Variation of vital signs with potential to influence the performance of qSOFA scoring in the Ethiopian general population at different altitudes of residency: A multisite cross-sectional study

Jonas Früh1‡, Andre Fuchs1,2‡, Tafese Beyene Tufa1,2,3, Loraine Früh1, Zewdu Hurissa3, Hans Martin Orth1,2, Johannes Georg Bode1,2, Kirsten Alexandra Eberhardt1, Dieter Häussinger1,2, Torsten Feldt1,2*

1 Hirsch Institute of Tropical Medicine, Asella, Ethiopia, 2 Department of Gastroenterology, Hepatology and Infectious Diseases, University Hospital, Heinrich Heine University, Düsseldorf, Germany, 3 College of Health Sciences, Arsi University, Asella, Ethiopia, 4 Bernhard Nocht Institute for Tropical Medicine, I. Department of Medicine, University Medical Center Hamburg-Eppendorf, Hamburg, Germany

‡ These authors contributed equally to this work and share first authorship.

* torsten.feldt@med.uni-duesseldorf.de

Abstract

Introduction

The physiological range of different vital signs is dependent on various environmental and individual factors. There is a strong interdependent relationship between vital signs and health conditions. Deviations of the physiological range are commonly used for risk assessment in clinical scores, e.g. respiratory rate (RR) and systolic blood pressure (BP.sys) in patients with infections within the quick sequential organ failure assessment (qSOFA) score. A limited number of studies have evaluated the performance of such scores in resource-limited health care settings, showing inconsistent results with mostly poor discriminative power. Divergent standard values of vital parameters in different populations, e.g. could influence the accuracy of various clinical scores.

Methods

This multisite cross-sectional observational study was performed among Ethiopians residing at various altitudes in the cities of Asella (2400m above sea level (a.s.l.)), Adama (1600m a. s.l.), and Semara (400m a.s.l.). Volunteers from the local general population were asked to complete a brief questionnaire and have vital signs measured. Individuals reporting acute or chronic illness were excluded.

Results

A positive qSOFA score (i.e. ≥2), indicating severe illness in patients with infection, was common among the studied population (n = 612). The proportion of participants with a
positive qSOFA score was significantly higher in Asella (28.1%; 55/196), compared with Adama, (8.3%; 19/230; p<0.001) and Semara (15.1%; 28/186; p = 0.005). Concerning the parameters comprised in qSOFA, the thresholds for RR (≥22/min) were reached in 60.7%, 34.8%, and 38.2%, and for BPsys (≤100 mmHg) in 48.5%, 27.8%, and 36.0% in participants from Asella, Adama, and Semara, respectively.

**Discussion**

The high positivity rate of qSOFA score in the studied population without signs of acute infection may be explained by variations of the physiological range of different vital signs, possibly related to the altitude of residence. Adaptation of existing scores using local standard values could be helpful for reliable risk assessment.

**Introduction**

The normal range of vital signs depends on various environmental and individual factors, and there is a strong interdependent relationship between vital signs and health condition [1, 2]. Deviations of physiological parameters, including respiratory rate (RR) and blood pressure (BP), from the normal range are used in several clinical scores, e.g. the qSOFA (quick sequential organ failure assessment) score. The qSOFA score has been developed as a tool for the identification of patients who are at greater risk for a poor outcome among patients with suspected infection outside the intensive care unit (ICU) [3]. It uses three criteria, assigning one point for low systolic blood pressure (BPsys ≤100 mmHg), high respiratory rate (≥22 breaths per min), or altered mentation (Glasgow coma scale (GCS) <15). Thus, the score ranges from 0 to 3 points and patients with 2 or more qSOFA points are likely to be septic and are at high risk for an unfavorable outcome [3]. Key advantages of the qSOFA are the easy and universal availability of the comprised parameters in clinical settings. A limited number of studies evaluating the performance of sepsis scores in low-resource health care settings have shown inconsistent results with mostly poor discriminative power and high variability across different study sites and settings [4–10]. Previous studies conducted elsewhere have mostly found lower BP, higher resting heart rate (HR) and lower peripheral oxygen saturation (SpO₂) in people living at high altitudes compared to lowlanders [11–13], although some studies have also documented a rise in BP with increasing altitude [14, 15].

Divergent physiological ranges of the applied vital signs in different populations could be a reason for heterogeneity of the performance of vital sign-dependent clinical scores as the qSOFA score in different populations. A possible cause of such heterogeneities may be attributed to adaptation mechanisms of the local population to higher altitudes. Thus, we investigated the potential influence of variations within the physiological range of vital signs in the general healthy population residing at different altitudes in Ethiopia on the performance of the qSOFA score.

