Practice guideline summary: Treatment of restless legs syndrome in adults


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ABSTRACT

Objective: To make evidence-based recommendations regarding restless legs syndrome (RLS) management in adults.

Methods: Articles were classified per the 2004 American Academy of Neurology evidence rating scheme. Recommendations were tied to evidence strength.

Results and recommendations: In moderate to severe primary RLS, clinicians should consider prescribing medication to reduce RLS symptoms. Strong evidence supports pramipexole, rotigotine, cabergoline, and gabapentin enacarbil use (Level A); moderate evidence supports ropinirole, pregabalin, and IV ferric carboxymaltose use (Level B). Clinicians may consider prescribing levodopa (Level C). Few head-to-head comparisons exist to suggest agents preferentially. Cabergoline is rarely used (cardiac valvulopathy risks). Augmentation risks with dopaminergic agents should be considered. When treating periodic limb movements of sleep, clinicians should consider prescribing ferrous sulfate with vitamin C (Level B). When nonpharmacologic approaches are desired, clinicians should consider prescribing vitamin C and E supplementation (Level B) and may consider prescribing ropinirole, levodopa, or exercise (Level C). Neurology® 2016;87:2585–2593

GLOSSARY

AE = adverse event; CI = confidence interval; ESRD = end-stage renal disease; FCM = ferric carboxymaltose; HD = hemodialysis; IRLS = International Restless Legs Syndrome Study Group rating scale; NIRS = near-infrared spectroscopy; PLMI = Periodic Limb Movement Index; PLMS = periodic limb movements of sleep; PSG = polysomnography; QoL = quality of life; RLS = restless legs syndrome; rTMS = repetitive transcranial magnetic stimulation; TST = total sleep time; WASO = wake after sleep onset.

This document summarizes information provided in the complete guideline, available as a data supplement at Neurology.org. References e1–e20, cited in this summary, are available at Neurology.org.

Restless legs syndrome (RLS) is a movement disorder characterized by an urge to move the legs or arms, commonly in response to uncomfortable dysesthesia. Clinically important RLS affects approximately 2.5% of adults in the United States and Northern Europe, with higher prevalence in women and with increasing age. RLS is classified as primary or secondary in origin, with secondary RLS attributed to comorbid iron deficiency, end-stage renal disease (ESRD), or pregnancy. Most patients with RLS also have periodic limb movements of sleep (PLMS). Clinical consequences of RLS include impairment in sleep quality
and quantity, mood and anxiety disorders, worsening health-related quality of life (QoL), and loss of work productivity. Augmentation is a major side effect related to long-term RLS treatment with dopaminergic medication and consists of iatrogenic worsening of RLS symptoms.

This practice guideline addresses the following question: What are safe and effective therapies, including both pharmacologic and nonpharmacologic approaches, for the symptoms and clinical consequences (disturbed sleep, PLMS, depression/ anxiety, and decreased QoL) of RLS in adults?

**DESCRIPTION OF THE ANALYTIC PROCESS** This practice guideline follows the methodologies outlined in the 2004 edition of the American Academy of Neurology’s guideline development process manual. A detailed description of the process is available in the full-length guideline at Neurology.org. For RLS efficacy, the International Restless Legs Syndrome Study Group rating scale (IRLS) was the preferred outcome, and a change of 3 points was considered clinically meaningful. For studies reporting polysomnography (PSG) results, the panel prioritized evaluating certain outcomes such as the Periodic Limb Movement Index (PLMI), total sleep time (TST), sleep efficiency, sleep latency, and wake after sleep onset (WASO). Outcomes related to subjective sleep outcomes, psychiatric symptoms, and QoL are described when available. The table presents selected adverse events (AEs), augmentation risks, and US Food and Drug Administration–approved doses for recommended medications.

Results of individual articles, including confidence intervals (CIs) and assessments of statistical significance and clinical relevance, are available in the full guideline at Neurology.org. Circumstances for which only 1 Class III study is available (for which no conclusions can be drawn, e.g., gabapentin) are also discussed only in the full guideline.

**ANALYSIS OF EVIDENCE** Dopamine agonists. **Ropinirole.** It is likely that ropinirole decreases IRLS scores at 12 weeks (meta-analysis of 2 Class I studies, of which 1 had sufficient precision independently). It is highly likely that ropinirole improves PLMS (2 Class I studies and 4 Class II studies using Medical Outcomes Study subscales). It is likely that ropinirole improves RLS-specific QoL at 12 weeks (1 Class I and 3 Class II studies). It is possible that ropinirole improves depression (meta-analysis of 1 Class II study and 1 Class I study with insufficient precision) and likely that it improves anxiety at 12 weeks (1 Class I study).

