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## Nonepileptic Seizures: An Updated Review

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### Abstract

Psychogenic nonepileptic seizures are a Functional Neurological Disorder/ Conversion Disorder subtype, which are neurobehavioral conditions at the interface of Neurology and Psychiatry. Significant advancements over the past decade have been made in the diagnosis, management and neurobiological understanding of PNES. This article reviews published PNES research focusing on semiologic features that distinguish PNES from epileptic seizures, consensus diagnostic criteria, the intersection of PNES and other comorbidities, neurobiological studies, evidence-based treatment interventions and outcome studies. Epidemiology and health care utilization studies highlight a continued unmet medical need in the comprehensive care of PNES. Consensus guidelines for diagnostic certainty are based on clinical history, semiology of witnessed typical event(s), and EEG findings. While certain semiologic features may aid the diagnosis of PNES, the gold standard remains capturing a typical event on video electroencephalography (EEG) showing the absence of epileptiform activity with history and semiology consistent with PNES. Medical-neurologic and psychiatric comorbidities are prevalent in PNES and should be assessed in diagnostic evaluations, and integrated into treatment interventions and prognostic considerations. Several studies, including a pilot multicenter, randomized clinical trial, have now demonstrated that a cognitive behavioral therapy informed psychotherapy is an efficacious treatment for PNES, and additional efforts are necessary to evaluate the utility of pharmacologic and other psychotherapy treatments. Neuroimaging studies, while requiring replication, suggest that PNES may occur in the context of alterations within and across sensorimotor, emotion regulation/

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processing, cognitive control and multimodal integration brain systems. Future research could investigate similarities and differences between PNES and other somatic symptom disorders.

### Keywords

Psychogenic Nonepileptic Seizures; Conversion Disorder; Cognitive Behavioral Therapy; Sertraline; Functional Neuroimaging; fMRI; Functional Neurological Disorder; Psychotherapy

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## INTRODUCTION

Functional Neurological Disorders (FND) / Conversion Disorders (CD) are highly prevalent neurobehavioral conditions at the interface of Neurology and Psychiatry. Psychogenic Nonepileptic Seizures (PNES) are a FND/CD subtype where individuals exhibit paroxysmal convulsive events and/or alterations in behavior and consciousness that resemble epileptic seizures (ES) but are not associated with changes in cortical activity<sup>1</sup>. PNES and other somatic symptoms were first introduced in the medical literature as “hysteria” and considered identifiable neurological conditions by Jean-Martin Charcot; early psychological theories were postulated by Sigmund Freud, Pierre Janet and others<sup>2</sup>. Modern day conceptualizations of PNES now integrate mind and brain<sup>3,4</sup>.

Over the past decade notable advances have been made in our ability to diagnosis and treat patients with PNES. Despite these achievements, epidemiology and health care utilization studies highlight a continued unmet medical need in the comprehensive interdisciplinary care of PNES. Advances in clinical neuroscience and systems-level neurobiological studies have started to elucidate a neurobiology for PNES, which offers promise in clarifying the underlying pathophysiology and identifying biomarkers guiding prognosis and treatment selection. In this article, we review recently published research in adults with PNES focusing on semiologic features that distinguish PNES from ES, newly published diagnostic criteria, the intersection of PNES and other psychiatric and neurological comorbidities, neurobiological studies, evidence-based treatment interventions, and outcome studies.

## THE SCOPE OF THE UNMET MEDICAL NEED

PNES occur worldwide, and in the United States up to 20% of civilians and up to 25% of Veterans diagnosed as having epilepsy actually have PNES<sup>5</sup>, making PNES as common as multiple sclerosis and Parkinson disease<sup>6</sup>. Comprehensive epidemiological investigations are lacking in PNES, however, the estimated prevalence of PNES is up to 33/100,000 individuals<sup>7</sup>. The incidence rate from a prospective study of 367,566 individuals followed over 3 years found that new-onset PNES was observed in 4.9/100,000/year<sup>8</sup>. As is typical for PNES, individuals were 80% women with a mean age of onset of 31 ( $\pm 15$ ) years. The majority of PNES individuals were unemployed and 46% were previously diagnosed with anxiety or depression; 57% of patients also had at least one additional medically unexplained symptom (MUS), highlighting the overlap between PNES and other FNDs. The coincidence of PNES with ES is approximately 10%<sup>9</sup>.

