to an altitude of 2,500 m or higher. With the development of modern transportation, more lowland children and adolescents are travelling to high altitudes on school expeditions, or for recreational purposes such as skiing, climbing, and mountaineering. However, previous large group studies on this population are rare. The reported incidence in children varies, and the susceptibility to AMS in this age group compared to adults is controversial [1,2,3]. Several factors contribute to the variations in incidence, including age, AMS definitions, modes and rates of ascent (trekking or by vehicles), and sleeping altitudes [3,4].

The cardinal symptom of AMS is headache, accompanied by other symptoms such as dizziness, lightheadedness, fatigue, weakness, nausea, vomiting, loss of appetite, and sleep disturbance [1,5]. Most AMS symptoms are mild; however, at altitudes above 3,000 m, there is a chance of developing high-altitude cerebral edema (HACE) and high-altitude pulmonary edema (HAPE). In previous reports on adults, HACE usually develops after at least 2 days at altitudes above 4,000 m, and the incidence is estimated to be 0.5–1.0% at 4,000–5,000 m. HAPE develops after 2 or more days at altitudes above 3,000 m, with an incidence of 0.2–2% at 4,500–5,500 m in subjects without a previous history of HAPE [5]. Reports of HAPE and HACE in children are rare, and are usually individual cases. Children who have an upper respiratory infection tend to have a greater chance of developing AMS, and the risk of HAPE, and occasionally HACE, also increases. Children with HAPE have coughs that gradually become worse, and rales on chest auscultation may also be present [1,6].

The diagnostic criteria of AMS were well established in 1991, at the Hypoxia and Mountain Medicine Symposium at Lake Louise, Canada, where a group of experts reached a consensus regarding the diagnosis [7]. The course of AMS is mostly benign and self-limited; it generally occurs within 6–12 hours of ascent [4,5,8,9] and resolves within 1–3 days after descent [5,10]. The major independent risk factors include lack of acclimatization [11,12], rapid ascent, high
sleeping altitude [6, 9, 13–15], and a previous history of acute mountain sickness [4,8,15]. However, in one recent study, a history of AMS in children was determined to have no predictive value. None of the children with AMS on re-exposure had AMS the first time, compared with 80% of reproducible AMS among adults [16]. Other factors that may affect the development of AMS are age, gender, residence altitude, concurrent medical illness [1,6,8], and general physical condition [8,9]. Young age is reported to be associated with the development of AMS [8,9,11,15,17–18], while some studies have found no difference between children and adults [2,19,20].

Over the past 30 years, many studies discussed AMS among the adult population. The AMS incidence has ranged from approximately 25% at moderate altitude [8,19,21] to 50–85% at high altitude [17,20,22,23]. However, few studies explored AMS exclusively among children and adolescents, and the results remain inconclusive. Imray et al. reported an incidence of 91.7% in adolescents aged 15–18 years trekking to 4,200–5,500 m in Machu Picchu, Peru [24]. Chan et al. reported an incidence of 59% in older children trekking to Jade Mountain, Taiwan (3,952 m) in 3 days [25]. A group of children and adolescents were studied after rapid ascent from 570 m to 3,450 m in a 3-hour train ride, with a reported incidence of 30–37.5% [3,4]. Theis et al. reported an incidence of 28% in children aged 9–14 years-old at a moderate altitude in Colorado (2,835 m). However, a similar proportion (21%) of children developed these symptoms at sea level. The author cites several important factors other than altitude that may affect the present of symptoms in children, such as travel fatigue, anxiety, and disruption of daily routine [26]. Similar phenomena were also reported in later studies [1].

The aim of this study was to examine the incidence and severity of AMS and associated symptoms in children trekking to Xue Mountain (3,886 m) in Taiwan. AMS was evaluated by trained physicians using the Lake Louise Score (LLS) questionnaire. The prevalence of LLS pertaining to each symptom was also analyzed in order to increase our understanding of AMS in this population.

**Methods**

**Study group**

This study was performed during a primary school yearly trekking event, and nearly all of the 6th grade students participated in the activity; few dropped out. There were 201 subjects initially; 1 was excluded due to a lack of informed consent, and 3 were excluded due to incomplete data, leaving a total of 197 subjects (age = 11.78 ± 0.43 years; 111 males and 86 females). All subjects were living at altitudes below 500 m. None of the subjects had high altitude exposure (> 2,500 m above sea level) in the previous 3 months. Only two subjects (who took herb medication and *Rhodiola*) had medication before ascent. The protocol was approved by the Institutional Review Board of Chang Gung Memorial Hospital, and written informed consent was obtained from all subjects and their parents.

**Study design**

This is a prospective observational study. The subjects, all from the same primary school, were divided into four different trekking groups according to their original classes. The children were separated into four cohorts of 55, 50, 47, and 45 individuals. The departure dates of the groups were separated by 1 day due to limited capacity in mountain huts. The medical team in each group consisted of one emergency physician and one nurse, both of whom were under the leadership of one medical director and one chief nurse for the duration of the trekking activity.

