Sleep Quality in Patients on Maintenance Hemodialysis and Peritoneal Dialysis

Maryam Masoumi, Afsoon Emami Naini, Rozita Aghaghazvini, Babak Amra¹, Ali Gholamrezaei²

ABSTRACT

Background: Sleep disturbances are common among uremic patients; however, limited data are available on predictors of sleep quality in this population. We assessed sleep quality in patients on hemodialysis (HD) and peritoneal dialysis (PD) and investigated predictors related to sleep quality.

Methods: Patients on maintenance HD and PD were consecutively included from two medical centers in Isfahan city (Iran). They completed the Pittsburgh sleep quality index (PSQI) and hospital anxiety and depression scale. Laboratory tests were done for iron state, kidney function, and electrolytes. Univariate and multivariate analyses were performed to find predictors of overall poor sleep quality.

Results: About 90 patients were evaluated (53 males, age = 54.2 ± 15.2 years, disease duration = 5.3 ± 4.5 years). Poor sleep quality was frequent in 86.6% of the cases in each group of HD and PD patients. Patients on HD had poorer sleep quality in terms of total PSQI scores and two dimensions of sleep latency and sleep efficiency (P<0.05). Anxiety (β = 0.232, P = 0.027), depression (β = 0.317, P = 0.004), and being on HD (β = 2.095, P = 0.009) were independent predictors of overall poor sleep quality.

Conclusions: Poor sleep quality is highly frequent in patients on maintenance dialysis and mood disorders and being on HD are predictive factors. Further studies are required for better understanding of risk factors associated with poor sleep quality and thus possible treatments in these patients.

Keywords: End-stage renal disease, hemodialysis, peritoneal dialysis, risk factors, sleep disorders

INTRODUCTION

Various types of sleep disturbances are common among patients with end-stage renal disease (ESRD), ranging from 50% to 80% by subjective manner and in up to 50% of the cases as objectively documented with polysomnography.¹,² Frequent sleep disorders in patients on hemodialysis (HD) and peritoneal dialysis (PD) include sleep apnea syndrome, restless legs syndrome, and periodic limb movement disorder.² The etiology of sleep disorders in patients on dialysis is not completely discovered; however, it is known to be multi-factorial. Various
conditions, including dialysis, medications, metabolic abnormalities, malnutrition, fatigue, muscle cramps, peripheral neuropathy, and emotional problems, all common in ESRD patients, can have effects on development of sleep disturbances in these patients.\cite{1,2,4,5}

The presence of sleep disorder is associated with lower quality of life and higher mortality in patients on dialysis.\cite{5‑7} Thus, it is important to investigate the risk factors associated with sleep disorders and improve the quality of sleep in ESRD patients in order to increase patients' quality of life and probably reduce associated mortality. Although separate studies have investigated the type and frequency of sleep disturbances in HD and PD patients, only few reports are available on the effects of dialysis type on sleep quality.\cite{8‑10} Furthermore, the results of previous studies on other risk factors have been controversial. Hence, we assessed sleep quality in patients on HD and PD and evaluated possible risk factors related to sleeping disturbances.

**METHODS**

**Patients and settings**

This cross-sectional study was conducted between January and December 2011 on adult patients with ESRD who were on maintenance HD or PD for at least 3 consecutive months in two academic medical centers in Isfahan city (Iran). Patients with a history of hospitalization within 4 weeks before the study were not enrolled. Consecutive sampling was done and with regard to the first type of error (\(\alpha\)) = 0.05, study power = 80%, and minimum expected difference of 10% in the frequency of poor sleep quality between the HD and PD patients based on previous studies,\cite{9} sample size was calculated as 45 cases in each group of the HD and PD patients. The study protocol was approved by the Ethics Committee of Isfahan University of Medical Sciences and an informed consent was obtained from all patients.