The economic and health care utilization costs incurred in caring for PNES are substantial. As one illustrative example, 15 of 28 PNES subjects accounted for 34 emergency room visits, and 12 of 28 individuals accounted for 66 inpatient hospital days pre-diagnosis<sup>10</sup>. In a retrospective study from Ireland, the annual estimated cost per patient with undiagnosed PNES was 20,995 euros<sup>11</sup>. While health care expenditures decrease substantially post-diagnosis<sup>10,11</sup>, multi-year delays in diagnosis following symptom onset, continued PNES events, unemployment, and medical disability perpetuate health care and economic costs.

Further compounding the problem, many health care providers report frustration when caring for patients with PNES. In a semi-structured interview of 75 health-care clinicians and workers in the Veterans Health Administration system, while some providers endorsed sentiments of hope, many expressed frustration over the complexity of PNES which included difficulties with patient acceptance of the diagnosis, uncertainties in treatment interventions, and a failure of cross disciplinary collaborations<sup>12</sup>. While neurologists increasingly recognize the scope of the unmet medical need, many remain uncomfortable discussing somatoform symptoms with patients<sup>13</sup>; psychiatrists and other mental health care providers may also potentially feel ill-equipped to treat individuals with PNES.

## **SEMILOGIC DIFFERENTIATION OF PNES FROM ES AND DIAGNOSTIC CRITERIA**

Differentiating PNES from ES can be clinically challenging, and several studies have investigated clinical signs that aid the diagnosis of PNES<sup>1,14</sup>. A retrospective review detailing signs that reliably distinguished PNES from ES suggested that a diagnosis of PNES was favored for events showing a long duration, a fluctuating course, asynchronous or side-to-side movements, ictal eye closure at onset, ictal crying and post-ictal recall of information when presented ictally (See Table 1). In addition, urinary incontinence and tongue biting do not reliably distinguish between ES and PNES<sup>15</sup>. A prospective study of 120 seizures in 35 consecutive subjects showed that video-documented preserved awareness, eye fluttering, and the modulation of event intensity by bystanders reliably predicted PNES; abrupt onset, ictal eye-opening and post-ictal confusion/sleep reliably predicted ES<sup>16</sup>. It is also worth noting that apart from differentiating PNES from ES, additional diagnostic considerations should also be evaluated and ruled out including paroxysmal movement disorders, panic attacks and physiologic forms of non-epileptic events including cardiac arrhythmias among other conditions.

While semiological features may suggest a diagnosis of PNES over ES, the gold standard for diagnosis is the recording of typical event(s) on video electroencephalography (vEEG) and noting a lack of epileptiform activity in the peri-ictal period with semology and history consistent with PNES. While this is the diagnostic work-up of choice, access to vEEG may not be available, and some patients may have sufficiently low event frequencies that hospital admission for long term monitoring is impractical. Recently, the International League Against Epilepsy (ILAE) Commission on Neuropsychobiology Nonepileptic Seizures Task Force published consensus guidelines on the minimum requirements to diagnosis PNES (See Table 2)<sup>17</sup>. Levels of diagnostic certainty are based on the clinical history, ictal semiology of

an event, and EEG findings. To meet criteria for *documented PNES*, the clinical history should favor PNES and habitual events must be recorded on vEEG demonstrating the absence of epileptiform activity. Lower levels of certainty (*clinically established, probable, possible*) are based on availability of diagnostic components.

Given the clinical need to clarify a diagnosis of PNES vs. ES, whenever possible, seizure induction protocols have drawn increasing interest. While ethical concerns have been raised regarding provocation techniques given that they may be perceived as misleading, studies suggest that use of conventional EEG activation procedures, including hyperventilation and photic stimulation, may be effective adjunctive techniques in the diagnosis of PNES<sup>18</sup>, making provocation unnecessary. Providing correct and explicit information about the potential utility of standard induction protocols may alleviate ethical concerns and aid diagnostic evaluations. Surface electromyography recordings may also potentially aid differentiation of convulsive ES and PNES events<sup>19</sup>. Approximately twice normal serum prolactin levels drawn 10-20 minutes following an ictal event can also help differentiate convulsive ES from PNES<sup>17</sup>.

## MEDICAL, NEUROLOGICAL and PSYCHIATRIC COMORBIDITIES

Apart from their seizures, adult patients with PNES have co-morbid medical, neurologic and psychiatric conditions that contribute to their overall symptom complex, prognosis and treatment responses. For example, in a two-year retrospective review of PNES (N=158) and ES (N=122), individuals with PNES were more likely to report a history of other medical somatic syndromes (e.g. fibromyalgia, chronic fatigue syndrome, chronic pain, irritable bowel syndrome) and more frequently endorsed chronic, intermittent medical conditions (e.g. migraines, asthma)<sup>20</sup>. Compared to ES, patients with PNES also more commonly endorsed complaints on a review-of-systems questionnaire<sup>21</sup>. Veteran<sup>22</sup> and civilian<sup>23</sup> populations with PNES frequently report a history of traumatic brain injury (TBI), and a distinct subset of patients have co-morbid intellectual disabilities<sup>9</sup>.