On Day 1, the subjects ascended from Taipei city (5 m above sea level) to the Xue Mountain trailhead (2,170 m) in approximately 6 hours by vehicles. The subjects then trekked 2 km from
the trailhead to Qika Hut (2,460 m) in about 2 hours. The evaluation of AMS symptoms and physical examinations were performed by medical staff at Qika Hut in the evening. On Day 2, the subjects trekked for 5 km and ascended to Sanliuju Hut (3,100 m) in approximately 6 hours. Once again, AMS scoring and physical examinations were performed after adequate rest more than 30 minutes. On Day 3, subjects trekked 3.8 km to the summit of Xue Mountain (3,886 m) in about 5 hours, stayed for less than 1 hour, and descended to Sanliuju Hut for another AMS evaluation. All participants then descended to the trailhead and were transported home by vehicle. All subjects received the same diet during the trip and care was taken to ensure adequate fluid intake. For subjects who met the criteria of AMS during trekking or at mountain huts, medication (acetaminophen, acetazolamide, metoclopramide, dexamethasone) was provided after evaluation by physicians.

Evaluation of AMS

The Lake Louise questionnaire was used to evaluate AMS, which included headache, dizziness/light-headedness, fatigue/weakness, gastrointestinal symptoms (nausea, vomiting, loss of appetite), and sleep disturbance. The sleep disturbance score reflected the sleep quality during the night before evaluation, but the sleep disturbance score of Day 1 was not incorporated in the analysis because all subjects slept at home (near sea level) on the night before trekking. Each symptom was scored on a scale ranging from 0 to 3, where 0 represented the absence of symptoms, 1 indicated mild symptoms, 2 indicated moderate symptoms, and 3 indicated severe and incapacitating symptoms. The sum of all five scores comprised the overall LLS, which ranged from a minimum of 0 to a maximum of 15. Subjects were considered to have AMS if they had a recent altitude gain at an elevation above 2,500 m and had headache, with at least one other symptom plus a total LLS ≥ 3. All evaluations were performed after more than 30 minutes of rest. To decrease subjective bias and the possibility of providing an incorrect answer just "for fun," serial assessments of AMS symptoms and physical examinations were performed by trained emergency physicians at specific locations, instead of being self-reported by subjects. The medical team consisted of emergency physicians and nurses who had expertise in trekking and high-altitude medicine. The medical staff not only helped the children to complete the LLS questionnaire but also observed the appearance and exercise performance of each child. Objective physical measures, such as resting oxygen saturation (SpO₂, %) and heart rate (HR, beats per minute), were also obtained.

Statistical analysis

Continuous data are expressed as means ± SD, and categorical data are presented as percentages. One-way analysis of variance followed by the Student-Newman-Keuls post hoc test was used to verify the difference in the LLS on different days. McNemar's test was used for between-day and within-group comparisons, and chi-squared test was used for between-group comparisons. The Wilcoxon signed rank test was used to detect within-group differences in the score for each symptom, and the two-tailed Mann-Whitney rank sum test was used to compare the score for each symptom between groups. P-values < 0.05 were considered statistically significant. All statistical analyses were performed using Sigma Plot software (ver 12; Systat Software, Inc, San Jose, CA, USA).

Results

The total number of subjects included in the analysis was 197 (age = 11.78 ± 0.43 years; 111 males and 86 females). In total, 80 (40.6%) of the 197 subjects experienced at least one episode of AMS (Table 1): 49 (24.9%) experienced one AMS episode, 29 (14.7%) had two AMS
episodes, and only 2 (1.0%) had AMS episodes on three occasions. One subject did not exhibit AMS on either the first or second evaluation; however, the LLS data of the third evaluation were missing. This subject attained the peak and did not receive any treatment during trekking. Thus, the subject was classified into the non-AMS group in terms of overall evaluation.

In addition, one subject had a LLS of 10, with all five symptoms presenting on Day 3, which were relieved by medication and immediate descent. None of the subjects developed HAPE or HACE during the trekking course.

Baseline characteristics of all subjects are listed in Table 1. The overall incidence of AMS was significantly greater among males than females ($p = 0.033$): 54 (48.6%) of the 111 males and 26 (30.2%) of the 86 females had AMS. Subjects with AMS had a higher body weight and body mass index (BMI) ($p = 0.008$). Logistic regression, with z-scores for age versus AMS, gender versus AMS, and BMI versus AMS showed that the odds ratio (OR) for age versus AMS was 0.89 ($z = -0.75, p = 0.455$); the OR for gender versus AMS was 1.98 ($z = 2.21; p = 0.027$), and the OR for BMI versus AMS was 1.46 ($z = 1.41; p = 0.016$). Only two subjects had medication before the ascent (herbal medications and *Rhodiola*), and they were both included in the analysis.

