Sleep related movement disorders?

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Nocturnal Frontal Lobe Epilepsy (NFLE)

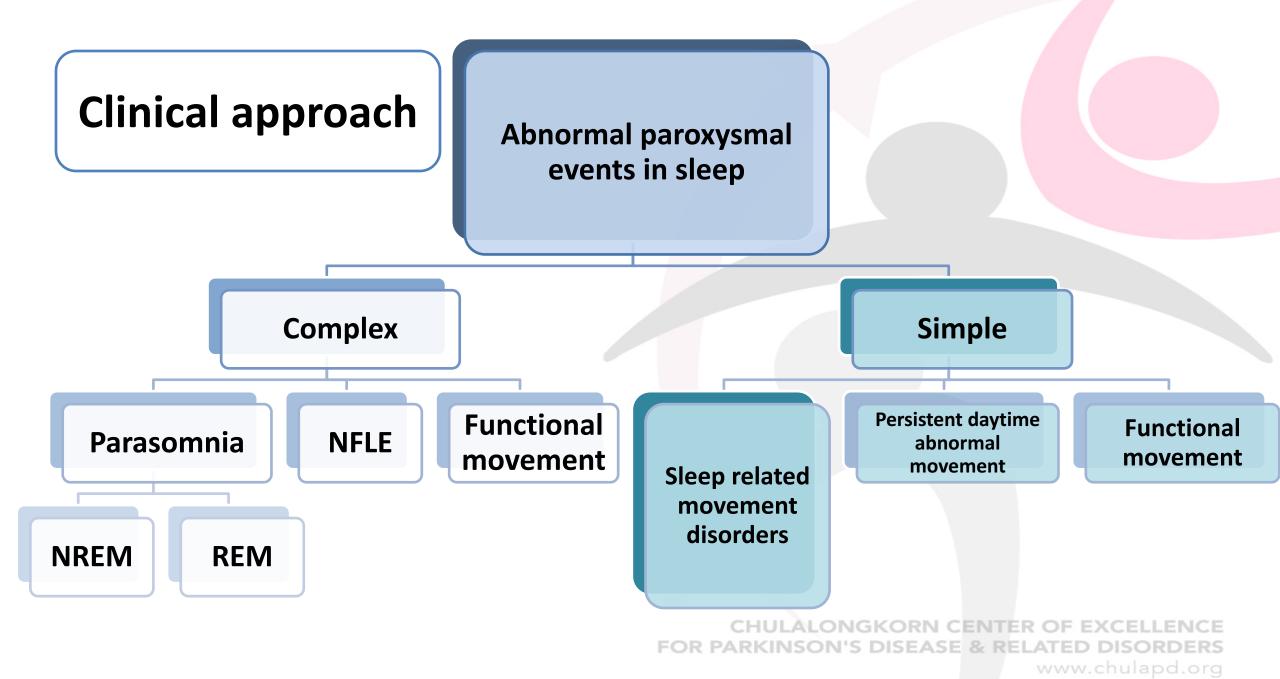
- Seizures in NFLE may have bizarre clinical features, with vocalization, complex automatisms, and ambulation (only in the bedroom)
- Typically of the hyper-motor type,
 - Repetitive, high-amplitude, and high-velocity movements of the trunk and proximal extremities and
 - Asymmetric tonic seizures having dystonic, dyskinetic, and repetitive proximal movements.
- Complex motor seizures in NFLE have hypermotor manifestations,
 - Marked agitation with body rocking, kicking, boxing, thrashing, pedaling, bending, hitting, running, spitting, and various types of vocalization that include shouting and swearing

Classification of movement disorders during sleep

Physiologic Motor Activity during sleep or at sleep onset	Pathologic Motor Activity during Sleep
Postural shifts, body& limb movements during sleep	Motor parasomnias
Physiologic fragmentary myoclonus	Sleep-Related Movement Disorders
Hypnic jerks	Isolated Sleep-Related Motor Symptoms (Apparently Normal Variants)
Hypnagogic foot tremor	Nocturnal seizures
Rhythmic leg movements	Miscellaneous Nocturnal Motor Hyperactivity
	Voluntary Movement Disorders (Sleep-related Dissociative disorders)

Abnormal paroxysmal events in sleep

- How to perform clinical approach of this abnormal paroxysmal events in sleep
 - Sleep related movement disorders
 - Parasomnias
 - Non REM parasomnias
 - REM parasomnias
 - Frontal Lobe Epilepsy and Parasomnias Scale (FLEP)
 - Other sleep related epilepsy
 - Other persistent daytime abnormal movements
 - Sleep-related Dissociative Disorders



Derry CP, Arch Neurol 2006.