Categorical psychiatric diagnoses and symptom-specific increases in anxiety, depression, and dissociation are linked to PNES. Patients with PNES often have mood, anxiety, dissociative and other somatic symptom disorders<sup>3</sup>, along with personality disorders, including clusters B and C personality disorders<sup>24</sup>.

Post Traumatic Stress Disorder (PTSD) is also associated with PNES. Estimates suggest that three-fourths of adults with PNES report prior traumatic experiences, including sexual abuse (~30%) and physical abuse (~25%)<sup>9,25</sup>. Individuals with PNES and prior sexual abuse exhibit an earlier event onset, greater diagnostic delay, more severe convulsions, emotional triggers and traumatic recollections among other symptoms<sup>26</sup>. In a study comparing PNES individuals with and without trauma, traumatized individuals were more likely to have co-morbid psychiatric conditions and dissociative symptoms<sup>27</sup>. Antecedent trauma has also been linked to other MUS in PNES<sup>9</sup>. Particular associations between PTSD and PNES have been identified in combat Veterans. In a retrospective review of 50 Veterans with PNES and 37 veterans with ES, PTSD symptoms preceded the diagnosis of PNES in 58%, and PTSD

comorbidity differentiated PNES and ES<sup>5</sup>. Veterans with PNES and combat related PTSD may exhibit more hypomotor or nonmotor PNES events<sup>28</sup>.

From a dimensional, symptom-specific perspective, patients with PNES exhibit abnormal neuropsychiatric and personality profiles. Associations between PNES and depression have been well established, although depression scores on psychometric measures do not reliably differentiate between PNES and ES<sup>29</sup>. Patients with PNES report dissociative symptoms (fragmentation of one's internal experience of the outside world (derealization) and/or fragmentation and compartmentalization of one's self-perception and body-schema (depersonalization)), and dissociation in PNES has been linked to depression, somatic symptoms and an external locus of control<sup>30</sup>. Alexithymia, the reduced ability to recognize and express emotions, has also been observed in PNES. It remains unclear, however, if alexithymia scores differentiate PNES and ES<sup>31</sup>. Beliefs about emotions are impaired in PNES compared to healthy subjects, with one study reporting that individuals with PNES described emotions as overwhelming, uncontrollable, shameful, irrational and damaging<sup>32</sup>. Cluster analyses suggest possible psychopathologic PNES subtypes; in a cohort of 43 PNES individuals, 11 were categorized as having high levels of psychopathology, somatization, alexithymia and impaired emotion regulation, while 32 subjects reported high somatization and depression scores but relatively normal emotion expression and regulation abilities<sup>33</sup>.

Personality measures, including the Personality Assessment Inventory (PAI) and the Minnesota Multiphasic Personality Inventory 2 (MMPI-2), have been widely used in PNES. In a large cohort of PNES (N=75) and ES (N=109) subjects, PNES patients endorsed significantly higher somatic, conversion, depressed, anxious and suicidal symptoms on the PAI<sup>34</sup>. Elevated conversion subscale (SOM-C) scores predict PNES vs. ES<sup>35</sup>. On the MMPI-2, PNES compared to ES show a "conversion V" pattern on personality testing, which consists of elevations in hypochondriasis and hysteria subscales relative to the depression subscale<sup>36</sup>. Predictive regression analyses suggest that dual use of the conversion and health concern subscales of the PAI and the hysteria subscale of the MMPI-2 may also aid PNES vs. ES distinctions<sup>37</sup>.

Neuropsychological studies have started to quantify cognitive profiles in PNES, however, they do not differentiate individuals with PNES from ES. Individuals with PNES frequently endorse cognitive symptoms including memory and concentration difficulties. It has been reported that PNES patients potentially overestimate their cognitive deficits; one study of PNES vs. ES showed that PNES patients endorsed greater word finding difficulties but performed better than ES on the Boston Naming Test<sup>38</sup>. Despite this finding, when controlling for effort and IQ, patients with PNES show some impairments in spatial working memory, attention and executive functions<sup>39</sup>. Several, particularly informative studies have probed the intersection of cognitive and emotional functions in PNES<sup>40-42</sup>. In a study of 20 subjects with PNES vs. healthy controls, those with PNES exhibited working memory deficits to an N-back task paired with facial distractors; following stress induction via the Cold Pressor Test, working memory performance exhibited more generalized impairments<sup>40</sup>. In this cohort, increases in salivary cortisol were associated with larger stress induced working memory impairments. In a study using an affectively valenced facial viewing task to probe aspects of set shifting, PNES compared to healthy subjects showed greater