**Materials and methods**

The study has been approved by the appropriate Ethical Review Committee (ERC) of the College of Health Sciences, Arsi University, Asella, Ethiopia (project number: A/CHS/RC/72/18). All volunteer healthy participants gave verbal consent after study procedures were thoroughly explained in local language by the study team before data acquisition. The method of verbal...
informed consent was approved by the Ethical Review Board, considering the high rate of illiteracy and that no invasive procedures were performed. The data were analyzed anonymously. For this multisite cross-sectional observational study, we selected three study sites in Ethiopian cities at different altitudes. Asella was the study site located at the highest altitude (2400 m a.s.l.). Second, we aimed for a study site at the lowest possible altitude within the country, inhabited by a population with similar descent. Since Ethiopia is a landlocked country with large shares of highlands and the areas with the lowest altitude in the country are barren, hostile deserts where hardly any people live, no inhabited area at sea level was available. Therefore, Semara, as one of the cities of the country located at the lowest altitude (400 m a.s.l.), was selected as study site. Third, for comparison at mid-level altitude, the city of Adama (1620 m a.s.l.) was chosen. In order to investigate comparable populations at the respective study centers and therefore avoid selection errors, we selected urban centers at the respective altitudes as study sites. Reliable data on socioeconomic differences between the study sites were not available.

The study was conducted between December 2018 and March 2019 among adult alert volunteers with a minimum age of 16 years (Fig 1). The selection of volunteers was randomly made among pedestrians on the street at each study site. Participants were included for measurements during daytime for two subsequent days in order to achieve a sufficiently large sample size. A mobile medical unit was set up in a tent in a busy area downtown at each of the study sites and random passers-by were invited to have their vital signs checked and to participate in our investigation. Apart from the age limit and the residency in the assigned area, no other initial study eligibility criterion was applied. This approach of random selection of subjects in a busy area of the respective city, populated by local people who pursue their ordinary activities was chosen in order to achieve inclusion of a representative cross-section of the local population. The physiological parameters like body temperature (T), BP_{sys}, RR, SpO₂ and HR were measured non-invasively with medical infrared thermometers, photo optic finger clip pulse oximeters, and aneroid sphygmomanometers. Height and weight were measured from which body mass index (BMI) was calculated by two trained nurses. Prior to these measurements, the volunteers were asked to come to rest in a designated waiting area for not less than 5 minutes, which is in conformity with official recommendations and international guidelines [16, 17]. To avoid measurement errors in pulse oximetry, the measurement was performed only on clean fingers without nail polish. The participants were asked to rest their arm during the measurement. In addition, data on socio-demographic background, current health condition, as well as chronic diseases were collected using a standardized questionnaire. In order to rule out potential influences on vital parameters by medical conditions, participants with acute or previously known chronic illnesses were excluded a posteriori. The qSOFA was calculated using the cutoffs of ≥22/min for RR, and ≤100 mmHg for (BP_{sys}). Since all participants were fully conscious and responsive during the study procedures, the GCS, assessing mental alteration, was graded unimpaired (15 points) in all participants. Continuous variables were expressed as median (interquartile range, IQR). Post-hoc sample size calculation and power analysis were performed using the approach by Cohen, J. and the R-Package “pwr” [18]. Multi-group comparisons were done using the Kruskal-Wallis Test or one-way ANOVA. Additionally, pairwise comparisons between group levels at the different study sites were performed and adjusted for multiple comparisons using the false discovery rate approach. Categorical variables were compared using either the χ² test or the Fisher exact test, as appropriate. To evaluate the association of the place of residency with the qSOFA score, a multiple ordinal regression model was used and adjusted for other covariates. An alpha of 0.05 was determined as the cutoff for significance. All statistical analyses were performed using R (version 3.6.3, R Foundation for Statistical Computing, Vienna, Austria).
A total of 612 participants were included in the final analysis. Forty-nine Participants (7.4% of the original study collective) were excluded prior to the final data analysis due to an acute (0.9%) or previously known chronic illnesses (6.5%), primarily arterial hypertension and lung diseases, including tuberculosis (see Fig 2). The mean age of included participants was 31.5 ±12.8 years and 30.6% of them were female (Table 1). All participants were fully alert and oriented and GCS was graded 15/15.

