Variation in the Low-frequency Fluctuation of the Cortex Among Patients with Paradoxical Insomnia: A Resting-State Functional Magnetic Resonance Imaging Study

Mortaza Afshani and Habibolah Khazaie

1Institute of Medical Science and Technology, Shahid Beheshti University, Tehran, Iran
2Psychiatry Department, Sleep Disorders Research Center, Kermanshah University of Medical Sciences, Kermanshah, Iran

*Corresponding author: Psychiatry Department, Sleep Disorders Research Center, Kermanshah University of Medical Sciences, Kermanshah, Iran. Tel: +98-38265255, Email: hakhazaie@gmail.com

Received 2022 August 04; Revised 2022 November 19; Accepted 2022 November 22.

Abstract

Background: Paradoxical insomnia is one of the most prevalent subtypes of insomnia disorder and its symptom includes a discrepancy between subjective and objective sleep measures.

Objectives: This study aimed to evaluate the amplitude differences of low-frequency fluctuation as a possible local characteristic in paradoxical insomnia by resting-state functional MRI among patients with paradoxical insomnia and healthy subjects.

Methods: A t-test was used in this case-control study to investigate possible changes in low-frequency fluctuation amplitude in 15 paradoxical insomnia patients and 48 healthy subjects.

Results: The results were far from the chance level by removing nuisance variables and controlling for multiple comparisons. In addition, there was a decrease in the amplitude of low-frequency fluctuation among people with paradoxical insomnia compared to the healthy group in the right region of the superior parietal lobule and precuneus regions.

Conclusions: Based on these results, brain signals appear to play an important role in the pathophysiology of paradoxical insomnia, and further research may identify the mechanism behind this type in the future.

Keywords: Insomnia Disorder, Paradoxical Insomnia, Functional Imaging, Low-frequency Fluctuation

1. Background

International classification of sleep disorders (ICSD), second edition defines insomnia disorder as difficulty in starting, maintaining, or stabilizing sleep when there is ample opportunity (1, 2). The patient usually reports daily disturbances associated with nocturnal sleep problems, such as fatigue, difficulty concentrating, impaired social and work functioning, unstable mood, daytime sleepiness, decreased motivation, physical symptoms, or excessive worry about sleep (3-5). There are some clinical and pathological subtypes of ID, including psychophysiological insomnia, idiopathic insomnia, and paradoxical insomnia (Par-I) (5). The symptoms of the Par-I condition include subjective complaints of severe insomnia, which do not correlate with objective sleep measures, such as polysomnography (1,3, 5, 6). Par-I patients usually underestimate their total sleep time, the period between sleep onset and waking, and the period between going to bed and sleeping (7, 8).

Recently, Krystal et al. compared the power of brain signals in different frequency bands and observed significant differences in alpha, beta, theta, and gamma bands between patients with Par-I and healthy individuals (9). Moreover, Harvey and Tang (10) hypothesized three possible mechanisms to potentially explain the discrepancy between subjective reports and objective measures as follows: (1) short-term awakenings during the night and early morning; (2) worry and selective attention to sleep-related concerns; and (3) misunderstand sleep as wakefulness. The misperception of sleep in Par-I may also be caused by the presence of local wakefulness and sleep (11, 12).

According to research on Par-I, local processes can have an essential role in the misperception of the sleep state. The purpose of this study was to evaluate possible differences in ALFF as a reliable marker of local processing (13), obtained by resting-state fMRI among 15 people with Par-I and 48 healthy people (HC).
2. Objectives

This study seeks to check whether two Par-I and HC groups significantly differ in the extracted ALFF.

3. Methods

A total of 79 participants were included in this study during 2018 - 2019. Patients with ID were selected from the Sleep Disorders Research Center, Kermanshah University of Medical Sciences, Iran. The study included HCs with good sleep quality and Pittsburgh Sleep Quality Index (PSQI) scores under five. Moreover, patients with Par-I were identified based on the mismatch between PSQI scores and nocturnal polysomnography (PSG) measurements. The reported mismatch involves a variance of at least 1 h difference for total sleep time (TST) or 15% for sleep efficiency (SE) between objective (PSG) and subjective (PSQI) measures besides ID patients with objective TST and SE above 6 h, 30 min, and 85%, respectively (6). The exclusion criteria were pregnancy, a history of other neurological and mental illnesses, and MRI imaging contraindications such as having metal in their bodies or claustrophobia. The Kermanshah University of Medical Science Ethics Committee approved the study, and all participants signed informed consent before participating. After excluding five patients with mild obstructive sleep apnea, two patients with comorbid periodic leg movements, two patients with brain tumors, one patient with hydrocephaly, two Par-I patients, and four HC due to excessive head motions (translation > 1.5 mm and rotation > 1.5 degrees), the analysis included 15 Par-I patients (11 male, mean ± SD age: 42.33 ± 12.23) and 48 HCs (24 male, age: 40.35 ± 12.67). The acquisition of images and the preprocessing of functional images were described in (14). Finally, ALFF maps were calculated for each Par-I and HC.