difficulty switching attention from emotion-related task demands<sup>42</sup>. One study comparing PNES individuals with PTSD, individuals with PNES and prior trauma without PTSD, and PNES subjects without previously experienced trauma showed that PNES subjects with PTSD exhibited lower episodic verbal memory performance than comparison groups<sup>41</sup>. Additional research is necessary to clarify the relationship between neurocognitive deficits and mood and anxiety symptoms in PNES, particularly whether neurocognitive deficits occur independent of mood symptoms.

## THE EMERGING NEUROBIOLOGY OF PNES

Systems-level functional and structural neuroimaging studies have started to elucidate the neurobiology of PNES<sup>3</sup>. Several functional magnetic resonance imaging (fMRI)<sup>43-47</sup> and one positron emission tomography (PET) study used resting state techniques to investigate neural circuit disturbances in PNES. In a region-of-interest “seed” based analysis, increased functional connectivity was observed between motor regions (precentral sulcus) and regions involved in emotional processing (anterior cingulate cortex (ACC), insula) and executive functions (inferior frontal gyrus, parietal cortex); among other findings, trait dissociation scores positively correlated with the functional connectivity strength between the precentral sulcus and the posterior insula<sup>43</sup>. Several studies used data-driven, multivariate functional connectivity analyses to demonstrate widespread functional connectivity alterations in emotion control, executive, fronto-parietal (attentional), sensorimotor and default mode networks<sup>44-46</sup>. Another study specifically evaluated functional connectivity patterns across insular subregions, observing that PNES compared to healthy subjects showed increased functional connectivity between the left ventral anterior insula and the left post-central gyrus and bilateral supplementary motor area (SMA); functional connectivity strength within the bilateral SMA positively correlated with PNES event frequency<sup>47</sup>. Functional connectivity strength between the SMA and ACC has also been shown to positively correlate with PNES frequency<sup>48</sup>. In comparison to healthy subjects, a 2-deoxy-2-[fluorine-18]fluoro-D-glucose PET study showed that PNES patients exhibited bilateral ACC and right inferior parietal lobule hypometabolism<sup>49</sup>.

Quantitative structural MRI have also been conducted in PNES<sup>50-52</sup>. Compared to matched controls, PNES patients showed ACC, SMA and pre and post central gyrus atrophy on cortical thickness and voxel-based-morphometry analyses<sup>50</sup>; bilateral precentral gyrus cortical thickness reductions were independently replicated along with observed increases in insular and orbitofrontal cortical thickness in PNES<sup>52</sup>. A diffusion tensor tractography study showed a right-ward asymmetry of the uncinate fasciculus (a tract connecting medial prefrontal and medial temporal regions) in PNES compared to healthy controls<sup>51</sup>. While functional and structural neuroimaging studies require further replication and control of potential confounding variables, these early results suggest that PNES may manifest in the context of alterations within and abnormal interactions across sensorimotor, emotion regulation/processing, cognitive control and multimodal integration neural circuits (See Figure 1 for plots of peak coordinates in the above discussed functional and structural neuroimaging studies). Research is needed to evaluate the specificity of PNES related neural circuit alternations in relation to neurologic (i.e. epilepsy, TBI) and psychiatric (PTSD, depression, other functional neurological disorders) control groups. Additional research



efforts are also needed to investigate if semiologic differences in PNES (i.e. convulsive vs. atonic events) are linked to distinct patterns of neural circuit alterations.

Electrophysiology and autonomic nervous system studies have also contributed to the understanding of PNES pathophysiology. In a cohort of 18 PNES and 18 healthy subjects, resting state high-density source EEG analyses showed that PNES patients exhibited decreased functional connectivity between the basal ganglia and cortical regions, along with reduced interhemispheric connectivity across paralimbic regions<sup>53</sup>. In a unique case of intracranial recordings during PNES, decreased power in the theta band was observed over the posterior parietal cortex<sup>54</sup>. Several investigations have probed autonomic profiles peri-ictally in PNES<sup>55</sup>; some noted increased sympathetic tone ictally in ES vs. PNES<sup>56</sup>, while others characterized pre-ictal and post-ictal autonomic changes as distinguishing between groups<sup>57</sup>.