Fig 1 shows that the daily AMS prevalence was significantly higher (29.4%) on Day 2 compared with Day 1 (5.6%; $p < 0.001$). On Day 3, 14 children did not reach Xue Mountain peak: 10 of them had at least one AMS episode during the trip, and 4 did not have AMS. The prevalence of AMS on Day 3 was calculated as follows: 45 (total Day 3 AMS cases) divided by (197–5) (subjects without AMS who failed to reach the peak, and the one subject with missing data) = 23.4%.

For the AMS group, the mean LLS was $1.00 \pm 1.55$ on Day 1, $4.09 \pm 1.97$ on Day 2, and $3.98 \pm 2.42$ on Day 3. The LLS on Day 2 and Day 3 was significantly higher than on Day 1 ($p < 0.001$). The mean LLS of all subjects was $1.77 \pm 2.08$. In subjects with AMS, the mean LLS ($3.02 \pm 2.46$) was significantly higher than that of the non-AMS group ($0.92 \pm 1.16; p < 0.001$). There were three LLS checkpoints for each subject, and 587 (591 minus the 4 subjects without AMS who failed to attain the peak) total occasions were analyzed over the whole trek. Subjects were classified into the AMS group if they experienced AMS at any checkpoint. An LLS of $< 2$

### Table 1. Subject characteristics.$^d$

|                | Total (n = 197) | AMS group (n = 80) | Non-AMS group (n = 117) | p-value$^d$ |
|----------------|-----------------|--------------------|-------------------------|-------------|
| Gender         |                 |                    |                         |             |
| Male, n$^a$(%) | 111 (56.3%)    | 54 (48.6%)         | 57 (51.4%)              | 0.033$¶$  |
| Female, n$^a$(%) | 86 (43.7%)    | 26 (30.2%)         | 60 (69.8%)              |             |
| Mean body weight, kg | 45.1 ± 10.8  | 47.8 ± 12.1        | 43.1 ± 9.3              | 0.008*     |
| Mean body height, cm | 152.0 ± 7.1  | 152.9 ± 7.7        | 151.3 ± 6.6             | 0.113      |
| Mean BMI$^b$, kg/m$^2$ | 19.4 ± 3.7   | 20.2 ± 4.0         | 18.7 ± 3.4              | 0.008*     |
| Menstruation$^c$, n $^a$(%, of females) | 13 (15.1%)   | 6 (23.1%)          | 7 (11.7%)               | 0.398      |

Abbreviations: AMS, acute mountain sickness; BMI, body mass index. Data are presented as mean ± SD.

$^a$ n = number of children

$^b$ BMI = weight in kilograms divided by the square of the height in meters.

$^c$ In the menstruation section, the total population represents female subjects (86).

$^d$ Chi-square test was used to determine differences in categorical variable between the AMS and non-AMS groups. Mann-Whitney U-test was used to determine differences in numerical variables (body weight, body height, BMI) between the two groups.

$¶$ $p < 0.05$ indicates that AMS incidence was significantly higher in males than in females.

$*$ $p < 0.05$ indicates a significant difference between the AMS and non-AMS groups.
was observed on 72.5% of all occasions, and an LLS ≥ 5 was found on only 12.5% of all occasions.

A total of 184 participants (93.4%) had at least one of the AMS-associated symptoms. The most prevalent symptom in all subjects was sleep disturbance (74.6%), followed by dizziness, headache, fatigue, and gastrointestinal (GI) symptoms (Table 2). Twenty-six (13.2%) subjects had a total LLS ≥ 3 but fell below the threshold for AMS due to lack of headache. A total of 115 (58.4%) subjects had headache during the trip, but 35 of these headaches (30.4%) did not meet the criteria for AMS. The prevalence of headache was significantly higher (46.2%) on Day 2, and decreased to 31.3% on Day 3 at the same altitude. Sleep disturbance was also most common on Day 2, while fatigue and GI symptoms became more prevalent on Day 3. Over 40% of subjects had dizziness on Day 2 and Day 3.

Fig 2 shows the prevalence of AMS-associated symptoms in males and females. More than half (67.7%) of the male subjects experienced headache compared with 45.4% of female subjects (p < 0.05). The prevalence of fatigue was also higher among males (54.0%) than females (34.9%) (p < 0.05).