International classification of Sleep disorders 3rd edition (ICSD-3)

Insomnia

Sleep related breathing disorders

Central disorders of hyper somnolence

Circadian rhythm sleep wake disorders

Parasomnias

Sleep- related movement disorders

Other sleep disorders

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Sleep related movement disorders

Characterized by relatively simple, usually stereotyped movements that disturb sleep

VS

Parasomnias

- Undesirable physical or experiential events that accompany sleep without disrupting sleep architecture.
- Complex behaviors during sleep that *appear purposeful and goal-directed*, but are outside the conscious awareness of the individual and are therefore inappropriate
- Consist of abnormal sleep-related movements, behaviors, emotions, perceptions, dreaming, and autonomic nervous system functioning
- Disorders of arousal, partial arousal, and sleep-stage transition

Sleep-related movement disorders

Restless legs syndrome

Periodic limb movement disorder

Sleep-related leg cramps

Sleep-related bruxism

Sleep-related rhythmic movement disorder

Benign sleep myoclonus of infancy

Propriospinal myoclonus at sleep onset

Sleep-related movement disorder due to a medical disorder

Sleep-related movement disorder due to a medication or substance

Sleep-related movement disorder, unspecified

Parasomnias

NREM-related parasomnias

Confusional arousals

Sleepwalking

Sleep terrors

Sleep-related eating disorder

REM-related parasomnias

REM sleep behavior disorder

Recurrent isolated sleep paralysis

Nightmare disorder

Other parasomnias

Exploding head syndrome

Sleep-related hallucinations

Sleep enuresis

Parasomnia due to a medical disorder

Parasomnia due to a medication or substance

Parasomnia, unspecified

Isolated Symptoms and Normal Variants

Excessive fragmentary myoclonus

Hypnagogic foot tremor and alternating leg muscle activation

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Sleep starts (Hypnic jerks)

Michael J, Chest. 2014

Restless leg syndrome (RLS)

• Willis-Ekbom disease (WED): A circadian disorder of sensory-motor integration manifested.

Criteria	Characteristics
The <i>urge</i> to move the legs	 Usually but not always accompanied by or felt to be caused by uncomfortable and unpleasant sensations in the legs "need to move," "crawling," "tingling," "rest-less," "cramping," "creeping," "pulling," "painful," "electric," "tension," "itching," "burning," "prickly Typical RLS sensations are felt deep inside the muscles and bones of the legs Bilateral legs (usually)
Begin or worsen during periods of rest or inactivity such as lying down or sitting	
Partially or totally <i>relieved by movement</i> , such as walking or stretching, at least as long as the activity continues	
Only occur or are worse in the evening or night than during the day	
The above features are not solely accounted for by other medical or behavioral conditions	Exclude: myalgia, venous stasis, leg edema, arthritis, leg cramps, positional discomfort, habitual foot tapping, and other nocturnal sensory-motor symptoms.

Restless leg syndrome (RLS)

Clinical features supporting the diagnosis of restless legs syndrome/Willis-Ekbom disease (RLS/WED).

The following features, although not essential for diagnosis, are closely associated with RLS/WED and should be noted when present:

- Periodic limb movements (PLM): presence of periodic leg movements in sleep (PLMS) or resting wake (PLMW) at rates or intensity greater than expected for age or medical/medication status.
- Dopaminergic treatment response: reduction in symptoms at least initially with dopaminergic treatment.
- Family history of RLS/WED among first-degree relatives.
- Lack of profound daytime sleepiness.^a

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a RLS/WED shares this characteristic with other hyperarousal conditions including insomnia disorder.

Pathogenesis

- Not fully understood
- Genetic factors
 - BTBD9, MEIS1, and MAP2K5/SKOR1 genes
- Dopaminergic, adenosinergic, and glutamatergic pathways involved
 - 'hyperdopaminergic state'
- Low central nervous system (CNS) iron stores may lead to downregulation of the adenosine 1 receptor (A1R)
 - Contributes to both a hyperdopaminergic and hyper-glutamatergic state
- Deficiency in CNS opioid receptors in patients with RLS compared with controls

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Restless legs syndrome associated with major diseases

A systematic review and new concept

Primary/Idiopathic

Familial & genetic risk factors

Other associated conditions

- Arthritis,
- Sensory neuropathy,
- Radiculopathy
- Diabetes,
- Renal disease
- Psychiatric disorders such as depression and
- anxiety is increased 1.5- to 2-fold in patients with RLS

Secondary/ symptomatic

- Iron deficiency anemia (IDA),
- Multiple sclerosis (MS),
- Polyneuropathy, and
- Parkinson disease (PD)
- Common chronic diseases such as
 - Arterial hypertension or
 - Headache or
 - Conditions such as inflammation and pregnancy.



