

*National Center on Sleep Disorders Research
and Office of Prevention, Education, and Control*

RESTLESS LEGS SYNDROME

DETECTION
AND MANAGEMENT
IN PRIMARY CARE

NATIONAL INSTITUTES OF HEALTH
NATIONAL HEART, LUNG, AND BLOOD INSTITUTE



RESTLESS LEGS SYNDROME

**DETECTION
AND MANAGEMENT
IN PRIMARY CARE**

*NATIONAL CENTER ON
SLEEP DISORDERS RESEARCH
NATIONAL HEART, LUNG,
AND BLOOD INSTITUTE
NATIONAL INSTITUTES
OF HEALTH*

*Produced in collaboration
with the Restless Legs Syndrome
Foundation through an unre-
stricted educational grant from
Pharmacia & Upjohn and
Boehringer Ingelheim.*

MARCH 2000

NIH PUBLICATION No. 00-3788

MEMBERS OF THE NATIONAL HEART, LUNG, AND BLOOD INSTITUTE WORKING GROUP ON RESTLESS LEGS SYNDROME

Michael Thorpy, M.D. (Chair)
Director
Sleep Disorders Center
Montefiore Medical Center
Bronx, New York

Bruce L. Ehrenberg, M.D.
Director
Clinical, EEG, and Sleep Laboratory
Tufts–New England Medical Center
Boston, Massachusetts

Wayne A. Hening, M.D., Ph.D.
Assistant Clinical Professor of Neurology
UMDNJ–Robert Wood Johnson
Medical School
New York, New York

Mark Mahowald, M.D.
Director
Minnesota Regional Sleep Disorders Center
Hennepin County Medical Center
Minneapolis, Minnesota

Beth A. Malow, M.D., M.S.
Assistant Professor
Department of Neurology
University of Michigan
Ann Arbor, Michigan

Barbara Phillips, M.D., M.S.P.H.
Director
Sleep Apnea Center
Samaritan Hospital
Lexington, Kentucky

Caroline Richardson, M.D.
Robert Wood Johnson Scholars Fellow
University of Michigan Medical School
Ann Arbor, Michigan

Caroline Wellbery, M.D.
Assistant Professor, Family Medicine
Fort Lincoln Family Medical Center
Georgetown University Medical Center
Washington, DC

NATIONAL INSTITUTES OF HEALTH STAFF

Mark Hallett, M.D.
Clinical Director
National Institute of Neurological
Disorders and Stroke
National Institutes of Health
Bethesda, Maryland

James P. Kiley, Ph.D.
Director
National Center on
Sleep Disorders Research
National Heart, Lung, and Blood Institute
Bethesda, Maryland

Charlotte McCutchen, M.D.
Program Director
Neuroscience Center
National Institute of Neurological
Disorders and Stroke
Bethesda, Maryland

Susan Rogus, R.N., M.S.
Coordinator, Sleep Education Activities
Office of Prevention, Education, and Control
National Heart, Lung, and Blood Institute
Bethesda, Maryland

SUPPORT STAFF

Susan Shero, R.N., M.S.
Prospect Associates
Silver Spring, Maryland

CONTENTS

- Introduction 7
- Consequences of RLS 7
- Prevalence 7
- Etiology
 - Primary RLS 7
 - Secondary Causes of RLS 7
 - Iron Deficiency 7
 - Neurologic Lesions 7
 - Pregnancy 7
 - Uremia 8
 - Drug-Induced 8
- Assessment and Diagnosis 8
 - Clinical Criteria 8
 - Physical Examination 9
 - Laboratory Tests 9
 - Differential Diagnosis 9
- Treatment 11
- When To Consider Referral 12
- Conclusion 12
- References 13
- Appendix: Sleep/Wake Profile 15
- Where To Get More Information 16

RESTLESS LEGS SYNDROME

DETECTION AND MANAGEMENT IN PRIMARY CARE

INTRODUCTION

Restless legs syndrome (RLS) is a common, underdiagnosed, and treatable condition. A neurologic movement disorder, RLS is often associated with a sleep complaint.¹ Patients with RLS can suffer an almost irresistible urge to move the legs, usually due to disagreeable leg sensations that are worse during inactivity and often interfere with sleep.² RLS can be described as an agitated inability to rest that can have a negative impact on the quality of life due to waking discomfort, chronic sleep deprivation, and stress. This publication provides science-based information about RLS and its assessment and management in the primary care setting.