**Pramipexole.** It is highly likely that pramipexole improves RLS symptoms as measured by the IRLS (3 Class I and 6 Class II studies over varying timeframes). It is likely that pramipexole improves PLMS (3 Class II studies and subjective sleep measures (1 Class I and 3 Class II studies, with one of the Class II studies showing limited improvement). It is possible that pramipexole improves depression and anxiety at 12 weeks in patients with moderate to severe RLS-related mood disturbance (1 Class II study).

**Rotigotine.** It is highly likely that the rotigotine patch improves RLS symptoms as measured by the IRLS (2 Class I and 3 Class II studies, up to 6 months in duration). It is likely that rotigotine improves PLMS (1 Class I study), but there is insufficient evidence to support or refute an effect on other objective sleep measures (1 Class I study that is not statistically significant but whose CIs include clinically important effects). It is likely that rotigotine improves sleep disturbance and subjective sleep quantity (meta-analysis of 1 Class I and 2 Class II studies with 1 of the Class II studies achieving statistical significance on its own and the other Class I and Class II studies achieving statistical significance together). Rotigotine possibly improves sleep adequacy (meta-analysis of 1 Class I and 2 Class II studies that requires all 3 studies to achieve significance). Rotigotine possibly improves RLS-specific QoL at 12 weeks (meta-analysis of 1 Class I and 2 Class II studies requiring all 3 studies to achieve significance).

**Cabergoline.** Cabergoline is rarely used, as there are concerns regarding fibrotic complications/cardiac valvulopathy (see Discussion in the full guideline at Neurology.org).

**Levodopa.** Levodopa (100–200 mg) possibly improves patient-reported RLS symptom severity (4 Class III studies, of which show a benefit alone and 2 of which show a benefit when combined in a meta-analysis to increase statistical precision). Levodopa possibly improves subjective sleep measures (4 Class III studies, with improvements in at least some subjective sleep measures) and the PLMI (3 Class III...
studies with sufficient precision and 1 Class III study with insufficient precision; meta-analysis showed significant effect). There is insufficient evidence to support or refute the effect of levodopa on QoL in RLS (2 Class III studies, only 1 with sufficient precision).

α2δ ligands. Gabapentin enacarbil. Gabapentin enacarbil is a slow-release gabapentin prodrug. It is highly likely that gabapentin enacarbil decreases IRLS scores (4 Class I studies with different study durations) and likely that it improves at least some subjective sleep measures (4 Class I studies) and likely that it improves at least some objective sleep measures other than the PLMI (1 Class I study). Because results of this Class I study were not statistically significant and CIs included both potentially clinically important and unimportant effects, there is insufficient evidence to support or refute the effect of gabapentin enacarbil on the PLMI. It is likely that gabapentin enacarbil improves RLS-specific QoL (1 Class I study) and mood (1 Class I study).

Pregabalin. Pregabalin likely improves IRLS scores at doses of at least 150 mg/d (1 Class I and 3 Class II studies; there is insufficient evidence to support or refute doses of 50–100 mg/d because analyses did not reach statistical significance but CIs

<table>
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<th>Table</th>
<th>Summary of interventions evaluated in idiopathic restless legs syndrome (RLS) with Level A–C recommendations</th>
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<tr>
<td>Intervention</td>
<td>FDA guidelines for starting dose, therapeutic dose, mg/d</td>
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<tr>
<td>Ropinirole</td>
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<td>Pramipexole</td>
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<td>Rotigotine patch (worn 24 h/d)</td>
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<tr>
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<tr>
<td>Pregabalin</td>
<td>Not FDA-approved for RLS</td>
</tr>
<tr>
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<tr>
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<td>tDCS</td>
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Abbreviations: AE = adverse event; FDA = US Food and Drug Administration; NA = not applicable; NIRS = near-infrared spectroscopy; PLMI = Periodic Limb Movement Index; rTMS = repetitive transcranial magnetic stimulation; tDCS = transcranial direct current stimulation.

* Level of evidence cited is the highest level of evidence identified for at least one subjective sleep rating; subjective sleep ratings are considered individually in the guideline text, with sometimes differing levels of evidence by measure. Refer to full guideline at Neurology.org for details on different subjective measures.