RLS mimic

RLS mimic	Characteristics			
Neuropathy	 Often worse at night and causes discomfort Not associated with an urge to move the legs Not relieved by movement 			
Akathisia	Positive for an urge/ need to move body or restlessness Does not have a circadian evening propensity The sensation of restlessness is typically diffuse without predilection for the legs			
Muscle cramps	 Predominantly or solely occur during sleep Result in painful muscle contractions Not associated with an urge to move 			
Habitual foot taping	 A behavioral phenomenon that is not associated with an urge to move, and It can be suppressed. 			
Positional discomfort	 From pressure that compresses nerves, limits blood flow, or stretches body tissue. Resolved by changing body position without requiring any repetitive movement No urge to move 			
Sleep starts (or hypnic jerks)	 Short, massive body movements during the sleep/wake transition Involve the extremities of both sides synchronously but without periodicity 			

RLS mimic

RLS mimic	Characteristics			
Sleep-related leg cramps	 Worse at night and are relieved by movement Involve a specific muscle and usually require stretching of the muscle more than moving of the leg to relieve symptoms Residual pain or sensitivity after the event 			
Painful legs and moving toes syndrome	 Severe pain in one or both feet A sensation of burning and associated with frequent, repetitive movements of the toes Pain is not necessarily increased at night or relieved by movement. 			
Vascular intermittent claudication	 Leg pain or discomfort Relieved by rest and worsens during a pro- longed upright position or walking. Absence of the typical circadian feature of RLS. 			
Pain related to the narrow spinal canal	 When lying in bed, pain is more pronounced in the supine position compared to a l position with bended knees. Combination with back pain, as well as with clinical signs of a myelopathy 			

BRIEF REPORT

Restless Leg Syndrome and Objectively-Measured Atherosclerosis in the Canadian Longitudinal Study on Aging

Key Words: restless leg syndrome; carotid intimamedia thickness; atherosclerosis; CLSA

Reported: RLS & major diseases

- Renal failure
- HT
- DM
- Obesity
- Thyroid dysfunction
- Depression
- Anxiety

USG Carotid intimamedia thickness

Excluded PLMS

RLS had significant carotid intima-media thickness more than non RLS patients

RLS is associated with a higher degree of atherosclerosis

Sympathetic hyperactivity, autonomic fluctuations, and hypertension in PLMS

Sleep deprivation & increase appetite/ sugar craving

RLS CHULALONGKORN CEN

Atherosclerosis

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Microcirculation impairment in the legs and/or global hypoxia might cause leg discomfort

(Non-pharmacological) Treatment

- Serum Iron : keep > 50-75 mcg/ml
- Transferrin saturation : keep > 20%
- Check exacerbating factors:
 - Lifestyle: Alcohol, smoking, sleep deprivation
 - Drugs:
 - Dopamine antagonist
 - Antidepressants SSRI, TCA
 - Anti-histamine
 - Recent opioid discontinuation
 - Blood loss
 - Pregnancy (iron)

Pharmacological Treatment

Drugs	Effective	Probably effective
Pramipexole	6 mo	1 year
Ropinirol	6 mo	1 year
Rotigotine	6 mo	5 years
Levodopa	_	2 years
Pregabalin	1 year	-
Gabapentin	_	_

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Dopamine agonists recommendation in RLS

			Side effects
D3 agents	Efficacy	Efficacy Safety	
Pramipexole	Efficacious at doses of 0.25, 0.50, and 0.75 mg	Acceptable risk with special monitoring for augmentation	Nausea, lightheadedness, fatigue, augmentation
Rotigotine	Efficacious at a dose of 2– 3mg	Acceptable risk with special monitoring for local site reactions and augmentation	Skin reaction, augmentation
0.78–4.6 mg spec		Acceptable risk with special monitoring for augmentation	Nausea, lightheadedness, fatigue, augmentation Winkelmann J, et al. Mov Disord 2018 Casoni F, Adv in Phar Vol 48. 2019

Treatment

Table 5

Clinical consensus of the benefits and risks for each pharmacologic treatment of RLS/WED.

	Levodopa	Nonergot DA		Ergot-based DA	α₂δ Ligand	Opioid	Clonazepam
		Short-acting	Long-acting				
The potential of the drug to cause it	the following adverse even	ts					
Augmentation	+++	++	+	++	0	NK	0
LoE	+++	++	NK	++	+	+	NK
ICD	0	+	0/+	NK	0	0	0
EDS	NK	++	+	++	+++	+	++
Negative mood	0	0	0	0	+	+	++
Weight gain	0	0	0	0	++	0	0
General toxicity	+	+	++	+++	+	++	+
The potential of the drug to have p	ositive effect on these para	imeters					
Subjective nighttime sleep	0	+	+	+	++	++	++
Classic nighttime RLS symptoms	+	++	++	++	++	++	0
QoL	NK	++	++	++	++	NK	NK
Pain reduction	+	+	+	+	++	+++	0

Abbreviations: RLS/WED, restless legs syndrome/Willis-Ekbom disease; DA, dopamine-receptor agonist; LoE, loss of efficacy; ICD, impulse control disorders; EDS, excessive daytime sleepiness; QoL, quality of life; NK, not known.

^{+++,} is very likely to affect this parameter; ++, is somewhat likely to affect this parameter; +, is slightly likely to affect this parameter; 0, has no effect on this parameter.

Initial Treatment: Factors that impact the choice of agent

Table 6
Clinical recommendations regarding factors that affect the selection of an agent for initial treatment in patients with restless legs syndrome/Willis-Ekbom disease.