CONSEQUENCES OF RLS

Direct results of RLS include discomfort, sleep disturbances, and fatigue.³ These consequences have a secondary impact on functioning by affecting occupational activities, social activities, and family life. Disrupted sleep and an inability to tolerate sedentary activities can lead to job loss, a compromised ability to enjoy life, and problems with relationships.

PREVALENCE

RLS is a common disorder. Although the exact prevalence is uncertain, limited studies have indicated that 2 to 15 percent of the population may experience RLS symptoms.^{4,5,6} This wide range may be due to differences in study methodologies.

Although the prevalence of RLS increases with age⁶, it has a variable age of onset and can occur in children.⁷ In patients with severe RLS, one-third

to two-fifths had their first symptom before age 20⁸, although the precise diagnosis of RLS was made much later.

ETIOLOGY

Primary RLS

RLS is a central nervous system (CNS) disorder.⁹ It is not caused by psychiatric factors or by stress but may contribute to or be exacerbated by these conditions. There is a high incidence of familial cases of RLS, suggesting a genetic origin for primary RLS.⁸ The exact mode of inheritance is unknown.^{8,10}

Secondary Causes of RLS

- **Iron deficiency.** RLS may be associated with iron deficiency. A patient's iron stores may be deficient without significant anemia. Recent studies have shown that decreased iron stores (indicated by ferritin levels below 50 mcg/L) can exacerbate RLS symptoms.^{11,12} Patients with newly diagnosed RLS or RLS patients with a recent exacerbation of symptoms should have their serum ferritin levels measured.
- **Neurologic lesions.** RLS has been reported in association with spinal cord and peripheral nerve lesions, although an exact pathological mechanism has not been identified. RLS may also emerge in patients with vertebral disc disease.⁸
- **Pregnancy.** RLS affects up to 19 percent of women during pregnancy.¹³ Symptoms can be severe but usually subside within a few weeks postpartum.

- **Uremia.** RLS occurs in up to 50 percent of patients with end-stage renal failure, and may be particularly bothersome during dialysis when the patient is confined to a resting position.^{14,15} Improvement in RLS symptoms has been seen after renal transplantation.¹⁶
- **Drug-induced.** There is some evidence from published case reports that RLS symptoms may be worsened or unmasked by medications such as tricyclic antidepressants,¹⁷ selective serotonin reuptake inhibitors (SSRIs),¹⁸ lithium,¹⁹ and dopamine antagonists.²⁰ Caffeine also has been implicated in the worsening of RLS symptoms.²¹

ASSESSMENT AND DIAGNOSIS

The diagnosis of RLS is based primarily on the patient’s history. Often, patients do not bring RLS symptoms to the physician’s attention; therefore, it can be helpful to include general sleep questions in the review of systems. See Appendix. When

RLS is suspected, more specific questions can be asked. See Table 1.

Symptoms are described by patients in many ways, ranging from mild to intolerable.²² See Table 2. Although most patients experience the sensations in their legs, the sensations may also occur in the arms or elsewhere. RLS symptoms are generally worse in the evening and night and less severe in the morning. RLS needs to be distinguished from sleep-related leg conditions such as nocturnal leg cramps.

Clinical Criteria

The following criteria for diagnosis of RLS are based on those developed by the International Restless Legs Syndrome Study Group.³

Minimal criteria include the following:

1. A compelling urge to move the limbs, usually associated with paresthesias/dysesthesias.

Table 1

RLS-RELATED QUESTIONS
■ Does the patient report “creeping, crawling, or uncomfortable, difficult-to-describe feelings” in the legs or arms that are relieved by moving or rubbing them?
■ Is there a correlation between RLS symptoms and time of day? Do the symptoms worsen with rest or inactivity?
■ Do sensations interfere with sleep onset or returning to sleep?
■ What daytime consequences does the patient report (e.g., fatigue, sleepiness, confusion, lack of attention)?
■ Does the bed partner report that the patient’s legs or arms jerk during sleep? (Relates to periodic limb movements in sleep.)
■ Does the patient have secondary causes of RLS such as low iron stores, diabetes mellitus, kidney disease, or pregnancy?
■ Are neurological symptoms or diagnoses present?
■ Is there a relationship between symptoms and medications, such as tricyclic antidepressants or SSRIs?
■ Was the onset of symptoms correlated with a change in medication?
■ Do family members report similar symptoms? Have family members been diagnosed with RLS?