**Assessments**

Based on patients’ documents and during an interview with a trained resident of internal medicine, a questionnaire, including demographic, disease, and dialysis characteristics, sleep quality, and psychological factor (anxiety and depression) was completed with the following parts:

**Pittsburgh sleep quality index**

The Pittsburgh sleep quality index (PSQI) is an 18-item self-rated questionnaire for the evaluation of the overall sleep quality within the preceding month. PSQI evaluates seven components of sleep quality, including subjective sleep quality, sleep latency, sleep duration, habitual sleep efficiency, sleep disturbances, use of medication for sleep, and daytime dysfunction due to poor sleep. Each component score is calculated and coded from 0 to 3, where 0 indicates “no difficulty,” and 3, “severe difficulty.” The global score ranges from 0 to 21, where scores of \(\leq 5\) indicate good sleep quality and scores of >5 indicates poor sleep quality.\cite{11}

**The hospital anxiety and depression scale**

The hospital anxiety and depression scale (HADS) is a brief and widely used measure of psychological distress in outpatient setting. The HADS contains 14 items and consists of two sub scales of anxiety and depression. Each item is rated on a 4-point scale, giving a maximum score of 21 for each of the anxiety and depression sub scales. Scores of 11 or more on either sub scale are considered to be a significant “case” of psychological morbidity, while scores of 8-10 represents a “borderline” and 0-7 a “normal” case.\cite{12,13} For the above-mentioned questionnaires, a linguistically validated and reliable Persian version was used in this study.\cite{14,15}

**Laboratory assessments**

Patients were tested for complete blood counts, iron state with serum iron, ferritin, total iron binding capacity (TIBC), and transferrin (and saturation), albumin, electrolytes, including calcium and phosphorus, creatinine, blood urea nitrogen (BUN), and C-reactive protein (CRP). Fasting blood samples were taken in the morning in almost all patients when they came to the hospitals for dialysis session or for visits, and then were sent to one clinical laboratory for analyses. In those who came for dialysis session, blood samples were taken before the dialysis.

**Data analyses**

Data were analyzed using the SPSS software for windows (version 16.0, Chicago IL., USA). Comparisons between categorical variables were done using the Fisher's exact or Chi-square tests. Student's \(t\)-test and Mann–Whitney tests were used for comparison of parametric and non-parametric
data, respectively. Multivariate regression analyses were also performed to find factors associated with sleep quality. Statistical significance was defined at the 95% level ($P<0.05$). Because not all comparisons have a predefined 1-tailed hypothesis, we reported 2-tailed $P$ values for all analyses.

**RESULTS**

During the study period, 45 patients on HD and 45 patients on PD were evaluated including 53 males and 37 females with the mean age of 54.2 ± 15.2 years and disease duration of 5.3 ± 4.5 years. Demographic data, disease characteristics, and laboratory tests’ results of the patients are presented in Table 1. Compared with patients on PD, those who were on HD had longer disease duration ($P<0.001$), lower body mass index (BMI) ($P=0.013$), lower calcium ($P=0.003$) and higher phosphate and calcium × phosphate product ($P<0.001$), lower BUN ($P=0.044$) and creatinine ($P<0.001$), and higher albumin ($P<0.001$) and ferritin levels ($P=0.001$). Furthermore, patients on PD were under iron supplementation more frequently than HD patients ($P=0.035$). Other variables were the same between the HD and PD patients.

**Psychological state and sleep quality**

Comparisons of psychological state and sleep quality between the two groups of HD and PD patients are presented in Table 2. The two groups were not significantly different regarding anxiety or depression scores ($P>0.05$). Poor sleep quality was frequent in 86.6% of the cases in each group. However, patients on HD had poorer sleep quality in terms of total PSQI scores and two dimensions of sleep latency and sleep efficiency ($P<0.05$).

**Predictors of sleep quality**

In univariate analyses, there was a significant association between anxiety/depression scores and PSQI total score ($r=0.463$ and 0.478, respectively, $P<0.001$). Age, gender, ESRD duration, comorbid diseases, or BMI were not significantly associated with PSQI total score, but there was a non-significant relationship between physical activity and total PSQI score ($r=-0.198, P=0.063$). Physical activity was also significantly associated with sleep quality in terms of “day dysfunction due to sleepiness” ($r=-0.429, P<0.001$). Scores of this component of sleep quality was also associated with CRP levels ($r=0.249, P=0.019$). Other laboratories tests results were not associated with PSQI total score or its components.