† Augmentation marked as yes if present in >2.4% at any timepoint in available studies (many of which are Class IV open-label long-term follow-up); the 2.4% cutoff was determined by averaging placebo augmentation responses from 3 studies (see text).

‡ Augmentation listed as unknown because studies describing augmentation were 12 weeks or less in duration and thus cannot reliably inform augmentation risks (augmentation typically develops after at least 6 months of treatment).

§ Oral studies were included only if patients had evidence of iron deficiency.
improves RLS symptoms, sleep adequacy, sleep duration, and RLS-specific QoL in patients with RLS who have not responded to other treatments (1 Class II study39).

Other medications and nutraceuticals. Other medications and nutraceuticals are discussed in the full guideline at Neurology.org.

Physical measures. Near-infrared spectroscopy (NIRS). NIRS is possibly effective in the treatment of primary moderate to severe RLS (1 Class II study vs sham66 and 1 Class II study showing no difference between 2 devices67).

Pneumatic compression. Pneumatic compression is likely effective in the treatment of patients with primary moderate to severe RLS (1 Class I study69).

Transcranial direct current stimulation. Cathodal and anodal transcranial direct current stimulation are probably ineffective for improving RLS symptoms in women with RLS who were drug-naive (one negative Class I study39).

Repetitive transcranial magnetic stimulation (rTMS). rTMS is possibly effective in the treatment of primary moderate to severe RLS (1 Class II study70).

Vibrating pads. Vibrating pads are possibly ineffective in treating RLS symptoms (meta-analysis of 2 Class II studies71-73 excluding a clinically important benefit) but possibly effective in treating subjective sleep outcomes (meta-analysis of 2 Class II studies74,75 where only one was sufficient to drive recommendations on its own). There is insufficient evidence to support or refute an effect of vibrating pads on QoL in RLS (meta-analysis of 2 Class II studies74,75 that is not statistically significant but where the CI includes a potentially clinically important effect).

Treatment of secondary RLS. There are many causes of secondary RLS. However, adequate evidence is available only for treatment of secondary RLS in patients with ESRD who are on hemodialysis (HD).

Ropinirole. Ropinirole 0.25 mg daily is possibly effective in the treatment of RLS symptoms associated with ESRD/HD (1 Class II study76).

Levodopa. Levodopa (100-200 mg) is possibly effective in treating PLMS associated with RLS (2 Class III studies77-79), but there is insufficient evidence to support or refute an effect of levodopa on RLS severity (2 Class III studies with insufficient precision/details80).

Opioid agonists. It is possible that prolonged-release oxycodone/naloxone (mean dose of oxycodone 21.9 ± 15.0 mg, naloxone 11.0 ± 7.5 mg) improves RLS symptoms, sleep adequacy, sleep...
PRACTICE RECOMMENDATIONS

1. In moderate to severe primary RLS, clinicians should consider prescribing a pharmacologic agent to reduce RLS symptoms. There is strong evidence to support the use of pramipexole, rotigotine, cabergoline, and gabapentin enacarbil (Level A); moderate evidence to support the use of ropinirole, pregabalin, and IV FCM (Level B); and weak evidence to support the use of levodopa (Level C). There are few head-to-head comparisons of these agents to suggest that one should be used preferentially, though in practice clinicians often decide on the basis of comorbidities or potential side effects such as augmentation with dopaminergic agents. When considering efficacy alone, clinicians may consider choosing cabergoline instead of levodopa (Level C). However, cabergoline is rarely used in clinical practice for RLS because of a risk of cardiac valvulopathy at higher doses. There is insufficient evidence to support or refute the preferential use of pregabalin instead of pramipexole (Level U).