Table 2

TERMS USED TO DESCRIBE RLS SENSATIONS		
Creeping	Indescribable	Electric current-like
Crawling	Pulling	Restless
Itching	Drawing	Painful
Burning	Aching	
Searing	Like water flowing	
Tugging	Like worms or bugs crawling under the skin	

2. Motor restlessness as seen in activities such as floor pacing, tossing and turning in bed, and rubbing the legs.
3. Symptoms worse or exclusively present at rest (i.e., lying, sitting) with variable and temporary relief by activity.
4. Symptoms worse in the evening and at night.

Other associated features commonly found in RLS but not required for diagnosis include the following:

- Sleep disturbance and daytime fatigue.
- Normal neurological exam in primary RLS.
- Involuntary, repetitive, periodic, jerking limb movements, either in sleep or while awake and at rest.

The last feature refers to periodic limb movements (PLM), also known as PLMS (periodic limb movements of sleep)²³ or nocturnal myoclonus,²⁴ which may be associated with RLS. PLM are stereotyped, repetitive flexions of the limbs (legs alone or legs more than arms) usually occurring during sleep. They occur periodically on an average of every 20 seconds. The most common movement is a dorsiflexion of the ankles and flexion of the knees or hips.

Physical Examination

The physical examination is usually normal in patients with RLS and is performed to identify secondary causes and rule out other disorders. The following are areas of particular importance:

- *A neurological exam* with emphasis on spinal cord and peripheral nerve function.
- *A vascular exam* to rule out vascular disorders.

Laboratory Tests

The following laboratory tests can identify possible secondary causes of RLS:

- Serum ferritin level (<50mcg/L)
- Serum chemistry to rule out uremia and diabetes

A sleep study (polysomography) is **not** routinely indicated in the workup of RLS²⁵ because RLS is diagnosed on the basis of history and clinical findings.

Differential Diagnosis

Differential diagnoses may include the following:

- Nocturnal leg cramps (typically painful, palpable, involuntary muscle contractions, often focal, with a sudden onset; usually unilateral).²⁶

Table 3

PHARMACOLOGIC TREATMENT FOR RLS		
Agent	Advantages	Disadvantages
Dopaminergic Agents Dopamine precursor combinations such as carbidopa-levodopa	Can be used on a “one time” basis or as circumstances may require. Useful for persons with intermittent RLS because dopamine agonists take longer to have an effect.	As many as 80 percent of patients on carbidopa-levodopa may develop augmentation.* Therapeutic effect may be reduced if taken with high-protein food. Can cause insomnia, sleepiness, and gastrointestinal problems.
Dopamine agonists such as Pergolide Pramipexole Ropinirole	Useful in moderate to severe RLS. Recent reports indicate high efficacy of dopamine agonists, but the role of their long-term use is unknown. ²⁹	Can cause severe sleepiness, ³⁰ which may limit its use during daytime. Agonists can cause nausea. To avoid this, slow dose increase is important, especially for pergolide.
Opioids such as Codeine Hydrocodone Oxycodone Propoxyphene Tramadol	Can be used on an intermittent basis. Can also be used successfully for daily therapy.	Can cause constipation, urinary retention, sleepiness, or cognitive changes. Tolerance and dependence possible with higher doses of stronger agents.
Benzodiazepines such as Clonazepam Temazepam	Helpful in some patients when other medications are not tolerated, and may help improve sleep.	Can cause daytime sleepiness and cognitive impairment, particularly in the elderly.
Anticonvulsants such as Carbamazepine Gabapentin	Can be considered when dopamine agonists have failed. May be useful in those with coexisting peripheral neuropathy and/or when RLS discomfort is described as pain.	Disadvantages vary depending on agent, but include gastrointestinal disturbance such as nausea, sedation, dizziness.
Iron (ferrous sulfate)	Use in patients with serum ferritin levels <50 mcg.	Ideal means of administration has not been established. Oral treatment may take several months to be effective and may be poorly tolerated.
Clonidine	May be useful in hypertensive patients.	Has the potential to cause hypotension, dermatitis, and sleepiness.