In multivariate analysis, with a stepwise model, anxiety ($\beta=0.232, P=0.027$), depression ($\beta=0.317, P=0.004$), and being on HD ($\beta=0.2095, P=0.009$) were independent factors associated with overall poor sleep quality (total PSQI score). Predictors of sleep quality with regards to different PSQI domains

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**Table 1**: Demographic, clinical, and laboratory characteristics of patients on hemodialysis and peritoneal dialysis

<table>
<thead>
<tr>
<th>Variables</th>
<th>HD $n=45$</th>
<th>PD $n=45$</th>
<th>$P$</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (year)</td>
<td>54.5±15.0</td>
<td>54.0±15.6</td>
<td>0.875</td>
</tr>
<tr>
<td>Gender (M/F) (%)</td>
<td>(55.5)/20</td>
<td>(62.2)/17</td>
<td>0.334</td>
</tr>
<tr>
<td>(44.4)</td>
<td>(37.7)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>ESRD duration (year)</td>
<td>7.4±5.3</td>
<td>3.3±2.2</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>BMI (kg/m²)</td>
<td>22.7±3.9</td>
<td>25.0±4.5</td>
<td>0.013</td>
</tr>
<tr>
<td>ESRD etiology (%)</td>
<td>18 (40)</td>
<td>18 (40)</td>
<td>0.570</td>
</tr>
<tr>
<td>Diabetes</td>
<td>11 (24.4)</td>
<td>13 (28.8)</td>
<td></td>
</tr>
<tr>
<td>Hypertension</td>
<td>2 (4.4)</td>
<td>4 (8.8)</td>
<td></td>
</tr>
<tr>
<td>Glomerulonephritis</td>
<td>8 (17.7)</td>
<td>7 (15.5)</td>
<td></td>
</tr>
<tr>
<td>Others</td>
<td>6 (13.3)</td>
<td>3 (6.6)</td>
<td></td>
</tr>
<tr>
<td>Comorbid (%)</td>
<td>31 (68.8)</td>
<td>39 (86.6)</td>
<td>0.074</td>
</tr>
<tr>
<td>Hypertension</td>
<td>20 (44.4)</td>
<td>20 (44.4)</td>
<td>1.000</td>
</tr>
<tr>
<td>Diabetes</td>
<td>13 (28.8)</td>
<td>15 (33.3)</td>
<td>0.820</td>
</tr>
<tr>
<td>Ischemic heart disease</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Hemoglobin (g/dL)</td>
<td>11.4±1.8</td>
<td>12.0±1.6</td>
<td>0.130</td>
</tr>
<tr>
<td>Albumin</td>
<td>4.0±0.4</td>
<td>3.4±0.4</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Iron (µg/dL)</td>
<td>79.7±32.4</td>
<td>81.6±34.2</td>
<td>0.784</td>
</tr>
<tr>
<td>TIBC (µg/dL)</td>
<td>291.2±42.6</td>
<td>282.5±62.7</td>
<td>0.445</td>
</tr>
<tr>
<td>Ferritin (ng/dL)</td>
<td>496.4±317.6</td>
<td>285.6±266.2</td>
<td>0.001</td>
</tr>
<tr>
<td>Transferrin (mg/dL)</td>
<td>186.6±38.3</td>
<td>194.8±42.5</td>
<td>0.340</td>
</tr>
<tr>
<td>Transferrin saturation (%)</td>
<td>44.4±20.3</td>
<td>43.9±21.5</td>
<td>0.906</td>
</tr>
<tr>
<td>Calcium (mg/dL)</td>
<td>8.16±1.04</td>
<td>8.71±0.59</td>
<td>0.003</td>
</tr>
<tr>
<td>Phosphate (mg/dL)</td>
<td>5.48±1.35</td>
<td>4.04±0.85</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Ca×Ph (mg²/dL²)</td>
<td>44.6±11.6</td>
<td>35.1±7.1</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Erythropoietin</td>
<td>5288.8±</td>
<td>4466.6±</td>
<td>0.399</td>
</tr>
<tr>
<td>dose (U/Wk)</td>
<td>4920.3</td>
<td>4261.8</td>
<td></td>
</tr>
<tr>
<td>CRP (mg/L)</td>
<td>6.6±10.9</td>
<td>6.9±11.6</td>
<td>0.892</td>
</tr>
<tr>
<td>Iron supplements (%)</td>
<td>10 (22.2)</td>
<td>19 (42.2)</td>
<td>0.035</td>
</tr>
</tbody>
</table>

Data are presented as mean±SD and number (%). ESRD=End-stage renal disease, HD=Hemodialysis, PD=Peritoneal dialysis, TIBC=Total iron binding capacity, BMI=Body mass index, TIBC=total iron binding capacity, CRP=C-reactive protein.
DISCUSSION