2. For patients with primary RLS for whom clinicians want to target sleep, clinicians should consider prescribing a pharmacologic agent that improves objective or subjective sleep parameters (or both). Evidence supports agents to different extents for subjective and objective outcomes.
   a. When targeting PLMS, specifically the PLMI as measured by PSG, there is strong evidence to support the use of ropinirole (Level A); moderate evidence to support the use of pramipexole, rotigotine, cabergoline, and pregabalin (Level B); and weak evidence to support the use of levodopa (Level C). There is insufficient evidence to support or refute the use of gabapentin enacarbil, FCM, or iron sucrose for PLMS (Level U). There is weak evidence (Level C) for using pramipexole in preference to pregabalin with regard to PLMI alone.
   b. With regard to other objective sleep measures (e.g., TST, sleep efficiency, sleep latency, and WASO), there is moderate evidence to support the use of ropinirole, gabapentin enacarbil, and pregabalin for at least some objective sleep measures (Level B). There is insufficient evidence to support or refute the use of pramipexole, rotigotine, cabergoline, or levodopa for these measures (Level U). There is weak evidence (Level C) for using pregabalin in preference to pramipexole with regard to objective sleep measures other than PLMI.
   c. With regard to subjective sleep measures, there is strong evidence to support the use of cabergoline and gabapentin enacarbil (Level A); moderate evidence to support the use of ropinirole, pramipexole, and pregabalin (Level B); weak to moderate evidence to support the use of rotigotine (Levels B and C); and weak evidence to support the use of levodopa (Level C), with the strength of evidence varying by measure and, sometimes, dose. There is insufficient evidence to support or refute the use of FCM for subjective sleep measures (Level U). There is moderate evidence to support the use of pregabalin instead of pramipexole with regard to subjective sleep outcomes (Level B).

3. For patients with RLS for whom clinicians want to target concomitant psychiatric symptoms, clinicians should consider ropinirole in the context of anxiety (Level B) and may consider ropinirole in the context of depression (Level C). In the context of moderate to severe RLS-related mood disturbance, clinicians may consider prescribing pramipexole for depression and anxiety (Level C). For overall mood, clinicians should consider prescribing gabapentin enacarbil (Level B).

4. For patients with RLS for whom clinicians want to select an agent that improves QoL, clinicians should consider prescribing ropinirole, pramipexole, cabergoline, gabapentin enacarbil, or IV FCM (Level B) and may consider prescribing rotigotine or pregabalin (Level C). There is insufficient evidence to support or refute the use of levodopa for improving QoL in RLS (Level U).

5. When avoidance of augmentation is a deciding factor, clinicians may consider prescribing pregabalin rather than pramipexole when considering 52-week treatment in light of lower augmentation rates with pregabalin (Level C). Clinicians may also consider prescribing cabergoline rather than levodopa when considering 30-week treatment in light of lower augmentation rates with cabergoline (Level C); however, this needs to be weighed against the risk of cardiac valvulopathy with high doses of cabergoline. There is insufficient evidence to support or refute which dopaminergic agents cause the least augmentation because augmentation rates are most commonly reported in long-term open-label Class IV studies (Level U). Results of these studies are summarized in the full guideline at Neurology.org but cannot support formal recommendations.

6. For patients with RLS who have not responded to other treatments, clinicians may consider...
When addressing RLS, clinicians and patients must first determine whether symptoms require treatment, the setting in which this practice guideline is relevant. Treatment should be considered if RLS symptoms interfere with sleep or daytime function to an important degree. Before determining the best treatment, it is important to first ensure there are no contributing factors to RLS symptoms (e.g., iron deficiency or serotonergic antidepressants). Because iron deficiency is a known contributor to RLS, can result in other complications, and may respond to iron supplementation, it is reasonable for clinicians to check iron studies in patients with RLS with new or worsening symptoms and treat the iron deficiency first if indicated.

There are important limitations in the evidence regarding RLS treatments. The clinical significance of some outcomes used in RLS trials, such as PLMI, is uncertain; thus conclusions drawn regarding these outcomes are of unknown clinical relevance. In addition, apart from the IRLS, clinically important differences for the measures used in RLS trials are unknown. Most of the studies are short-term trials, often 12 or fewer weeks, whereas clinical treatment of RLS is ongoing over years. Conclusions regarding long-term efficacy and risks are difficult to develop because of the open-label nature of many longer duration studies. Short-term trials are less able to inform risks associated with prolonged medication exposure, such as augmentation occurring with dopaminergic medications. Augmentation is a major concern and an important consideration when choosing a treatment approach. Long-term risks with other treatment approaches, such as opioid use, are also important to consider.

Though some patients have RLS symptoms intermittently, the value of PRN medications is unknown.1 Additionally, there are no data to guide the approach to cases where monotherapy is not adequately effective or clinicians want to use multiple agents to minimize doses of dopaminergic agents. Clinical trials of RLS medications generally exclude patients with common comorbid conditions such as mood and anxiety disorders and peripheral neuropathy. Generalizability of studies to populations with these disorders is uncertain.49 Certain populations with secondary RLS, such as pregnant women, are also under-studied.