*Augmentation is a worsening of RLS symptoms in the course of therapy. Symptoms may be more severe and start earlier in the day (e.g. afternoon rather than evening) than before treatment began and may spread to different parts of the body. Augmentation, which can start soon after therapy is begun or not until months or years later, has also been reported with dopamine agonists and may occur with other medications.

- Akathisia (excessive movement, without specific sensory complaints; often does not correlate with rest or time of day and usually results from medication such as neuroleptics or other dopamine blocking agents).²⁷
- Peripheral neuropathy (can cause leg symptoms that are different from RLS; they are usually not associated with motor restlessness, nor helped by movement, and have no evening or night time worsening. Sensory complaints are typically numbness, tingling, or pain. Small fiber sensory neuropathies, as seen in diabetes, are often confused with RLS. Patients with neuropathies may have both neuropathic and RLS symptoms.)
- Vascular disease (such as deep vein thrombosis).
- Severity of symptoms. (Some patients with mild symptoms may elect not to use medications; others may benefit from levodopa or a dopamine agonist. Patients with severe symptoms may require a strong opioid.)
- Frequency or regularity of symptoms. (Patients with infrequent symptoms may benefit from a single effective p.r.n. dose of a medication such as an opioid or levodopa.)
- Presence of pregnancy or comorbid illnesses. (There are no controlled clinical trials that have assessed the safety and efficacy of medications for RLS or PLM during pregnancy.²⁸)
- Renal failure. (In these patients, pharmacologic agents are generally safe, but less frequent doses may be needed if drugs are renally excreted. In addition, for dialysis patients, some medications are dialyzable [e.g., gabapentin] and others are not [e.g., propoxyphene].²⁸)

TREATMENT

The severity of RLS varies from patient to patient. Although pharmacologic treatment is helpful for many RLS patients, those with mild symptoms may not need medications. Since no single medication or combination of medications will work predictably for all patients, treatment must often be individualized. Physicians and patients may need to work together over time to find the medication or combination of medications and the dosages that will work best. See Table 3 for a list of pharmacologic agents and their advantages and disadvantages. Therapy for RLS constitutes an “off label” use of these pharmacologic agents.

The selection of pharmacologic agents is influenced by a number of factors, including:

- Age of the patient. (For example, benzodiazepines may cause cognitive impairment in the elderly.)

Dopaminergic agents are the first-line drugs for most RLS patients. It is important for the primary care physician to educate the patient about the nature and actions of the drugs that are prescribed, including side effects and the uncertainty of long-term effects. For example, when prescribed dopaminergic agents, the patient should be informed that although these medications are usually used to treat Parkinson’s disease, they also help to relieve RLS symptoms.

RLS medications have received approval from the Food and Drug Administration for other uses. In many cases, the therapeutic doses to treat RLS are much lower than those required for the original uses. The starting dose is usually very low and is gradually increased until effective. In addition to the medications listed in Table 3, agents such as vitamin E, folate, and magnesium may be useful. Although many nonpharmacologic treatments have been reported by patients to be helpful, there is no scientific evidence that they are useful in the treatment of RLS.

WHEN TO CONSIDER REFERRAL

For most patients, RLS can be effectively managed by the primary care physician. If the primary care physician encounters difficulty managing RLS symptoms in a patient, referral to, or consultation with, a movement disorders specialist or a sleep specialist may be helpful.

CONCLUSION

The primary care physician plays a central role in the identification and treatment of RLS. Incorporating sleep- and RLS-related questions into the general review of systems can be helpful in diagnosing RLS. An important aspect of treatment is listening to and supporting patients and carefully evaluating their complaints. Most patients with RLS can obtain symptomatic relief with commonly prescribed medications and support.