Factor that impacts the choice of agent	Treatment choice		
Time of day (daytime disturbance)	Preferably a long-acting agent		
	 Twice a day dosing of a short-acting agent 		
Sleep disturbance disproportionate to other symptoms of RLS/WED	α ₂ δ Ligand		
Comorbid insomnia	α ₂ δ Ligand		
Pregnancy risk	 Avoid both dopaminergic agents and α2 δ ligands 		
	 Consider the use of iron 		
Impaired renal function	 Select a drug that is not renally excreted 		
Increased risk for falls	Dopamine-receptor agonist		
Painful restless legs	α ₂ δ Ligand		
Comorbid pain syndrome	α ₂ δ Ligand		
History of or current ICD	α ₂ δ Ligand		
History of or current alcohol or substance abuse	Dopamine-receptor agonist or α2δ ligand		
Severe symptoms of RLS/WED	Dopamine-receptor agonist		
Excess weight, metabolic syndrome, or obstructive sleep apnea	Dopamine-receptor agonist		
Availability ²	Dopamine agonist or α ₂ δ ligand		
Cost ^b	Dopamine agonist or α ₂ δ ligand		
Comorbid depression	Dopamine-receptor agonist		
Comorbid generalized anxiety disorder	α₂δ Ligand		
Daytime sleepiness	 Investigate the cause 		
Higher potential for drug interactions	 Select drug that is not hepatically excreted 		

Restless legs syndrome (RLS) / Willis—Ekbom disease with augmentation

 RLS = a sensorimotor disorder with leg discomfort and the irresistible urge to move affected body parts with relieve by movement & usually occur in the evening.

Augmentation is the condition that

An earlier onset of the symptom or

An expansion to other body parts or

Paradoxical response to pharmacological treatment or

have shorter duration of the treatment effect than during early treatment days.

Risk factors for augmentation

High dosage of dopaminergic Rx and short acting dopamine agonist drugs



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Other Risks of Augmentation

- Increased risk of augmentation in:
 - low iron stores
 - Greater severity of RLS/WED symptoms prior to initiation of treatment
 - Possibly a family history of RLS or lack of neuropathy (Primary RLS)

PLMD Combination of the extension of big toe and partial flexion of ankle, knee and sometimes hip Lower extremities > upper extremities Typically, patient is unaware of movement • Arousal may precede, coincide with or follow the movement. PLMS is associated with hypertensive autonomic arousals

Periodic limb movement disorder

Accompany in

- RLS, RBD, OSA, nacrolepsy, MSA, DRD
- General medical conditions such as
 - Renal failure, anemia, CHF, peripheral neuropathy
- In patients treated with some medications such as
 - Tricyclic antidepressants, neuroleptics and serotonin reuptake inhibitors

Associated features

- Higher rates of mood disorders,
 - Anxiety, attention deficits, oppositional behaviors and
- parasomnias

Periodic limb movement disorder: treatments

• Treatment:

- Dopaminergic drugs:
 - Dopamine agonists esp. pramipexole and ropinirole
 - Levodopa
- Gabapentin
- Clonazepam

Sleep related leg cramps

Differential diagnosis:

RLS:

- Leg discomfort during the sleep and sometimes complain of cramping sensation in RLS
- Actual spasm and hardening of muscle: critical differentiating factor
- Much briefer in leg cramp (a few second to sometimes several minutes) than RLS (persist for hours)

Dystonia:

- Electrophysiological study: on-going co-contraction of agonist and antagonist muscles.
- Leg cramp can be relieved by stretching, while dystonia can not.
- Treatment:
 - Local massage, stretching and movement of the limbs

Sleep related bruxism

Drugs that may induce bruxism (awake or sleep)

- Anti dopaminergic drugs
- Selective serotonin reuptake inhibitors
- Calcium antagonists
- Alcohol, caffeine or cigarettes
- Cocaine
- Amphetamine
 - 3% in elderly

Clinical subtypes

Primary, idiopathic: no medical/ dental causes

Secondary: associated with other disorders

- Children with cerebral palsy, mental retardation, ADHD
- Adults with abnormal movements such as facio-mandibular
 myoclonus, Parkinson's disease, RBD, tardive dyskinesia, dementia,
 and sleep related breathing disorders esp. OSA

latrogenic: Treatment-induced

Sleep related bruxism

- Pathophysiology: Multifactorial
 - Physiologic central nervous system (oromotor, sleep-wake regulation, catecholaminergic)
 - Autonomic nervous system
 - Psychosocial factors
 - Genetic factors
- Non-REM stage 1,2 (80%) but can occur in all stage

Sleep related bruxism

• Management: no specific cure exists

Behavioral strategies

- Psychological or physiological relaxation: sleep hygiene
- Biofeedback techniques

Dental strategies

- Occlusal appliances

Pharmacological strategies

- Benzodiazepine
- Muscle relaxants
- Dopaminergic drugs
- Botulinum toxin injection
- Avoid Selective Serotonin Reuptake Inhibitor (SSRI): increase sleep related bruxism

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Others

- Benign sleep myoclonus of infancy
- Propriospinal myoclonus at sleep onset
- Sleep related movement disorder due to a medical disorder
- Sleep related movement disorder due to a medication or substance
- Sleep related movement disorder, unspecified