A chi-square test was conducted to assess potential gender differences between the two groups. The t-test was performed to calculate differences between age, PSQI, and head motion differences between the groups. Moreover, a t-test was used to check the possibility of ALFF differences between Par-I and HC. The effects of age and gender were controlled using a regressor, and multiple comparisons and the family-wise error (FWE) were used to prevent arbitrary results.

4. Results and Discussion

A total of 48 HC subjects (24 male, mean ± SD (age): 40.35 ± 12.67) and 15 Par-I patients (11 male, age: 42.33 ± 12.23) were included in the study. Table 1 presents the demographic characteristics of all subjects. Additionally, a t-test was used to examine possible differences between Par-I and HC groups, and no significant differences were found (P-value = 1.000). Moreover, the result of the chi-square test indicated no significant differences in gender between the two groups (P-value = 0.112). Furthermore, t-tests revealed significant differences between the two groups in PSQI (P = 0.001), but not in head motion (P = 1.000) (Table 1).

This study evaluated possible ALFF differences between patients with Par-I and HCs using the t-test. Multiple comparison corrections (FWE) revealed that ALFF decreased in superior parietal and precuneus regions in Par-I patients as compared to HC patients (Figure 1). The results were significant at the level of P_{FWE} < 0.001, and no significant differences were found in increased ALFF for Par-I compared to HC after multiple comparisons.

The purpose of this study was to assess whether ALFF differs between patients with Par-I and HC, based on resting-state fMRI images. The results indicated a significant ALFF decrease in the Par-I group compared to the healthy group. This decrease was evident in the superior parietal lobule and precuneus regions of the brain, which is in line with previous studies.

Koenigs et al. examined the role of parietal lobe regions in human working memory (15). Based on an extensive assessment of cognitive function and a sample of 19 patients with changes in the shape of the parietal lobe regions, they found that: (1) changes in the parietal lobe regions are related to deficits in information manipulation, and reorganization; (2) patients with parietal lobe changes have difficulty manipulating and reorganizing information in working memory for auditory-verbal and visual-spatial stimuli. Based on their findings, the parietal lobe regions are essential to manipulating and reorganizing information in working memory. In addition, previous studies have indicated the role and effects of sleep on memory areas (16). Furthermore, Li et al. investigated the functional relationship between brain regions among people with insomnia and HC and indicated changes in the functional connection between the parietal lobe and other brain parts compared to HC (17).

This network is assumed to be critical in regulating conscious processes, awareness, mental self-

| Table 1. Demographics and Clinical Variables of the Participants * |
|--------------------|---------|---------|--------|
| Variables | HC (48) | PDI (15) | PValue |
| Gender (F, M) | 24/24 | 4/11 | 0.112 |
| Age | 40.35 ± 12.67 | 42.33 ± 12.23 | 1.000 |
| Head motion | 0.091 ± 0.052 | 0.088 ± 0.052 | 1.000 |
| PSQI | 3.33 ± 1.79 | 16.47 ± 2.03 | < 0.001 |

Abbreviation: PSQI, Pittsburgh sleep quality index.
* Values are expressed as mean ± SD unless otherwise indicated.
representation, and inferential information processing. Moreover, Levenson et al. found changes in these areas in insomnia disorder (18), as well as a relationship between precuneus structural and variation in brain function with the quality of sleep among depressive patients (19). Finally, further studies should investigate the role of this network in Par-I by patients with equal total sleep time due to the uncovered role of the parietal lobe in organizing memory and relation between impaired memory and Par-I (3, 20).

This study should be interpreted with caution due to the low number of included subjects. The leak of this information prevented us from analyzing other covariates, such as anxiety and depression factors, in order to unravel the pathophysiology of the parietal lobe. Furthermore, a 1.5 T scanner was used, while nowadays, scanners with 3T have been used to increase the signal-to-noise ratio and precision.

4.1. Conclusions

According to our findings, brain signals play a differential role in paradoxical insomnia’s pathophysiology, and these findings may lead to further analysis of the underlying mechanism in the near future. Additionally, future studies should consider more complicated methods like machine learning with more subjects to study ID subtypes.

Acknowledgments

We would like to express our appreciation to the study’s participants and their families, and specially thank the Sleep Disorders Research Center’s staff for their assistance in data collection.

Footnotes

Authors’ Contribution: M. A. contributed to design, analyze, and conceptualization of the idea. H. K. collected data. M. A. and H. K. drafted the manuscript and revised the manuscript for intellectual content and collected data.

Conflict of Interests: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

Data Reproducibility: The dataset presented in the study is available on request from the corresponding author during submission or after its publication upon reasonable request.

Ethical Approval: This study is approved under the ethical approval code of IR.KUMS.REC.1398.971 (webpage of ethical approval code is: ethics.research.ac.ir/IR.KUMS.REC.1398.971).

Funding/Support: This study was supported in part by grant 97327 (Habibolah Khazaie).

Informed Consent: All participants signed informed consent before participating.

References