In patients with RLS symptoms requiring treatment, choosing the most appropriate intervention requires an individualized approach including regard for patient factors, such as the most prominent symptoms (e.g., presence of sleep disturbance, because of varying strength of evidence by outcome), comorbidities relating to RLS (e.g., mood), other comorbidities (such that an agent may be used preferentially to treat more than one indication or avoided because of a presumed higher risk of side effects), age, side effect profile, augmentation risks, and patient preferences (e.g., pharmacologic or nonpharmacologic approaches). In addition to AEs commonly reported in trials, some agents for RLS have less common but important risks, including cardiac valvulopathy with cabergoline and impulse control disorders with dopamine agonists.

Given the chronicity of RLS, long-term risks of augmentation with dopaminergic agents are relevant for many patients. Scant data exist to guide the decision-making process relating to augmentation.120 For patients on dopaminergic agents, careful reassessment of changes in the time of symptom onset, anatomical distribution, total medication dose, and

prescribing prolonged-release oxycodone/naloxone (where available) for RLS symptoms, subjective sleep symptoms, and QoL (Level C), but potential benefits need to be weighed against known opioid risks.

7. There is insufficient evidence to support or refute the use of gabapentin, iron sucrose, oxycodone, clonazepam, bupropion, clonidine, selenium, rifaximin, botulinum neurotoxin, valproic acid, carbamazepine, or valerian in the treatment of RLS (Level U).

8. For patients or clinicians wanting to use nonpharmacologic approaches to treat RLS, clinicians should consider prescribing pneumatic compression before usual symptom onset (Level B) and may consider prescribing NIRS or rTMS (where available) (Level C). Clinicians may consider prescribing vibrating pads for subjective sleep concerns (Level C) but not for RLS symptoms (Level C against). Clinicians may also choose not to consider transcranial direct current stimulation for RLS symptoms (Level C against). There is insufficient evidence to support or refute use of acupuncture in RLS (Level U).

9. In patients with RLS and serum ferritin ≤75 µg/L, clinicians should consider prescribing ferrous sulfate with vitamin C for improvement of RLS symptoms (Level B).

10. In patients with secondary RLS associated with ESRD on HD, clinicians should consider prescribing vitamin C and E supplementation (alone or in combination) (Level B) and may consider prescribing ropinirole, levodopa, or exercise (Level C). There is insufficient evidence to support or refute the use of gabapentin or IV iron dextran in RLS associated with ESRD/HD (Level U). There is also insufficient evidence to support or refute the use of gabapentin or levodopa preferentially over the other in this population (Level U).

CLINICAL CONTEXT When addressing RLS, clinicians and patients must first determine whether symptoms require treatment, the setting in which this practice guideline is relevant. Treatment should be considered if RLS symptoms interfere with sleep or daytime function to an important degree. Before determining the best treatment, it is important to first ensure there are no contributing factors to RLS symptoms (e.g., iron deficiency or serotonergic antidepressants). Because iron deficiency is a known contributor to RLS, can result in other complications, and may respond to iron supplementation, it is reasonable for clinicians to check iron studies in patients with RLS with new or worsening symptoms and treat the iron deficiency first if indicated.

There are important limitations in the evidence regarding RLS treatments. The clinical significance of some outcomes used in RLS trials, such as PLMI, is uncertain; thus conclusions drawn regarding these outcomes are of unknown clinical relevance. In addition, apart from the IRLS, clinically important differences for the measures used in RLS trials are unknown. Most of the studies are short-term trials, often 12 or fewer weeks, whereas clinical treatment of RLS is ongoing over years. Conclusions regarding long-term efficacy and risks are difficult to develop because of the open-label nature of many longer duration studies. Short-term trials are less able to inform risks associated with prolonged medication exposure, such as augmentation occurring with dopaminergic medications. Augmentation is a major concern and an important consideration when choosing a treatment approach. Long-term risks with other treatment approaches, such as opioid use, are also important to consider.

Though some patients have RLS symptoms intermittently, the value of PRN medications is unknown.1 Additionally, there are no data to guide the approach to cases where monotherapy is not adequately effective or clinicians want to use multiple agents to minimize doses of dopaminergic agents. Clinical trials of RLS medications generally exclude patients with common comorbid conditions such as mood and anxiety disorders and peripheral neuropathy. Generalizability of studies to populations with these disorders is uncertain.49 Certain populations with secondary RLS, such as pregnant women, are also under-studied.

