Restless legs syndrome (RLS) is a movement disorder characterized by an imperative urge to move the legs associated with uncomfortable sensations, paraesthesias and motor restlessness. These symptoms worsen during rest, mainly at night, with at least temporary relief brought on by activity. It has a circadian pattern and af-
Restless legs syndrome in Parkinson’s disease
Guerrero et al.
Arq Neuropsiquiatr 2010;68(6)

Effects onset and continuity of sleep due to the unpleasant sensations at bedtime. The prevalence rates of RLS vary with age, gender, causes (primary or secondary) and in different ethnic populations. In general population its prevalence range between 0.8% and 15%. Most people who are affected are middle-aged or older women.

The pathophysiology of RLS remains unclear and it is either genetic or secondary to other medical conditions, such as neuropathy, iron deficiency, renal failure, pregnancy and, possibly, rheumatoid arthritis. In most cases, serum ferritin and iron-binding saturation should be obtained for serum iron deficiency, and electrolytes should be obtained for renal failure. Familial cases of RLS demonstrating autosomal dominant pattern of inheritance are well-known described. Recent clinical and functional imaging data suggest impaired central dopaminergic transmission. Thus, dopaminergic drugs have been widely accepted as an efficacy therapy for RLS, and dopamine medications currently represents the first-line treatment for this disorder.

RLS was first described by Ekbom in 1945 and it has remained as a defined clinical diagnosis since then. Diagnostic criteria was established by the International Restless Legs Syndrome Study Group (IRLSSG). The aim of the present study was to investigate the prevalence, clinical characteristics and biochemical correlations of restless legs syndrome amongst Parkinson’s disease (PD) patients.

METHOD
This study was carried out in a Movement Disorder Outpatient Clinic in the Federal University of Sergipe from January 2007 to December 2007. Forty-eight (N=48) patients with PD were recruited. The ethic committee of the Federal University of Sergipe has approved this study, and all subjects gave their informed consent to take part in the investigation.

The diagnosis of PD was made previously by a specialist in movement disorders using the UKPD Society Brain Bank criteria. PD patients with diabetes mellitus, renal failure, anemia and rheumatoid arthritis were excluded from the study. The diagnosis of RLS was established by face-to-face interviews based upon the criteria of the International Restless Legs Syndrome Study Group. The criteria comprises a set of four questions: [1] desire to move the limbs usually associated with paresthesias/dysesthesias; [2] motor restlessness; [3] symptoms which are worse or exclusively present at rest (i.e., lying and sitting) with at least partial and temporary relief by activity; and [4] symptoms are worse in the evening or night. The diagnosis is made when all the above four criteria were met.

After making the diagnosis of RLS, PD patients were divided into two groups: those with RLS and those without RLS. Clinical characteristics assessed in both groups were age, gender, duration of PD, severity (Hoehn and Yahr scale) and activities of daily living scale (Schwab and England scale) of PD. Biochemical parameters such as hemoglobin, s-iron, s-ferritin and creatinine were obtained.

Following, comparisons of clinical and biochemical variables between each group were made. SPSS (v. 15.0) statistical software was used and statistical analyses by Mann-Whitney test, Chi-square, Fisher exact test and Student t test.

RESULTS
A total of 48 patients were recruited. The estimated prevalence of RLS found among PD patients in our Movement Disorder Clinic was 18.75% (9 patients).

Table 1. Comparison of clinical and biochemical findings between Parkinsonian patients with and without restless legs syndrome (Outpatient in University Hospital/ Federal University of Sergipe - January 2007 to December 2007).