RBD | Recurrent isolated sleep paralysis

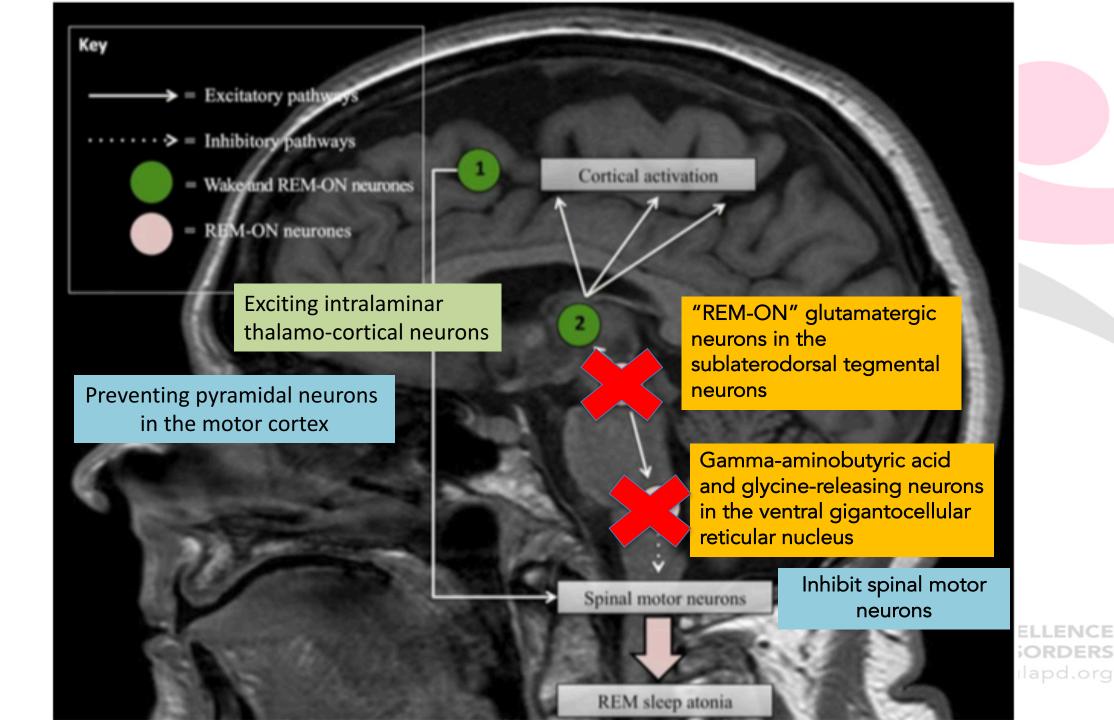
Nightmare disorders

Overlap parasomnia

Status dissociatus

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Movements in RBD

- Short and abrupt,
- Sometimes jerky, and
- Mostly related to the extremities.
- Complex behaviors with violent actions (punching or kicking) or bed falls may occur.
- Dream content is usually unpleasant and may involve the individual being attacked, chased, or threatened (by a person or animal)

RBD characteristics

REVIEW

REM Sleep Behavior Disorder: Motor Manifestations and Pathophysiology

Isabelle Amulf, MD, PhD*

Sleep disorders unit, Pitié-Salpêtrière Hospital, Pierre and Marie Curie University, Inserm U975, CRICM, Paris, France

- Behaviors during RBD are complex and varied.
 - Gesturing, reaching, grabbing, arm flailing, slapping,
 - Punching, kicking, sitting up, and leaping from bed
 - The forceful and violent aspect of these motor behaviors,
 - Often distal than proximal
- Usually associated with vivid, unpleasant, and active dreams
- Fighting or fleeing in response to danger (91%),
- Nonviolent elaborate behaviors during RBD, which were found in 18% of patients with PD
- In contrast to sleepwalking, only a minority of patients with RBD (3%)
- Occasionally stand up and walk and run, and most patients have the eyes closed

"Acting out" of dream content

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Vocalization in RBD

- Mumbling,
- Talking,
- Shouting,
- Swearing profanities,
- Laughing, and
- Crying
- Modulate their voices according to the dream context

Dreams and sleep behaviours

- Eye usually closed
- No major differences in dream content were found between patients with PD who did or did not have RBD
- Dream recall is quite variable, individuals are usually easily awakened and often remember their dreams, especially if questioned immediately after the behaviour.
- The dreams reported tend to have *negative emotional* content, or are often action-packed and aggressive but they can also be pleasant.
- Falling out of bed is frequent, but deliberately leaving the bed is rare.