For details on vital parameters see Table 2 and Fig 3. Interestingly, the majority of vital parameters differed significantly across the sites (Table 2). The median respiratory rate was significantly higher in Asella compared with the other sites (22 [IQR 20–24], 20 [IQR 18–22], and 21 [IQR 19–23] /min in Asella, Adama, and Semara, respectively), whereas the median systolic blood pressure was lower in participants from Asella (110 [IQR 100–113], 110 [IQR 100–120], and 110 [IQR 100–120] mmHg in Asella, Adama, and Semara respectively).
regard to the qSOFA, RR threshold was reached in 60.7% (77/196), 34.8% (80/230) and 38.2% (71/186) in Asella, Adama and Semara, and BP threshold was reached in 48.5% (95/196), 27.8% (64/230) and 36.0% (67/186), respectively. Remarkably, in Asella, at high altitude, the median RR in the analyzed healthy population reached the qSOFA score threshold (Fig 3). As presented in Table 3, across all sites 16.7% (102/612) of participants scored 2 points in the qSOFA score (RR $\geq 22$ and BP$_{sys} \geq 100$ mmHg). In particular, the qSOFA score reached 2 points in 28.1% (55/196) of participants in Asella, in 8.3% (19/230) of participants in Adama and 15.1% (28/186) of participants in Semara (see Fig 4). Notably, the distribution of the qSOFA score in Asella was significantly different from Adama and Semara ($p<0.001$, Table 3 and Fig 4). Also, when adjusting for the covariates age, sex and BMI in the multiple ordinal regression model, Asella as area of residency was significantly associated with an elevated

![Fig 2. Flow chart of inclusions and exclusions at the different study sites. A total of 661 participants were recruited. After screening for any sickness or disease, the remaining participants used for the analysis was 612.](https://doi.org/10.1371/journal.pone.0245496.g002)

Table 1. Demographic parameters.

|                  | Total (n = 612) | Asella (n = 196) | Adama (n = 230) | Semara (n = 186) | p-value (group differences) |
|------------------|----------------|------------------|----------------|----------------|-----------------------------|
| Age in years     | Mean ± SD      | 31.5 ± 12.7      | 26.7 ± 10.6    | 36.0 ± 13.4    | 31.1 ± 12.1                | <0.001                     |
| Sex              | male, n (%)    | 425 (69.4)       | 125 (63.8)     | 148 (64.3)     | 152 (81.7)                 | <0.001                     |
|                  | female, n (%)  | 187 (30.6)       | 71 (36.2)      | 82 (35.7)      | 34 (18.3)                  |                            |
| Ethnicity        | Oromo, n (%)   | 293 (47.9)       | 155 (79.1)     | 132 (57.4)     | 6 (3.2)                    | <0.001                     |
|                  | Amhara, n (%)  | 179 (29.2)       | 21 (10.7)      | 52 (22.6)      | 106 (57.0)                 |                            |
|                  | Afar, n (%)    | 58 (9.5)         | 0 (0)          | 0 (0)          | 58 (31.2)                  |                            |
|                  | Gurage, n (%)  | 24 (3.9)         | 7 (3.6)        | 17 (7.4)       | 0 (0)                      |                            |
|                  | Tigray, n (%)  | 14 (2.3)         | 3 (1.5)        | 8 (3.5)        | 3 (1.6)                    |                            |
|                  | Wolayita, n (%)| 8 (1.3)          | 0 (0)          | 2 (0.9)        | 6 (3.2)                    |                            |
|                  | Silete, n (%)  | 7 (1.1)          | 4 (2.0)        | 3 (1.3)        | 0 (0)                      |                            |
|                  | Somali, n (%)  | 1 (0.2)          | 0 (0)          | 0 (0)          | 1 (0.5)                    |                            |
|                  | Other, n (%)   | 17 (2.8)         | 2 (1.0)        | 10 (4.3)       | (5 (2.7)                   |                            |
|                  | not stated, n (%)| 11 (1.8)      | 4 (2.0)        | (2.6)          | 1 (0.5)                    |                            |

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qSOFA score compared to the location Adama (aOR 3.26 [2.20–4.86], p < 0.001, Table 4), whereas no difference between Semara and Adama was observed. Furthermore, male gender was significantly associated with a lower qSOFA score (p = 0.004).