In patients with RLS symptoms requiring treatment, choosing the most appropriate intervention requires an individualized approach including regard for patient factors, such as the most prominent symptoms (e.g., presence of sleep disturbance, because of varying strength of evidence by outcome), comorbidities relating to RLS (e.g., mood), other comorbidities (such that an agent may be used preferentially to treat more than one indication or avoided because of a presumed higher risk of side effects), age, side effect profile, augmentation risks, and patient preferences (e.g., pharmacologic or nonpharmacologic approaches). In addition to AEs commonly reported in trials, some agents for RLS have less common but important risks, including cardiac valvulopathy with cabergoline and impulse control disorders with dopamine agonists.

Given the chronicity of RLS, long-term risks of augmentation with dopaminergic agents are relevant for many patients. Scant data exist to guide the decision-making process relating to augmentation.120 For patients on dopaminergic agents, careful reassessment of changes in the time of symptom onset, anatomical distribution, total medication dose, and
medication timing are indicated at least yearly. In the absence of evidence, it is reasonable to consider discontinuing a patient’s dopaminergic medication in the setting of clinically important augmentation and switching to a nondopaminergic agent or a longer-acting dopaminergic medication.

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**CONFLICT OF INTEREST** The American Academy of Neurology (AAN) is committed to producing independent, critical, and truthful clinical practice guidelines (CPGs). Significant efforts are made to minimize the potential for conflicts of interest to influence the recommendations of this CPG. To the extent possible, the AAN keeps separate those who have a financial stake in the success or failure of the products appraised in the CPGs and the developers of the guidelines. Conflict of interest forms were obtained from all authors and reviewed by an oversight committee prior to project initiation. The AAN limits the participation of authors with substantial conflicts of interest. The AAN forbids commercial participation in, or funding of, guideline projects. Drafts of the guideline have been reviewed by at least 3 AAN committees, a network of neurologists, Neurology® peer reviewers, and representatives from related fields. The AAN Guideline Author Conflict of Interest Policy can be viewed at aann.com. For complete information on this process, access the 2004 AAN process manual.9

**AUTHOR CONTRIBUTIONS**

Dr. Winkelman: study concept and design, acquisition of data, analysis or interpretation of data, drafting/revising the manuscript, critical revision of the manuscript for important intellectual content, study supervision.

Dr. Armstrong: study concept and design, acquisition of data, analysis or interpretation of data, drafting/revising the manuscript, critical revision of the manuscript for important intellectual content, study supervision.

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Dr. Gronseth: study concept and design, acquisition of data, analysis or interpretation of data, drafting/revising the manuscript, critical revision of the manuscript for important intellectual content.

J. Winkelman currently serves on scientific advisory boards for Merck and Flex Pharma and has served on scientific advisory boards for UCB, Impax, Pfizer, Lacerma, Lundpold Pharmaceuticals, GlaxoSmithKline, Boehringer-Ingeheim, Xenoprot, Zan Inc., Sunovion, Takeda, Jazz, and Neurogen; currently performs neurophysiology studies as part of his practice; currently serves as a journal editor for the following publications: Sleep, Sleep Medicine, and CNS Drug; has received honoraria from or served on speakers bureaus for the following organizations: Boehringer-Ingeheim, GlaxoSmithKline, Pfizer, Sepracor (now Sunovion), Takeda, Lundpold Pharmaceuticals, Novartis, Neurogen, and UCB (Schwarz Pharma); has received research support from Boehringer-Ingeheim, GlaxoSmithKline, UCB (Schwarz Pharma), Sepracor (now Sunovion), and Pfizer; holds stock in Flex Pharma; receives publishing royalties for the following publications: Foundations of Psychiatric Sleep Medicine (Cambridge University Press, 2010) and an UpToDate chapter on nocturnal leg cramps; receives government research support from the National Institute of Mental Health (1R01MH097972-01A1, PI), and has given expert testimony for legal cases representing generic manufacturers of pharmaceuticals approved for the treatment of insomnia and narcolepsy. M. Armstrong receives compensation from the American Academy of Neurology (AAN) as an evidence-based medicine methodologist and serves on the Level of Evidence editorial board for Neurology but is not compensated financially. R. Allen has served on a volunteer basis for the International Restless Legs Syndrome Study Group and the World Association of Sleep Medicine, has served on scientific advisory boards for Pfizer, GlaxoSmithKline,
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