<table>
<thead>
<tr>
<th></th>
<th>Positive RLS group (n=9)</th>
<th>Negative RLS group (n=39)</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>67.11±9.47</td>
<td>63.92±11.46</td>
<td>0.443</td>
</tr>
<tr>
<td>Masculine gender (%)</td>
<td>77.80</td>
<td>87.20</td>
<td>0.471</td>
</tr>
<tr>
<td>PD duration (years)</td>
<td>9.55±5.45</td>
<td>7.35±3.59</td>
<td>0.277</td>
</tr>
<tr>
<td>Hoehn and Yahr</td>
<td>2.83±0.93</td>
<td>2.51±1.19</td>
<td>0.309</td>
</tr>
<tr>
<td>Schwab and England</td>
<td>65.55±27.43</td>
<td>71.02±26.33</td>
<td>0.640</td>
</tr>
<tr>
<td>Hemoglobin (g/dl)</td>
<td>13.41±1.71</td>
<td>13.55±1.28</td>
<td>0.829</td>
</tr>
<tr>
<td>Serum iron (mcg/dl)</td>
<td>78.50±35.05</td>
<td>90.64±33.88</td>
<td>0.306</td>
</tr>
<tr>
<td>Serum ferritin (ng/ml)</td>
<td>145.87±72.09</td>
<td>157.70±138.53</td>
<td>0.530</td>
</tr>
<tr>
<td>Urea (mg/dl)</td>
<td>37.50±9.29</td>
<td>34.84±14.88</td>
<td>0.083</td>
</tr>
<tr>
<td>Creatinine (mg/dl)</td>
<td>1.04±0.26</td>
<td>1.04±0.27</td>
<td>0.971</td>
</tr>
</tbody>
</table>

Positive RLS group: group of parkinsonian patients with restless legs syndrome; Negative RLS group: group of parkinsonian patients without restless legs syndrome; PD: Parkinson’s disease; Normal range of hemoglobin: 13-17 g/dl; Normal range of serum iron: 35-150 mcg/dl; Normal range of serum ferritin: 16.4-253 ng/ml; Normal range of urea: 15-39 mg/dl; Normal range of creatinine: 0.6-1.3 mg/dl.
Table 2. Comparison of Hoehn and Yahr grades between Parkinsonian patients with and without restless legs syndrome (Outpatient in University Hospital/ Federal University of Sergipe - January 2007 to December 2007).

<table>
<thead>
<tr>
<th>Hoehn e Yahr scale</th>
<th>Positive RLS group (%)</th>
<th>Negative RLS group (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>≤ 2.5</td>
<td>3 (33%)</td>
<td>25 (64%)</td>
</tr>
<tr>
<td>&gt;2.5</td>
<td>6 (67%)</td>
<td>14 (36%)</td>
</tr>
</tbody>
</table>

Positive RLS group: group of parkinsonian patients with restless legs syndrome. Negative RLS group: group of parkinsonian patients without restless legs syndrome.

On those diagnosed with RLS (positive RLS group), the mean age was 67.11±9.47 years and the mean duration of PD was 9.55±5.45 years. 77.80% were men and 22.20% were women (Table 1).

The mean Hoehn and Yahr (UPDRS part V) score found was 2.83±0.93 and in the Schwab and England scale (UPDRS part VI) was 65.55±27.43% (Table 1). Only three (N=3) patients (33%) showed Hoehn and Yahr (UPDRS part V) grades lower than 2.5 (Table 2).

Data obtained by biochemical variables, such as hemoglobin, s-ferritin, s-iron and creatinine are registered on Table 1. Only one (N=1) patient showed s-ferritin value lower than 45 ng/ml on positive RLS group.

On the other group of PD patients, those without RLS (negative RLS group), the mean age found was 63.92±11.46 years and the mean duration of PD was 7.35±3.59 years. 87.20% were men and 12.80%, women (Table 1).

In the negative RLS group, the mean Hoehn and Yahr score found was 2.51±1.19 and in the Schwab and England scale was 71.02±26.33% (Table 1). Up to 64% patients showed Hoehn and Yahr scores lower than 2.5 (Table 2).

Biochemical parameters assessed on negative RLS group are reported in Table 1.

No significant differences between the two groups regarding age, gender, duration of PD and severity of PD (Hoehn and Yahr scale), activities of daily living scale (Schwab and England scale) were observed. Biochemical parameters did not differ between on those PD patients with RLS and on those without RLS (Table 1).

**DISCUSSION**

The prevalence rate of RLS in PD patients appears to be higher than in general population. It may vary from one survey to another since earlier studies have not used the full set of four diagnostic criteria of the IRLSSG. Any series not using the full criteria simply were not considered valid. This study was performed following that questionnaire during all face-to-face interviews.