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RBD1Q Questionnaire



NIH Public Access

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A Single-Question Screen for REM Sleep Behavior Disorder: A Multicenter Validation Study

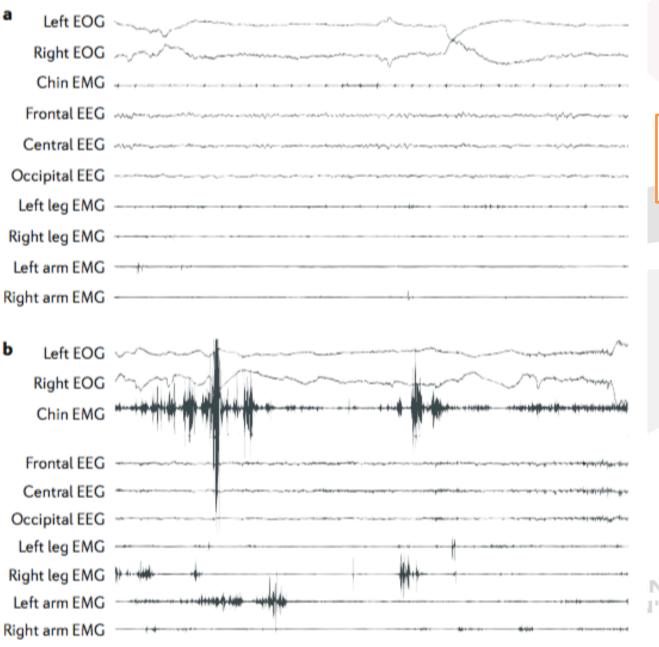
Yes- No Question

"Have you ever been told, or suspected yourself, that you seem to 'act out your dreams' while asleep (for example, punching, flailing your arms in the air, making running movements, etc.)?"

A sensitivity of 93.8% and a specificity of 87.2% chulapd.org

Other RBD questionnaires

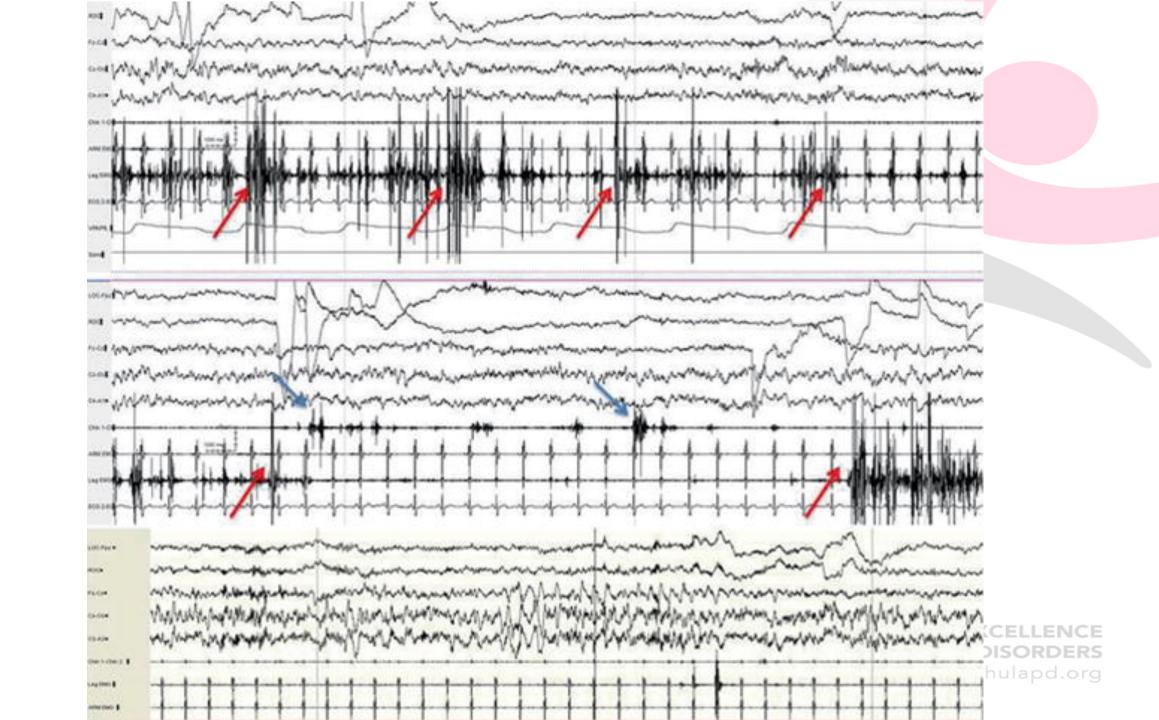
- REM behavior disorder screening questionnaire (RBDSQ):
 - 10- item questionnaire with scores ranging from 0 to 13.
 - Sensitivity of English version in PD for >/=6 cutoff 68-90%, specificity 63-82.8%
- REM sleep behavior disorder questionnaire Hong Kong (RBDQ-HK):
 - 13 item questionnaire.
 - Sensitivity 82.2%, specificity 86.9% mixed (idiopathic and secondary)
 RBD population
- Mayo sleep questionnaire (MSQ) item 1:
 - Single question.
 - Sensitivity in PD 90.3%, specificity 87.9% among PD patients meeting ICSDII criteria
- Innsbruck REM sleep behavior disorder inventory:
 - five item questionnaire.
 - In mixed (idiopathic and secondary) RBD population, sensitivity 91.4%,
 specificity 85.7%. For single RBD summary question, sensitivity 74.3%.
 specificity 92.9%.