The daily prevalence and score for each symptom are shown in Figs 3 and 4. On Day 1, the main symptoms in both groups were headache and dizziness, with only a few GI symptoms reported (Fig 3A). Fewer than 7% of the non-AMS group experienced AMS symptoms, and the LLS for each symptom in both groups were all less than 0.4 on Day 1 (Fig 4A). On Day 2, headache was the most prevalent symptom (82.5%) among the AMS group, followed by sleep disturbance and dizziness, while in the non-AMS group, the main symptom was sleep disturbance (46%) followed by dizziness, headache, and fatigue (prevalence < 30% for all) (Fig 3B). Sleep disturbance had the highest LLS in both groups (AMS: 1.29 ± 0.92, non-AMS: 0.39 ± 0.38).
On Day 2 (Fig 4B). On Day 3, more than 50% of the AMS group experienced headache, dizziness, fatigue, and sleep disturbance (Fig 3C). Sleep disturbance symptoms remained the most prevalent and severe, while GI symptoms the least severe in both groups (Fig 4C). On all three days, the LLS and prevalence of all symptoms were significantly higher in the AMS versus non-AMS group (*p < 0.05) (Fig 4A–4C).

| Overall prevalence (%) of AMS-associated symptoms on different days. |
| --- |
| Headache | Dizziness | Fatigue | GI symptoms | Sleep disturbance |
| Day 1 | 14.7 | 14.2 | 11.2 | 3.6 | N/A |
| Day 2 | 46.2* | 40.6¶ | 27.9 | 12.7 | 57.9* |
| Day 3 | 31.3 | 41.7¶ | 32.3* | 22.4* | 49.5 |
| Overall | 58.4 | 61.9 | 45.7 | 28.9 | 74.6 |

Abbreviations: AMS, acute mountain sickness; GI symptoms, gastrointestinal symptoms.

* Day 1: Qika Hut, 2,460 m. Day 2: Sanliuju Hut, 3,100 m. Day 3: Sanliuju Hut, after reaching the summit of Xue Mountain (3,886 m).

¶ The sleep disturbance score reflected the sleep quality during the night before evaluation, but the sleep disturbance score of Day 1 is not taken into account because all subjects slept at home (near sea level) on the night before trekking.

* McNemar’s test was used to determine the difference in prevalence of each symptom from day to day.

* p < 0.05 indicates that the prevalence of dizziness was higher on Day 2 and Day 3 compared with Day 1. However, there was no significant difference between Day 2 and Day 3.

https://doi.org/10.1371/journal.pone.0183207.t002

Fig 2. Prevalence of AMS-associated symptoms in males and females. The prevalence of headache and fatigue was significantly higher among males than females. More than half (67.7%) of the male subjects experienced headache compared to 45.4% of females. The prevalence of fatigue was also higher among males (54%) than females (34.9%). The chi-squared test was used to detect differences in prevalence between male and female subjects. *p < 0.05 was considered statistically significant. * p < 0.05. AMS, acute mountain sickness; GI symptoms, gastrointestinal symptoms (nausea, vomiting, poor appetite).
Fig 3. Daily prevalence of AMS-associated symptoms in the AMS and non-AMS groups. A—Day 1 (Qika Hut, 2,460 m): 26% of the subjects in the AMS group experienced headache and dizziness. Fewer than 7% of the non-AMS group experienced AMS symptoms, and none experienced GI symptoms at this altitude. B—Day 2 (Sanliuju Hut, 3,100 m): headache was the most prevalent symptom (82.5%) in the AMS group, followed by sleep disturbance. In the non-AMS group, the main symptom was sleep disturbance (46%) followed by dizziness, headache, and fatigue (prevalence < 30% for all). C—Day 3 (Sanliuju Hut, 3,100 m, after reaching the summit of Xue Mountain, 3,886 m): more than 50% of the AMS group experienced headache, dizziness, fatigue, and sleep disturbances. Sleep disturbance was the most prevalent symptom in
On Day 1, the HR was 113.69 ± 11.51 in the AMS group and 112.37 ± 12.22 in the non-AMS group, while the SpO₂ (%) was 91.28 ± 2.24 in the AMS group and 91.42 ± 2.40% in the non-AMS group. On Day 2, the HR was 114.70 ± 11.21 in the AMS group and 111.38 ± 11.92 in the non-AMS group, while the SpO₂ was 87.59 ± 3.88% in the AMS group and 87.99 ± 4.50% in the non-AMS group. On Day 3, the HR was 119.65 ± 14.85 in the AMS group and 117.82 ± 11.90 in the non-AMS group, while the SpO₂ was 87.49 ± 6.31% in the AMS group and 87.73 ± 4.62% in the non-AMS group. There were no significant differences in HR and SpO₂ between the groups on any day.

A total of 73 (37.1%) children received medication during the trip. Among the AMS group (n = 80), 45 (56.3%) received medication. Of the non-AMS group (n = 117), 28 (23.9%) were given either acetaminophen for headache/dysmenorrhea, or metoclopramide for GI symptoms such as nausea/vomiting. Notably, we gave acetazolamide and/or dexamethasone only to children who met the criteria for AMS (headache plus an LLS ≥ 3).