**Discussion**

Within the cohort of healthy adult volunteers, the high proportion of individuals reaching the cutoff for a positive qSOFA score was surprising. The qSOFA score has been developed as an easy-to-use bedside score in order to quickly identify individuals at risk of a poor outcome among patients with an infection [19]. In previous studies, positive qSOFA criteria showed a similar prognostic significance compared with more complex tools as the SOFA, MEDS (Mortality in Emergency Department Sepsis) or APACHE II (Acute Physiology And Chronic Health Evaluation II) scores [20]. The specificity to predict mortality among patients with an infection of the qSOFA score in sub-Saharan African cohorts was reported to be 82% (95% Confidence Interval [CI] 76–88) and 81% (95% CI 78–85) while sensitivity was much lower (55% [95% CI 23–83] and 44% [95% CI 33–55], respectively) [5, 7].

All study participants were alert pedestrians, were not apparently mentally altered, as assessed during study procedures, and did not report to suffer from chronic illness. Thus, all patients were considered to have an unimpaired GCS of 15. The RR was significantly higher

![Graphs showing vital signs](https://doi.org/10.1371/journal.pone.0245496.g003)
and the SpO₂ significantly lower in Asella, the site situated at the highest altitude a.s.l., whereas there was no significant difference between the sites at lower altitudes. Although the rate of positive qSOFA was also remarkably high at those sites, we postulate that the high altitude of 2400 m a.s.l. might contribute to the high rate of unspecific positive qSOFA score values. As it has been suggested by other authors [5], an adaption of existing scores to various settings might be necessary to improve the performance. Alternatively, to improve the impaired performance of clinical scores at different altitudes, a constant conversion factor could possibly be derived from future cohort analyses.

Table 3. Proportions of the qSOFA at the different study sites.

| qSOFA-score, n (%) | Total (n = 612) | Asella (n = 196) (A) | Adama (n = 230) (B) | Semara (n = 186) (C) | p-value (group differences) | p-value (Single group comparisons) |
|-------------------|-----------------|---------------------|---------------------|---------------------|-----------------------------|-----------------------------------|
|                   |                 | A|B | A|C | B|C | A|B | A|C | B|C |
| 0                 | 218 (35.6)      | 37 (18.9)            | 105 (45.7)          | 76 (40.9)           | <0.001                      | <0.001 <0.001 0.089               |
| 1                 | 292 (47.7)      | 104 (53.1)           | 106 (46.1)          | 82 (44.1)           |                             |                                   |
| 2                 | 102 (16.7)      | 55 (28.1)            | 19 (8.3)            | 28 (15.1)           |                             |                                   |

qSOFA: Quick Sequential Organ Failure Assessment

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Fig 4. Proportions of qSOFA categories according to study site.

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The finding of a positive qSOFA score in the normal population was mostly common in Asella, located at 2400 m a.s.l. It has been shown elsewhere, that the qSOFA can be a reliable predictor of mortality, also in resource-limited settings [6, 21, 22] but perhaps not at extreme elevations. Our hitherto deviating results can only partly be explained by comparatively high altitudes in Ethiopia, since also at lower altitude, in Semara at 400 m a.s.l., a positive qSOFA was frequently found and was present in 15.1% of the local population. However, the median RR in the normal healthy population at Asella reached the threshold of the qSOFA score and even though the median BP\textsubscript{sys} was 110 mmHg at all three sites, there was a significant difference in the IQR between the three sites, being lowest in participants from Asella. These findings explain the high rate of healthy individuals reaching a positive qSOFA. On the other hand, body temperature and HR, both parameters not included in the qSOFA score, showed no or hardly any differences between the study sites at different altitudes.

A limited specificity of the score in the studied population has already been described in previous studies conducted in countries with limited resources in Sub-Saharan Africa [5, 10, 23, 24] and may be partially explained by deviations of the standard values of different vital signs like RR or BP\textsubscript{sys} although optimal thresholds remain uncertain. Our results support previous findings questioning the accurate applicability of the qSOFA not only in resource-limited settings, but also in more developed settings [8, 25, 26]. With a high proportion of positive qSOFA criteria in a normal population, the score fails to serve as specific tool for the identification of septic patients at risk for adverse outcomes.

Our findings might be limited due to the fact that our data was recorded as a single cross-sectional assessment and do not reflect physiological variation of the parameters within individuals. Nevertheless, also vital parameters used to calculate sepsis scores are usually assessed once at a certain time point in clinical settings. The analysis of multiple individuals within our cohort reduces the risk for selection bias. Since all study procedures were performed using volunteers, the results could be influenced by volunteer bias. However, since the sampling methods did not differ between the study sites, this bias cannot explain the apparent differences between the different study groups. No extrinsic motivation in form of any compensation was offered for participants. The conducted post-hoc power analysis indicated a sufficiently large group size to test the study objective.