REFERENCES

1. Meissner HH, Riemer A, Santiago SM, Stein M, Goldman MD, Williams AJ. Failure of physician documentation of sleep complaints in hospitalized patients. *West J Med* 1998;169(3):146-149.
2. Ekblom KA. Restless legs syndrome. *Neurology* 1960;10:868-873.
3. Walters AS. Toward a better definition of the restless legs syndrome. *Mov Disord* 1995;10(5):634-642.
4. Ekblom KA. Restless legs. A report of 70 new cases. *Acta Med Scand* 1950;246(suppl):64-68.
5. Strang RR. The symptom of restless legs. *Med J Aust* 1967;1:1211-1213.
6. Lavigne GJ, Montplaisir JY. Restless legs syndrome and sleep bruxism: prevalence and association among Canadians. *Sleep* 1994;17(8):739-743.
7. Walters AS, Picchietti DL, Ehrenberg BL, Wagner ML. Restless legs syndrome in childhood and adolescence. *Pediatr Neurol* 1994;11(3):241-245.
8. Walters AS, Hickey K, Maltzman J, et al. A questionnaire study of 138 patients with restless legs syndrome: the 'Night-Walkers' survey. *Neurology* 1996;46:92-95.
9. Chokroverty S, Jankovic J. Restless legs syndrome. A disease in search of identity. *Neurology* 1999;52:907-910.
10. Montplaisir J, Boucher S, Poirier G, Lavigne G, Lapierre O, Lésperance P. Clinical, polysomnographic, and genetic characteristics of restless legs syndrome: a study of 133 patients diagnosed with new standard criteria. *Mov Disord* 1997;12(1):61-65.
11. Sun ER, Chen CA, Ho G, Earley CJ, Allen RP. Iron and the restless legs syndrome. *Sleep* 1998;21(4):371-377.
12. Aul EA, Davis BJ, Rodnitzky RL. The importance of formal serum iron studies in the assessment of restless legs syndrome. *Neurology* 1998;51:912.
13. Goodman JD, Brodie C, Ayida GA. Restless leg syndrome in pregnancy. *BMJ* 1988;297(6656):1101-1102.
14. Wetter TC, Stiasny K, Kohnen R, Oertel WH, Trenkwalder C. Polysomnographic sleep measures in patients with uremic and idiopathic restless legs syndrome. *Mov Disord* 1998;13(5):820-824.
15. Winkelman JW, Chertow GM, Lazarus JM. Restless legs syndrome in end-stage renal disease. *Am J Kidney Dis* 1996;28(3):372-378.
16. Yasuda T, Nishimura A, Katsuki Y, Tsuji Y. Restless legs syndrome treated successfully by kidney transplantation—a case report. *Clin Transpl* 1986;12:138.
17. Garvey MJ, Tollefson GD. Occurrence of myoclonus in patients treated with cyclic antidepressants. *Arch Gen Psychiatry* 1987;44:269-272.
18. Bakshi R. Fluoxetine and restless legs syndrome. *J Neurol Sci* 1996;142:151-152.
19. Terao T, Terao M, Yoshimura R, Abe K. Restless legs syndrome induced by lithium. *Biol Psychiatry* 1991;30:1167-1170.
20. Ward NG. Akathisia associated with droperidol during epidural anesthesia. *Anesthesiology* 1989;71:786-787.
21. Lutz EG. Restless legs, anxiety and caffeinism. *J Clin Psychiatry* 1978;39:693-698.
22. Ondo W, Jankovic J. Restless legs syndrome: clinicoetiologic correlates. *Neurology* 1996;47:1435-1441.

23. American Sleep Disorders Association. International classification of sleep disorders, revised: diagnostic and coding manual. Rochester, Minnesota: American Sleep Disorders Association, 1997. Pgs. 65-68.
24. Lugaresi E, Cirignotta F, Coccagna G, Montagna P. Nocturnal myoclonus and restless legs syndrome. *Adv Neurol* 1986;43:295-307.
25. Chesson AL Jr., Ferber RA, Fry JM, et al. The indications for polysomnography and related procedures. *Sleep* 1997;20(6):423-487.
26. Riley JD, Antony SJ. Leg cramps: differential diagnosis and management. *Am Fam Physician* 1995;52(6):1794-1798.
27. Walters AS, Hening W, Rubinstein M, Chokroverty S. A clinical and polysomnographic comparison of neuroleptic-induced akathisia and the idiopathic restless legs syndrome. *Sleep* 1991;14(4):339-345.
28. Hening W, Allen R, Earley C, Kushida C, Picchiotti D, Silber M. The treatment of restless legs syndrome and periodic limb movement disorder. *Sleep* 1999;22(7):970-999.
29. Montplaisir J, Nicolas A, Denesle R, Gomez-Mancilla B. Restless legs syndrome improved by pramipexole. A double-blind randomized trial. *Neurology* 1999;52:938-943.
30. Frucht S, Rogers JD, Greene PE, Gordon MF, Fahn S. Falling asleep at the wheel: motor vehicle mishaps in persons taking pramipexole and ropinirole. *Neurology* 1999;52:1908-1910.