**Discussion**

In Taiwan, the number of trekkers going to Xue Mountain has increased steadily over the past two decades, to more than 25,000 per year. The increase in popularity is associated with the convenience of reaching the trailhead by vehicle, good trail conditions for most of the year, and the subtropical climate of Taiwan. The number of children and adolescents going to high altitudes also increased, and it has become more important to have a better understanding on the incidence, risk factors, and common symptoms of AMS among this population. The AMS incidence increased with altitude [10,22,24,27]. In a study led by Imray et al., 91.7% of adolescents had AMS while trekking to 4,200–5,500 m [24]. Major et al. reported a similar incidence (84%) in a group of adolescents ascending to 5,200 m during a 23-day expedition [23]. In previous studies, it was unclear whether young age was a risk factor of AMS [8,9,11,15,17–18]. Gonggalanzi et al. found an AMS incidence of 36.7% among tourists (mean age = 37.2 ± 14.4 years) visiting the city of Lhasa, which is at 3,658 m [9]. Moraga et al. reported an AMS incidence of 27% in 15 tourists with a mean age of 31.9 years and 50% in 10 tourists with a mean age of 15.4 years and ascending to 3,500 m by truck [18]. In a study conducted at the Swiss high altitude research station at Jungfrau (3,450 m), AMS prevalence was lower in children than in adolescents and adults on Day 1 (8–10 hours within ascent), but not on Day 2. The cumulative prevalence of AMS was 30%, 37%, and 45% in children, adolescents, and adults, respectively (p < 0.001) [3]. In our investigation, the overall incidence of AMS among children aged 11–12 years and trekking to Xue Mountain (3,886 m) was 40.6% (Table 1). Our results are comparable to those of several previous studies of children travelling to similar altitudes [4,14]. In addition, the trekking route, sleeping altitude, and the 3-day ascent profile of Xue Mountain is similar to that of the Jade Mountain (3,952 m) trail in Taiwan. The AMS incidence in children was higher compared with adult climbers on Jade Mountain, which was reported to be 32–36% by Wang et al. and Chen et al. [11,15]. Although AMS was common among children, our study revealed that the AMS symptoms were mostly mild. For all
Fig 4. LLS for each symptom according to group on days 1–3. A—Day 1 (Qika Hut, 2,460 m): the gastrointestinal (GI) symptom score was lowest in the AMS group. No subject in the non-AMS group experienced GI symptoms at this altitude. B—Day 2 (Sanliujiu Hut, 3,100 m): the sleep disturbance score was the highest LLS in both groups (AMS: 1.29 ± 0.92, non-AMS: 0.68 ± 0.83). In the AMS group, headache and dizziness were the most severe symptoms after sleep disturbance. C—Day 3 (Sanliujiu Hut, 3,100m, after reaching the Xue Mountain summit, 3,886 m): similar to previous days, the sleep disturbance symptoms were the most severe, and GI symptoms the least severe, in both groups. On all 3 days, all of the LLS were...
occasions, 72.2% of subjects had LLSs < 2. None of the subjects in our study developed HAPE or HACE during the trekking course.

The rate of ascent and sleeping altitude also have an important impact on the incidence of AMS [1,5,6,8,13–15]. However, the sleeping altitudes differed among previous studies, as did the ascent rate and method (by foot, vehicle, or airplane [4,9,14,21]). There were also other confounding factors, such as age, gender, physical ability, and concurrent illness [1,6,8]. Therefore, it was difficult to compare these studies [3]. The purpose of our study was to evaluate the incidence and severity of AMS among children for this particular trekking route.

Although the prevalence and severity of AMS increases with a recent gain in altitude, there was evidence that subjects experienced less AMS and AMS-associated symptoms after a previous stay at high altitudes [15,20,24,28]. In teenagers travelling between 2,400 m and 5,500 m, the LLS decreased despite the subjects ascending to over 5,000 m on the 13th to 18th days of the expedition [24]. Wang et al. found that the most common site of AMS symptom development on Jade Mountain was the midway overnight hut, the Paiyun Lodge (3,400 m), but not on the summit or on the day after reaching the summit [15]. In our study, the prevalence of AMS increased from 5.6% on Day 1 to 29.4% on Day 2, and slightly decreased (23.4%) despite the effort to reach the summit of Xue Mountain (3,886 m) on Day 3 (Fig 1). There was also a trend of decreasing AMS severity at the same altitude (the mean LLS of AMS group was 4.09 ± 1.97 on Day 2, and 3.98 ± 2.42 on Day 3). The decrease in AMS prevalence and severity was mostly due to the decrease in headache prevalence and severity (Figs 3B, 3C and 4B and 4C), which was most common (46.2%) on Day 2 and decreased significantly to 31.3% on Day 3 (Table 2).

Although there was a decreasing trend in AMS prevalence and severity on Day 3, the comparison between Day 2 and Day 3 revealed no statistical difference. The effects of medication and/or acclimatization may have contributed to this finding.