Participants reporting any form of chronic disease were excluded from further analysis to rule out the possibility of changes in vital signs caused by illness. However, Patel et al. were able to show that self-reporting leads to limitations in the reliability of chronic disease detection [27]. Thus, among the participants classified as healthy in this study, there may have been

| Table 4. Factors associated with the qSOFA at the different study sites. |
|-------------------------|---------|-----------|--------|
| Covariates              | qSOFA  | 95% CI    | p-value |
| Age in years            | 0.99   | (0.98–1.01)| 0.247  |
| Female                  | 1      |           |        |
| Male                    | 0.61   | (0.43–0.86)| 0.004  |
| BMI in kg/m\textsuperscript{2} | 0.98   | (0.94–1.02)| 0.295  |
| Adama                   | 1      |           |        |
| Asella                  | 3.26   | (2.20–4.86)| <0.001 |
| Samara                  | 1.32   | (0.89–1.96)| 0.171  |

qSOFA: Quick Sequential Organ Failure Assessment, aOR: adjusted Odds Ratios, CI: Confidence Interval, BMI: Body Mass Index

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individuals potentially suffering from a chronic disease. Since this limitation applies equally to all study centers, no distortion of the study results in the comparison of the study centers is to be expected.

The European Society of Cardiology, the American Heart Association and others suggest having a patient rest for 5 minutes before measuring BP [16, 17, 28]. This approach was followed during our study. However, there are other data which indicate that a longer resting time of 10 or even 25 minutes might be necessary for reliable stabilization of BP [29, 30]. To circumvent a resulting error, no volunteers apparently exhausted by physical activity were included and the same procedure was followed at all study sites.

Also, the differences among demographic parameters such as age, gender and BMI between the study sites have to be considered as possible limitation of the study, but for this very reason the ordinal regression model was adjusted using these parameters and it was shown that Asella as place of residence is associated with a higher probability of a false positive qSOFA score, regardless of age, gender and BMI. Possibly, this finding could also be confounded by different socio-economic and environmental conditions (e.g. climate, air pollution) at the three study sites with Adama as a metropolis, Asella as a major district town and Semara as rather remote city. The varying BMI at the different study populations could reflect the respective economic strength and might be interpreted as an indication of different lifestyles of the population at the study sites. However, this assumption is based on personal observations and reliable data to support this hypothesis are insufficient.

**Conclusion**

Our study indicates that the applicability of the qSOFA score or other clinical scores based on examination of the vital signs BP and RR may be adversely affected by shifts in the range of normal values of the vital signs, e.g. as an adaptation mechanism for altitude. As it has previously been suggested by the international Sepsis-3 Task force, the qSOFA needs further investigation and validation especially in resource-limited health care settings [19]. High altitude might potentially be a relevant factor, since large populations of around 389 million people live in altitudes above 1,500 m, especially in Mexico, South America, the South-Central Asian Highlands, and Eastern Africa (Kenya, Ethiopia) [31]. Adaption of scores based on physiological parameters, as the qSOFA, according to local variances could improve the performance of these scores.

**Supporting information**

S1 Dataset. Minimal dataset.
(XLSX)

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**Author Contributions**

**Conceptualization**: Jonas Früh, Andre Fuchs, Tafese Beyene Tufa, Loraine Früh, Hans Martin Orth, Torsten Feldt.

**Data curation**: Jonas Früh, Loraine Früh, Zewdu Hurissa.
Formal analysis: Jonas Früh, Andre Fuchs, Tafese Beyene Tufa, Kirsten Alexandra Eberhardt, Torsten Feldt.

Investigation: Jonas Früh, Loraine Früh.

Methodology: Jonas Früh, Andre Fuchs, Tafese Beyene Tufa, Zewdu Hurissa, Torsten Feldt.

Project administration: Andre Fuchs, Tafese Beyene Tufa, Loraine Früh, Zewdu Hurissa, Torsten Feldt.

Resources: Torsten Feldt.

Software: Kirsten Alexandra Eberhardt.

Supervision: Andre Fuchs, Tafese Beyene Tufa, Hans Martin Orth, Johannes Georg Bode, Dieter Häussinger, Torsten Feldt.

Validation: Andre Fuchs, Hans Martin Orth, Kirsten Alexandra Eberhardt, Torsten Feldt.

Visualization: Jonas Früh, Kirsten Alexandra Eberhardt.

Writing – original draft: Jonas Früh, Andre Fuchs.

Writing – review & editing: Andre Fuchs, Tafese Beyene Tufa, Hans Martin Orth, Johannes Georg Bode, Kirsten Alexandra Eberhardt, Dieter Häussinger, Torsten Feldt.

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