APPENDIX: SLEEP/WAKE PROFILE

- How has the patient been sleeping recently?
(Ask patient and bed partner.)

SUGGESTED QUESTIONS FOLLOWING A SLEEP COMPLAINT

- When did the problem begin?
(To determine acute vs. chronic insomnia)
- Does the patient have a psychiatric or medical condition that may cause insomnia?
- Is the sleep environment conducive to sleep (relative to noise, interruptions, temperature, light)?
- Does the patient report “creeping, crawling, or uncomfortable, difficult-to-describe feelings” in the legs or arms that are relieved by moving them? (Relates to restless legs syndrome.)
- Does the bed partner report that the patient’s legs or arms jerk during sleep?
(Relates to periodic limb movement in sleep.)
- Does the patient snore loudly, gasp, choke, or stop breathing during sleep?
(Relates to obstructive sleep apnea.)
- Is the patient a shift worker? What are the work hours? (Relates to circadian sleep disorders/sleep deprivation.)
- What are the bedtimes and rise times on weekdays and weekends? (Relates to poor sleep hygiene and sleep deprivation.)
- Does the patient use caffeine, tobacco, or alcohol? Does the patient take over-the-counter or prescription medications (such as stimulating antidepressants, steroids, decongestants, beta-blockers)? (Relates to substance-induced insomnia.)

SIGNS OF SLEEPINESS

- What daytime consequences does the patient report?
- Does the patient report dozing off or difficulty staying awake during routine tasks, especially while driving?

WHERE TO GET MORE INFORMATION

- **National Center on Sleep Disorders Research (NCSDR)**
The NCSDR, located in the National Heart, Lung, and Blood Institute, supports research, scientist training, dissemination of health information, and other activities on sleep disorders and related concerns. The NCSDR also coordinates sleep research activities with other Federal agencies and with public and nonprofit organizations.

National Center on Sleep Disorders Research
Two Rockledge Centre
Suite 10038
6701 Rockledge Drive, MSC 7920
Bethesda, MD 20892-7920
(301) 435-0199
(301) 480-3451 (fax)

NCSDR Web site: <http://www.nhlbi.nih.gov/about/ncsdr>

- **National Heart, Lung, and Blood Institute (NHLBI) Health Information Network**
The NHLBI Health Information Network acquires, analyzes, promotes, maintains, and disseminates programmatic and educational information related to sleep and sleep disorders, as well as cardiovascular, pulmonary, and blood diseases. Write for a list of available publications or to order additional copies of this brochure.

NHLBI Health Information Network
P.O. Box 30105
Bethesda, MD 20824-0105
(301) 592-8573
(301) 592-8563

NHLBI Web site: <http://www.nhlbi.nih.gov>

- **Restless Legs Syndrome Foundation, Inc.**
The foundation is a nonprofit organization dedicated to helping the public, patients, families, and physicians better understand RLS.

Restless Legs Syndrome Foundation, Inc.
819 Second Street, S.W.
Rochester, MN 55902-2985

RLS Foundation Web site: www.rls.org

Discrimination Prohibited: Under provisions of applicable public laws enacted by Congress since 1964, no person in the United States shall, on the grounds of race, color, national origin, handicap, or age, be excluded from participation in, be denied the benefits of, or be subjected to discrimination under any program or activity (or, on the basis of sex, with respect to any education program or activity) receiving Federal financial assistance. In addition, Executive Order 11141 prohibits discrimination on the basis of age by contractors and subcontractors in the performance of Federal contracts, and Executive Order 11246 states that no federally funded contractor may discriminate against any employee or applicant for employment because of race, color, religion, sex, or national origin. Therefore, the National Heart, Lung, and Blood Institute must be operated in compliance with these laws and Executive Orders.

U.S. DEPARTMENT OF HEALTH AND
HUMAN SERVICES

Public Health Service
National Institutes of Health
National Heart, Lung, and Blood Institute

NIH Publication No. 00-3788
March 2000