Normal control REM sleep

RBD
EMG activity at chin
TA
FDS

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EMG activity during REM sleep

Phasic activity: a short burst of electromyography (EMG) activity lasting 0.1–5.0 s that is more than twice as high as the background EMG amplitude. Can be measured in 3 s mini-epochs or 30 s epochs.

Tonic activity: EMG activity increased by at least a factor of two or four compared with baseline in more than half of the epoch. Can be measured in 3 s mini-epochs or 30 s epochs.

Any activity: either phasic or tonic EMG activity. In addition, tonic and phasic muscle activity lasting between 5 and 15 s can be scored and can be measured in 3 s mini-epochs or 30 s epochs.

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Differential diagnosis RBD-like feature

- History of dream enactment behavior and daytime sleepiness with no evidence of REM sleep without atonia on PSG → OSA
 - Treatment with CPAP → eliminate the behavior (possible increased pressure for REM atonia prior REM sleep fragmentation)
- NFLE
- PLMS
- NREM parasomnias (such as sleepwalking, sleep terrors and confusional arousals, nightmares and sleep-related seizures (mainly sleep-related hypermotor epilepsy)

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NREM PARASOMNIA

Disorders of arousal (DOAs)

Confusional arousals

Sleep walking

Sleep terrors

sleep-related eating disorder (SRED)

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Confusional arousal

- Onset: peak onset before 5 years
- Abrupt onset out of slow wave sleep in the first third of the night, but may also occur out of stage N2
- Appears confused with vacant look and may have automatic and inappropriate behavior (few seconds to minutes), behavior may be inappropriate& violent but no autonomic hyperactivity or signs of fear like STs
- Multiple confusional episodes in the same night: uncommon
- Part of spectrum of sleep walking and sleep terror: all may occur in the same individual

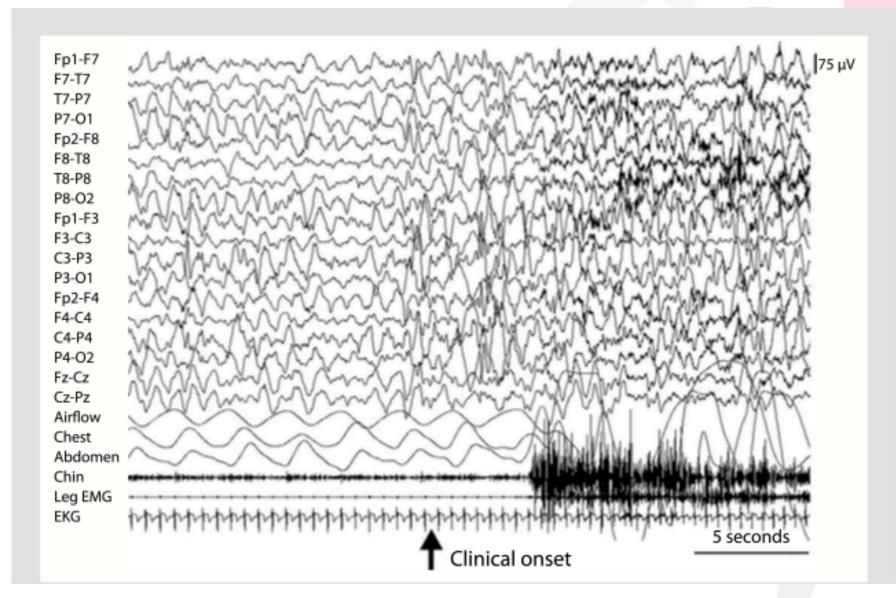
Sleep walking (Somnambulism)

- Most common in children between 5-12 years of age.
 - Sometimes it persists in adulthood, rarely begins in adult (1/3).
- Simply sitting up in bed, picking at the covers to walking, eye open but clumsy, more complex & violent& longer in adult.
- Episodes last less than 10 minutes, or terminated by returning to bed or simply lying down and continuing sleep.
- Injury, violent actions but they can negotiate their way/ may have sleep-related sexual behavior or violence.
- Family history: positive
- Precipitating factors:
 - sleep deprivation, fatigue, sedatives, concurrent illness

Sleep terror (Pavor Nocturnus)

- Peak onset is between 5 and 7 years of age
 - Sometimes it persists into adulthood
- Non-REM stage (slow wave sleep)
- The spell begins with abrupt set of intense autonomic and motor symptoms
 - A loud piercing scream
 - Confuse and fearful/ diaphoretic/ tachycardia
 - Usually sits up in the bed but can not communicate
- Many pts have history of sleep walking
- Precipitating factors: same as sleep walking

Disorders of Arousal From Non-REM Sleep



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NREM PARASOMNIA

	Confusional arousal	Sleep walking	Sleep terrors
Sleep stage at onset	N3	N3	N3
Autonomic hyperactivity	No	No	Prominent
Loud scream	No	No	Yes
Ambulation out of bed	No	Yes	No
Confusion during episode	Yes	Yes	Yes
Amnesia (partial/complete)	Yes	Yes	Yes