Sleep disturbances are common among ESRD patients and are associated with decreased quality of life and higher mortality rates.\(^\text{[5–7]}\) Having knowledge about the risk factors of sleep disorders is helpful in designing preventive and therapeutic strategies for these patients. We evaluated the association of dialysis type (HD vs. PD) and other possible risk factors with sleep quality in patients on dialysis. The prevalence of poor sleep quality in our patients was similar to other reports from our society (87.8%).\(^\text{[16]}\) Using the same sleep measure (PSQI), the frequency of poor sleep quality is reported from 57% to 88.5% in other societies.\(^\text{[9,17–19]}\) This variation between different populations highlights the role of internal as well as external factors affecting the quality of sleep. Although previous studies evaluated different possible risk factors in patients on dialysis, only few of them included both HD and PD patients. Turkmen et al., found similar sleep quality scores between the HD and PD patients.\(^\text{[8]}\) Another study in the same population (Turkey) also found a non-significant difference between the HD and PD patients in the frequency of poor sleep quality (88.5% vs. 78.0%).\(^\text{[9]}\) The study by Holley et al., also did not found difference between HD and PD patients in the frequency or severity of sleep disturbances.\(^\text{[10]}\) In our study, the overall frequency of poor sleep quality was similar between the HD and PD patients, but the total PSQI score was higher in the HD patients. This difference was confirmed by the multivariate analysis, controlling for different factors, which showed independent association of being on HD with poorer sleep quality. On the other hand, patients on PD had higher BMI, more frequent hypertension, and lower albumin and ferritin levels, and these factors are supposed to be associated with poor sleep quality based on some reports.\(^\text{[9,17,18,20,21]}\) Regarding dialysis shift, Hsu et al., found a beneficial effect of evening HD, compared with morning and afternoon HD, on sleep quality and reduction of daytime symptoms,\(^\text{[22]}\) while another study did not find an association in this regard.\(^\text{[23]}\) Our sample was not large enough for such comparison as we only have one patient not receiving evening HD. Unfortunately, we did not have access to data on dialysis adequacy for all patients and precise analysis of such an association remains for future studies. Furthermore, in future studies other factors related to HD compared with PD must be considered as possible mechanisms by which patients on HD have poorer quality of sleep compared with patients on PD.

With regards to possible risk factors of poor sleep, we evaluated the association of demographic characteristics, clinical and laboratory parameters, and depression score with sleep quality in patients on dialysis. The most significant predictors of poorer sleep quality were anxiety, hemodialysis, sleep efficiency, depression, and female gender. This is consistent with other studies.\(^\text{[2,10]}\) Anxiety, depression, and female gender are non-modifiable factors associated with poor sleep quality. On the other hand, sleep efficiency, which is modifiable, was associated with sleep quality. The hemodialysis treatment was another important predictor of poor sleep. The sleep efficiency of patients receiving evening HD was significantly better than those receiving morning/fixed HD. This finding is consistent with other reports.\(^\text{[6,24]}\) Although not statistically significant, the sleep efficiency was higher in patients receiving PD compared with HD. These findings are consistent with previous reports.\(^\text{[10,22]}\) Having knowledge about the risk factors of sleep disorders is helpful in designing preventive and therapeutic strategies for these patients. We evaluated the association of demographic characteristics, clinical and laboratory parameters, and depression score with sleep quality in patients on dialysis. The most significant predictors of poorer sleep quality were anxiety, hemodialysis, sleep efficiency, depression, and female gender. This is consistent with other studies.\(^\text{[2,10]}\) Anxiety, depression, and female gender are non-modifiable factors associated with poor sleep quality. On the other hand, sleep efficiency, which is modifiable, was associated with sleep quality. The hemodialysis treatment was another important predictor of poor sleep. The sleep efficiency of patients receiving evening HD was significantly better than those receiving morning/fixed HD. This finding is consistent with other reports.\(^\text{[6,24]}\) Although not statistically significant, the sleep efficiency was higher in patients receiving PD compared with HD. These findings are consistent with previous reports.\(^\text{[10,22]}\) Having knowledge about the risk factors of sleep disorders is helpful in designing preventive and therapeutic strategies for these patients.
sleep quality other than type of dialysis, similar to several other studies,[24‑26] we found that mood disorders (anxiety and depression) are important factors independently associated with sleep quality. Mood disorders are highly prevalent in patients on dialysis, describing high prevalence of sleep disturbance in these patients.[27] Although mood disorders have direct effects on sleep disorders,[28] this association in ESRD patients may also be mediated by increased (neuro) inflammatory burden due to mood disorders resulting in other comorbidities such as cardiovascular and pulmonary disorders.[29] An association between age and sleep quality in ESRD patients is also reported by some studies.[18,25,30] In our study, however, age was independently associated with only one of the sleep quality domains, “day dysfunction due to sleepiness.” Although age is found to be independently associated with sleep quality, this association may be, at least in part, mediated by the presence of comorbid diseases, less physical activity, and nutritional status in older patients.[18,31‑33] We also found female gender an independent predictor of sleep latency. Female gender was associated with poor sleep quality in one other study,[30] but the other one found an association between male gender and sleep quality.[34] While male gender is found to be associated with sleep apnea syndrome,[35,36] female gender is an independent risk factor for restless legs syndrome.[37,38] and both of these syndrome are common among dialysis patients and are associated with poor sleep quality.[19,38,39] Thus, the association between gender and sleep quality may be mediated by the presence of other sleep disorders.