A RLS prevalence of 18.75% was estimated among this series of PD patients. Of 48 PD patients, 9 fulfilled the diagnosis criteria. Ondo et al. and Gomez et al. had found prevalence rates of 20.8% and 21.9%, respectively. Although, Nomura et al. and Krishnan et al. were not able to replicate these findings in their series of PD patients estimating prevalence rates of 12% and 7.9%, respectively. The higher prevalence noted among our group of PD patients may corroborate for the hypotheses of similar pathology between RLS and PD.

PD patients with RLS were older than those without RLS but there was no statistical difference between RLS and non-RLS groups (positive RLS: 67.11±9.47 years; negative RLS: 63.92±11.46 years; p=0.443). This finding is consistent with that reported by Gómez et al., who found no statistical difference (positive RLS: 69.8±8.8 years; negative RLS: 68.4±9.7 years) among their series of 114 patients with PD, and also with Calzetti et al. (positive RLS: 71.0±4.5 years; negative RLS: 68.0±8.0). In an Indian survey, they reported a significant difference of mean age between PD patients with and without RLS (positive RLS: 63.70±7.8 years; negative RLS: 57.37±10.04 years; p=0.05). Nomura and associates, however, concluded that PD patients without RLS seemed to be younger than those with RLS (positive RLS: 59.7±2.7 years; negative RLS: 70.1±0.8 years; p=0.0001).

The subjects included in the positive RLS group were mostly men (seven of nine patients). Krishnan et al. e Tan et al. have found similar gender distribution. This result is not noted by others and RLS has been found to be more prevalent in female. Studies have reported prevalence two or three times higher among female subjects with RLS. Gómez et al. found 25 (21.9%) patients with RLS among their PD patients and on this group, 17 (68%) were female and 8 (32%) male.

In the present study, most of the RLS patients (67%) were in an advanced stage of PD (Hoehn and Yahr scale score ≥ 2.5). The mean score in the positive RLS group was 2.83±0.93 and the mean score in the negative RLS was 2.51±1.19. Although, there was no significant difference between those two groups. Similar findings have been reported by others as Krishnan et al. and Ondo et al.

Previous studies raised the possibility that low level of serum ferritin may be related with the occurrence of RLS in PD patients since iron has a role in both biosynthesis and transmission of monoamines. It is known that low iron level leads to dopaminergic dysfunction. Its deficiency decreases density of dopamine D2 and D1 receptors in the caudate-putamen. In the series of 18 patients with RLS and 18 controls, O’Keeffe et al. found serum ferritin levels significantly lower in the patients with RLS (positive RLS: 33 ng/ml; negative RLS: 59 ng/ml; p<0.01). Although, in this study no statistical difference of serum ferritin level was found between positive RLS group and negative RLS group. Ondo et al. and Nomura et al. have reported similar findings.
Only one patient with RLS had serum ferritin levels <45 ng/ml. Winkelmann et al. recommend iron replacement if ferritin levels are <50ng/ml, even though this level is considered to be in normal range. The same authors have reported that lower ferritin levels are also associated with greater RLS severity and decreased sleep efficiency, with increasing waking time after sleep onset. O’Keeffe et al. treated patients who had serum ferritin levels <45 ng/ml with iron replacement obtaining partial relief of RLS symptoms.

This study suffers from certain limitations collecting biochemical data since most of the sample of PD patients comprises subjects mostly from the suburbs of Aracaju, the countryside as well as nearby states, Alagoas and Bahia. This fact is responsible for the reduced number of patients.

All patients were interviewed and examined personally by the author in order to avoid bias during the development of our study with false-positive or false-negative findings.

In conclusion, the present study suggests that RLS occurs more frequently in patients with PD. Moreover, there was no statistical difference comparing clinical and biochemical characteristics between subjects with RLS and without RLS. It seems that secondary causes of RLS did not play a central role in the pathophysiology of RLS in this group of parkinsonian patients.

We reiterate the importance of diagnosis of RLS in patients with PD, since disturbances in sleep is significantly more frequent in parkinsonian patients than in healthy population and it may worsen their quality of life. Studies with larger sample are necessary to explore the relationship of RLS in brazilian patients with PD.

ACKNOWLEDGMENT – The authors thank all patients for participating in the study.

REFERENCES