Most studies found no difference in AMS incidence between males and females [4,14–15,21,25,29], while some studies reported that female sex was more susceptible to the development of AMS [8,13,20,24,27–28]. However, our study showed that males had a significantly higher incidence of AMS than females (Table 1). There was also a trend toward males’ having a higher overall prevalence of AMS-associated symptoms, with headache and fatigue being significantly more prevalent than in females (Fig 2). Higher AMS incidence was also seen in subjects with a higher BMI. Gender (male) and BMI can be considered as two independent risk factors for AMS. The pubertal stage of each child was not evaluated, since this was not part of our study protocol. Most of the girls had reached puberty (and almost all of them had periods), while half of the boys had not experienced a voice change at the time of the activity. Hormones may play a role in acclimatization to high altitude, but the underlying mechanism remains unknown and requires further investigation.

A total of 73 (37.1%) children received medication during the whole trekking course. Among the AMS group (n = 80), 45 (56.3%) received medication. Of the non-AMS group (n = 117), 28 (23.9%) received medication (either acetaminophen for headache/dysmenorrhea or metoclopramide for nausea/vomiting). We gave acetazolamide and/or dexamethasone only to children who met the criteria for AMS (headache and an LLS ≥ 3) so that the administration
of acetazolamide and/or dexamethasone would not affect the overall incidence of AMS. The use of acetaminophen for treating headache may have been a confounding factor, since headache is required for an AMS diagnosis. Less than 30% of the children experienced GI symptoms, which were also the least prevalent AMS-associated symptom. The administration of metoclopramide may also have affected the result; however, the effect was minimal. In addition, we studied an annual trekking activity of primary school children. To ensure the safety of the children, we administered treatments immediately if needed. Due to ethical concerns, we could not permit the possibility of the children experiencing severe AMS for research purposes. Administering medication may lead to underestimating of the overall incidence of AMS, as well as the daily prevalence and severity of AMS. Of note, one subject had taken herbal medicine before activity, and one took Rhodiola as a prophylaxis to AMS. In a recent randomized double-blind crossover trial, Rhodiola extract was ineffective in reducing the incidence or severity of AMS compared with a placebo [30].

Headache was the most prevalent AMS-associated symptom in most previous studies [8,15,17,24]. On the other hand, Bloch et al. and Gonggalanzi et al. reported fatigue to be the most common symptom at around 3,500 m [4,9]. In our study, 115 (58.4%) subjects experienced at least one episode of headache during the trip (Table 2). Furthermore, 35 of these did not have other symptoms or an LLS ≥ 3. Twenty-six (13.2%) subjects had an LLS ≥ 3 but did not meet the criteria for AMS. These may indicate early or mild AMS. The most common symptom of AMS in our study was sleep disturbance, followed by dizziness, headache, fatigue, and GI symptoms (Table 2). We found that up to 74.6% of children had sleep disturbance during the trip (Table 2), which was a higher rate than in other studies [8,22,24]. Similar findings were also reported in children and adult trekkers on Jade Mountain (3,952 m) [15,25]. This may be due to the crowded environment of Qika Hut and Sanlijiu Hut, which contain a maximum of 130 and 110 people, respectively, in two large rooms. Different mountain trekking groups have different times of arrival and departure, and the noise may disturb the sleep of other teams. A similar situation was observed by Horiuchi et al. on Mount Fuji (3,776 m), where climbers staying overnight at a lodge were more likely to have poor sleep quality and a reduction in total sleep time [29]. Moreover, it was likely that cold weather, and not being accustomed to sleeping bags, contributed to the higher sleep disturbance scores. Several children also reported that they missed their homes and parents at night, which made it difficult to fall asleep. The other AMS-associated symptoms may be the result of travel fatigue and physical effort expended while trekking. Children traveled in vehicles for 6 hours, and trekked for 2 hours on Day 1; thus, they were likely to have symptoms, such as fatigue or nausea, due to travel that might have imitated the milder forms of AMS. These special considerations, together with other factors, such as boredom and disruption of daily routine, among this population have also been reported in previous literature [1,26].

Finally, medical teams were on standby and trekked with each group. We also established a “high-altitude clinic” at Sanlijiu Hut (3,100 m), where a senior emergency physician and an experienced nurse were stationed to deliver medical services at night during the trek. Once a subject developed AMS or other unbearable symptoms, the physicians would evaluate and treat them immediately. Early detection of AMS might have prevented the children from experiencing serious complications. This was the first established medical service provided for children at high altitudes in Taiwan. More than 2,000 children have benefited from the medical service since 2009 [25,31]. Although we measured HR and SpO2, there were no significant differences in either measure between the AMS and non-AMS group on any day. We did not discuss this in this article, as the effect of acetazolamide may have influenced these physiologic measures.
Study limitations