With regards to laboratory tests’ results, we did not find an association between calcium/phosphate and sleep quality, while in one study, poor sleep quality was associated with higher serum phosphate levels,[34] and other studies found that higher amount of calcium × phosphate product is associated with sleep quality.[18] This association may be justified by higher frequency of comorbidities in patients with elevated calcium × phosphate product, though further investigations are required in this regard.[18] Similar to several other reports we found an independent association between CRP levels and sleep quality.[40‑42] This association corroborated to the role of inflammation in the pathogenesis of sleep disturbances. On the other hand, the inflammatory state in dialysis patients might be due to sleep related breathing disorders.[43] Indeed, it might be a bidirectional relationship between sleep disturbance and inflammatory state in ESRD patients, which can also justify the association of sleep disorders with an increased mortality rate of dialysis patients.

Anemia is another possible mechanism that has long been considered as a risk factor for sleep disturbance in dialysis patients. An association between hemoglobin levels,[26] iron deficiency,[16] and transferrin saturation[17] has been indicated by previous reports. In our study, higher transferrin level (that may indicate an iron deficient state) was a predictor of poor overall sleep quality and sleep disturbance. However, other indicators of iron state were not associated with sleep quality in our study and a precise conclusion is not possible from our data in this regard. Anyway, the full evaluation of iron status in ESRD patients with sleep disorder and appropriate therapy when results indicate an iron deficiency state seems to be warranted.

While several studies showed high prevalence of sleep disorders among patients on dialysis, few reports are available on treatment strategies, which could be attributed to the unclear pathophysiology of sleep disorders in these patients. Also, the presence of frequent comorbidities and drug interactions limit the use of sleep medications in ESRD patients. However, considering the current known risk factors, some trials have been done with two main approaches; therapeutic interventions and dialysis modifications. In one study, a course of cognitive-behavioral therapy resulted in improvement of sleep quality and psychological state.[44] Another study with intradialytic aerobic exercise found the same results.[45] In these studies, sleep quality improvement has been associated with decrease in inflammation and oxidative stress markers, highlighting the role of inflammation and oxidative stress in sleep disturbances in patients on dialysis.[45,46] The beneficial effects of cognitive-behavioral therapy and (intradialytic aerobic) exercise have also been reported by other studies.[47,48] Thus, regarding high prevalence of mood disorders in ESRD patients on dialysis and its effects on sleep quality, psycho-behavioral therapies could be good candidates for the treatment of mood/sleep disorders in ESRD patients. Regarding dialysis-related factors, skin temperature changes following dialysis have possible effects on nocturnal
sleep, and using cool dialysate during HD showed improved sleep during the night following treatment in one trial.\[49\] The underlying mechanism in this trial might be decreasing sympathetic activation and maintaining normal nocturnal skin temperature.\[49\] Several studies also reported beneficial effects of nocturnal dialysis, which provides superior dialysis based upon dose, duration, and frequency.\[50‑53\] Correction of melatonin rhythm\[51,54\] and better fluid and uremic clearance are mentioned as possible mechanisms in this regard.\[52\] Anyway, much more studies are required for the treatment of sleep disorders in ESRD patients.

**CONCLUSIONS**

The results of the present study showed that poor sleep quality is highly frequent in uremic patients on maintenance dialysis, and mood disorders and being on HD are major risk factors for poor sleep quality in these patients. Furthermore, we found that age, female gender, lower physical activity, inflammation, and an iron deficient state are independent predictors of quality of sleep in some of its dimensions. Further studies are required for better understanding of risk factors associated with poor sleep quality and thus possible treatments in these patients. With current knowledge, psycho-behavioral therapies could be good candidates for the treatment of mood/sleep disorders in ESRD patients on maintenance dialysis.

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