PSG is not required in NREM parasomnia

Sleep-related movement disorders VS Nocturnal epilepsy

Feature	NFLE	Arousal disorders	RBD
Age at onset	Variable, typically 1 st -2 nd decade	Usually 1 st decade	Over 50 years
Sleep stage of origin	NREM (N1,2, sleep wake transition)	NREM (N3)	REM
Timing of episode	Anytime	First third of sleep period	Last third of sleep period
Duration of episode	5-60 seconds	2-30 minutes	Seconds to 2 minutes
Frequency of episode	Nightly clusters	Sporadic, rare cluster	Sporadic, rare cluster
Onset and offset	Sudden	Gradual	Sudden
Seminology	Highly stereotyped, hypermotor, asymmetric tonic/dystonic	Not stereotyped, variable complexity	Not highly stereotyped, vocalizations, self-protective behaviors, dream recall
Level of consciousness during episode	Usually preserved	Variable	Poorly responsive
Postictal confusion	Typically absent	Present	Absent
Risk of injury	Low	High	Moderate
PSG with EEG	Epileptic activity < 50%	Slow-wave sleep arousals, rhythmic delta pattern	REM sleep without atonia

Sleep-related Dissociative Disorders

- Emerge at the transition from wake to sleep or shortly following awakening with EEG evidence of wakefulness
- Psychiatric comorbidities
 - Mood disorders, post- traumatic stress disorder, and a history of sexual abuse
- Episodes are non- stereotyped and feature screaming, running, and self-mutilating, violent behaviors that may represent a reenactment of prior traumatic events.
- Can last from minutes to an hour or longer, waxing & waning
- 70% developed between 2nd and 4th decades
- Injury & female → common
- Psychogenic non-epileptic seizures (PNES)
 - Arise from awake
 - Seizure overlap (pseudosleep)

Sleep-related Dissociative Disorders

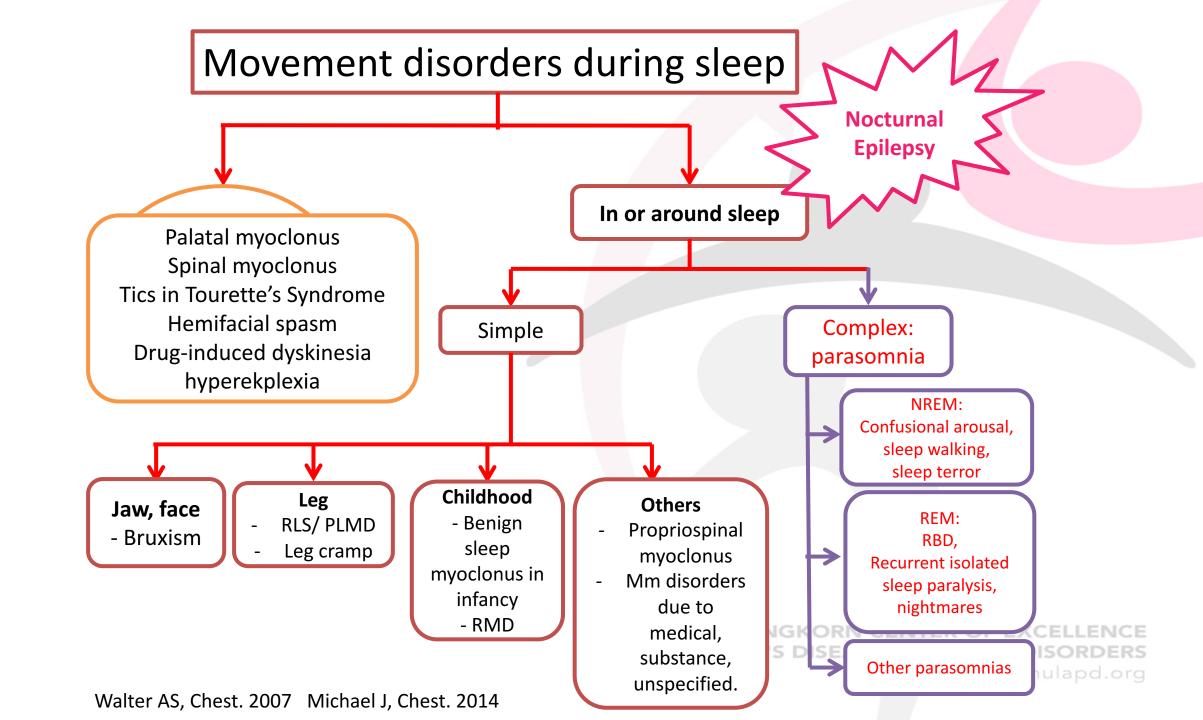
Motor manifestations

- Jactitation (restless tossing in bed),
- Asynchronous movements,
- Side- to-side head movements,
- Pelvic thrusting,
- Opisthotonic posturing,
- Prolonged body flaccidity,
- Preserved awareness during bilateral motor activity.
- Ictal eye closure and jaw clenching

Affective manifestations

- Vocalizations,
- Ictal moaning and crying,
- Emotive speech,
- Ictal stuttering,
- Heart rate elevations
- Postictal crying

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Thank you for your attention