The major limitation of this study was the medication administrated during the trip. This may have led to underestimating of the overall incidence of AMS. For ensuring the children's safety, we treated subjects with AMS and other complaints. We administrated acetaminophen for headache, metoclopramide for nausea/vomiting, acetazolamide only when subjects met the criteria for AMS, and dexamethasone for severe AMS. The medication was given at any place or time along the trekking route once a child developed AMS or other symptoms. Since the medication treatment plan was not part of our study protocol, we did not analyze the time of treatment or any subsequent effects of medication. Thus, the daily AMS prevalence and severity were also possibly subject to bias and underestimation. Second, four subjects were unwilling to ascend to the summit or (because of other discomfort that did not meet the criteria for AMS) descended to Qika Hut. These four non-AMS subjects were excluded from the analysis of AMS prevalence and LLS on Day 3, which may also have affected the results. Third, the mean BMI of the males (19.79 ± 4.29 kg/m²) was slightly higher than that of the females (18.83 ± 2.75 kg/m²); however, there was no significant statistical difference (p = 0.453). Although we have adjusted the associations among gender, BMI, and AMS using logistic regression and z-scores, this may still have affected the results regarding the association between BMI and AMS prevalence. Finally, it is possible that some confounding factors, such as poor sleep conditions, changes in routine and diet lifestyle, and absence of family members may have accounted for the symptoms such as sleep disturbance.

Conclusions

The AMS incidence among children trekking to Xue Mountain was 40.6%. Gender (male) and a higher BMI can be considered as two independent risk factors for AMS. Although AMS was common among children, the symptoms were mostly mild. Sleep disturbance was the most common symptom. The effects of medication and/or acclimatization may explain the lower incidence of AMS and headache on Day 3 compared to Day 2.

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References

1. Pollard AJ, Niermeyer S, Barry P, Bärtsch P, Berghold F, Bishop RA, et al. Children at high altitude: an international consensus statement by an ad hoc committee of the International Society for Mountain Medicine, March 12, 2001. High Alt Med Biol. 2001; 2(3):389–403. https://doi.org/10.1089/15270290152608561 PMID: 11682018

2. Stokes SH, Kalson NS, Frost H, Tyrrell-Marsh I, Whitehead AG, Davies A, et al. Adolescents with altitude: young people do well on Kilimanjaro. Arch Dis Child. 2009; 94(7):562–563.

3. Kriemler S, Bürgi F, Wick C, Wick B, Keller M, Wiget U, et al. Prevalence of acute mountain sickness at 3500 m within and between families: a prospective cohort study. High Alt Med Biol. 2014; 15(1):28–38. https://doi.org/10.1089/ham.2013.1073 PMID: 24559431

4. Bloch J, Duplain H, Rimoldi SF, Stuber T, Kriemler S, Allemann Y, et al. Prevalence and time course of acute mountain sickness in older children and adolescents after rapid ascent to 3450 meters. Pediatrics. 2009; 123(1):1–5. https://doi.org/10.1542/peds.2008-0200 PMID: 19117853

5. Bärtsch P, Swenson ER. Acute high-altitude illnesses. N Engl J Med. 2013; 368(24):2294–2302. https://doi.org/10.1056/NEJMcp1214870 PMID: 23758234

6. Neumann K. Children at altitude. Travel Med Infect Dis. 2007; 5(2):138–141. https://doi.org/10.1016/j.tmaid.2006.01.011 PMID: 17298923

7. Roach RC, Bärtsch P, Hackett PH, Oelz O. The Lake Louise acute mountain sickness scoring system. Hypoxia and molecular biology. Burlington VT: Queen City Press. 1993;272–274.

8. Honigman B, Theis MK, Koziol-McLain J, Roach RC, Yip R, Houston C, et al. Acute mountain sickness in a general tourist population at moderate altitudes. Ann Intern Med. 1993; 118(8):587–592. PMID: 8452324

9. Gongglangzi, Labasangzhu, Nafstad P, Stigum H, Wu T, Haldorsen ØD, et al. Acute mountain sickness among tourists visiting the high-altitude city of Lhasa at 3658 m above sea level: a cross-sectional study. Archives of Public Health. 2016; 74(23):1–7.

10. Imray CH, Booth A, Wright A, Bradwell A. Acute altitude illness. BMJ. 2011; 343:1–10.

11. Chen HC, Lin WL, Wu JY, Wang SH, Chiu TF, Weng YM, et al. Change in oxygen saturation does not predict acute mountain sickness on Jade Mountain. Wilderness Environ Med. 2012; 23(2):122–127. https://doi.org/10.1016/j.wem.2012.03.014 PMID: 22656657

12. Weng YM, Chiu YH, Lynn JJ, Li WC, Wang SH, Kao WF, et al. Different duration of high-altitude pre-exposure associated with the incidence of acute mountain sickness on Jade Mountain. Am J Emerg Med. 2013; 31(7):1113–1117. https://doi.org/10.1016/j.ajem.2013.03.028 PMID: 23688567

13. Murdoch DR. Altitude illness among tourists flying to 3740 meters elevation in the Nepal Himalayas. J Trav Med. 1995; 2(4):255–256.

14. Pradhan S, Yadav S, Neupane P, Subedi P. Acute mountain sickness in children at 4380 meters in the Himalayas. Wilderness Environ Med. 2009; 20(4):359–363. https://doi.org/10.1580/1080-6032-0.004.0359 PMID: 20030445

15. Wang SH, Chen YC, Kao WF, Lin YJ, Chen JC, Chiu TF, et al. Epidemiology of acute mountain sickness on Jade Mountain, Taiwan: an annual prospective observational Study. High Alt. Med. Biol. 2010; 11(1):43–49. https://doi.org/10.1089/ham.2009.1063 PMID: 20367488

16. Rexhaj E, Garcin S, Rimoldi SF, Duplain H, Stuber T, Allemann Y, et al. Reproducibility of acute mountain sickness in children and adults: a prospective study. Pediatrics. 2011; 127(6):e1445–8. https://doi.org/10.1542/peds.2010-3756 PMID: 21536612

17. Hackett PH, Rennie D, Levine HD. The incidence, importance, and prophylaxis of acute mountain sickness. Lancet. 1976; 308(7996):1149–1155.

18. Moraga FA, Osorio JD, Vargas ME. Acute mountain sickness in tourists with children at Lake Chungara (4400 m) in northern Chile. Wilderness Environ Med. 2002; 13(1):31–35. PMID: 11929059

19. Yaron M, Niermeyer S, Lindgren KN, Honigman B. Evaluation of diagnostic criteria and incidence of acute mountain sickness in preverbal children. Wilderness Environ Med. 2002; 13(1):21–26. PMID: 11929057

20. Lawrence JS, Reid SA. Risk determinants of acute mountain sickness and summit success on a 6-day ascent of Mount Kilimanjaro (5895 m). Wilderness Environ Med. 2016; 27(1):78–84. https://doi.org/10.1016/j.wem.2015.11.011 PMID: 26948557
21. Moraga FA, Pedreros CP, Rodríguez CE. Acute mountain sickness in children and their parents after rapid ascent to 3500 m (Putre, Chile). Wilderness Environ Med. 2008; 19(4):287–292. https://doi.org/10.1580/06-WEME-BR-084.1 PMID: 19099320

22. Maggiorini M, Bühler B, Walter M, Oelz O. Prevalence of acute mountain sickness in the Swiss Alps. BMJ. 1990; 301(6756):853–855. PMID: 2282425

23. Major SA, Hogan RJ, Yeates E, Imray CH. Peripheral arterial desaturation is further exacerbated by exercise in adolescents with acute mountain sickness. Wilderness Environ Med. 2012; 23(1):15–23. https://doi.org/10.1016/j.wem.2011.12.008 PMID: 22441084

24. Imray CH, Kennedy CH, Pattinson K, Brearey S, Wright A. Self-assessment of acute mountain sickness in adolescents: a pilot study. Wilderness Environ Med. 2004; 15(3):202–206. PMID: 15473461

25. Chan CW, Lin YC, Chiu YH, Weng YM, Li WC, Lin YJ, et al. Incidence and risk factors associated with acute mountain sickness in children trekking on Jade Mountain, Taiwan. J Trav Med. 2016; 23(1):1–5.

26. Theis MK, Honigman B, Yip R, McBride D, Houston CS, Moore LG. Acute mountain sickness in children at 2835 meters. Am J Dis Child. 1993; 147(2):143–145. PMID: 8427234

27. Dallimore J, Rowbotham EC. Incidence of acute mountain sickness in adolescents. Wilderness Environ Med. 2009; 20(3):221–224. https://doi.org/10.1580/07-WEME-OR-119R3.1 PMID: 19737028

28. MacInnis MJ, Carter EA, Freeman MG, Pandit BP, Siwakoti A, Subedi A, et al. A prospective epidemiological study of acute mountain sickness in Nepalese pilgrims ascending to high altitude (4380 m). PLoS One. 2013; 8(10):e75644. https://doi.org/10.1371/journal.pone.0075644 PMID: 24130729

29. Horiuchi M, Endo J, Akatsuka S,Uno T, Jones TE. Prevalence of acute mountain sickness on Mount Fuji: a pilot study. J Trav Med. 2016; 23(4):1–5.

30. Chiu TF, Chen LL, Su DH, Lo HY, Chen CH, Wang SH, et al. Rhodiola crenulata extract for prevention of acute mountain sickness: a randomized, double-blind, placebo-controlled, crossover trial. BMC Complement Altern Med. 2013;31; 13:298. https://doi.org/10.1186/1472-6882-13-298 PMID: 24176010

31. Wang SH, Huang KF. Medical service for the children high-altitude group trekking activity. Am J Emerg Med. 2013; 31(7):1142–1144. https://doi.org/10.1016/j.ajem.2013.03.043 PMID: 23702062