A few studies have investigated serologic biomarkers in PNES. Positive correlations were observed between baseline salivary cortisol levels and negative attentional bias during performance of a masked emotional Stroop task<sup>58</sup>; in this same cohort, negative attentional bias also correlated with past sexual trauma<sup>59</sup>. Increases in basal diurnal cortisol levels in PNES compared to healthy controls were mediated mainly by patients reporting past sexual trauma<sup>60</sup>. Reductions in serum brain derived neurotrophic factor, implicated in synaptic reorganization and neurogenesis, have also been reduced in samples with PNES and with ES, compared to healthy controls<sup>61</sup>.

## TREATMENT AND OUTCOME STUDIES

In the past decade, significant progress has been made in the evidence-based management of PNES<sup>62</sup> (See Table 3). In a pilot randomized controlled trial of 12-week, cognitive-behavioral therapy (CBT) plus standard medical care (SMC) vs. SMC alone, conducted in 66 patients with PNES, intention to treat analyses showed a statistically significant reduction in seizure frequency at the end of the treatment period associated with CBT<sup>63</sup>. At a 3-month follow-up, however, mean seizure frequency between the treatment arms did not remain statistically significant. A pilot, double-blind, randomized, placebo-controlled trial of flexible-dose sertraline conducted over 12 weeks, showed that PNES subjects assigned to the sertraline treatment displayed a 45% reduction in seizure frequency as compared to an 8% seizure frequency increase to placebo<sup>64</sup>. Recently a multicenter, randomized clinical trial was conducted in 38 patients with PNES comparing four treatment arms: flexible-dose sertraline only, CBT-informed psychotherapy (CBT-ip) only, CBT-ip with flexible-dose sertraline and SMC<sup>65</sup>. Within-group analyses for each group showed that the two psychotherapy treatment arms exhibited a significant reduction in seizures (51% in the CBT-ip only arm and 59% in the CBT-ip plus sertraline arm), while the sertraline and the SMC did not show a significant reduction in seizures. The CBT-ip only group also showed significant improvements in depression, anxiety, quality of life and global functioning. These findings support the use of a manualized CBT-ip PNES Workbook<sup>66</sup>. The intervention is being disseminated for treatment by a range of clinicians, from epileptologists/neurologists, psychiatrists, psychologists, and social workers<sup>67</sup>.

A randomized controlled trial of a post-PNES diagnosis group psychoeducation program (3 successive monthly, 1.5 hour group sessions) also showed that patients in the psychoeducation intervention reported improvements in perceived psychosocial functioning and a trend towards decreased seizure-related emergency room visits or hospitalizations, however, seizure frequency reduction was not observed<sup>68</sup>. Lastly, a randomized controlled trial of immediate vs. delayed withdrawal of antiepileptic drugs (AED) in PNES showed that immediate AED withdrawal was associated with a reduction in seizures, decrease use of rescue medications, and greater perceived internal locus of control<sup>69</sup>. The long-term benefits of therapy and other treatments can be examined in randomized, controlled clinical trials along with other potential pharmacologic and psychotherapy treatment interventions<sup>70,71</sup>.

Outcome and predictor of outcome studies have been performed in PNES. Given that the initial stages of treatment include accurate diagnosis and effective communication of the diagnosis, it is noteworthy that diagnostic acceptance by patients improves outcomes<sup>72</sup>. Effective communication alone, however, is insufficient to improve outcomes in most patients<sup>73</sup>. In a retrospective study of 260 consecutively evaluated PNES patients measuring outcome at 6 and 12 months post diagnosis, 18% showed an increase in seizures, 38% were event free and the majority of patients continued to have seizures<sup>74</sup>. Women, individuals with anxiety/depression, and patients receiving social security benefits were less likely to become event-free. Post diagnosis, many also developed other MUS functional neurological symptoms<sup>75</sup>. In addition to psychosocial and psychiatric variables impacting outcome, patients with PNES with a history of TBI exhibit worse outcomes, as exemplified by increased likelihood of having major depression, behavioral impulsivity, PTSD, receiving disability and lower global functioning<sup>23</sup>.

## CONCLUSIONS

PNES are highly prevalent, complex neuropsychiatric symptoms. Over the past 5-10 years, notable advancements have been made in the diagnosis and management of this previously enigmatic condition. Neurobiological studies using systems-level, clinical neuroscience techniques are elucidating an emergent neurobiology for PNES, which will hopefully help reduce the stigma associated with this condition and facilitate additional research efforts to identify diagnostic biomarkers of prognosis and treatment response. Future investigations will also help clarify gender differences, and similarities and differences between PNES and other somatic symptom subtypes.

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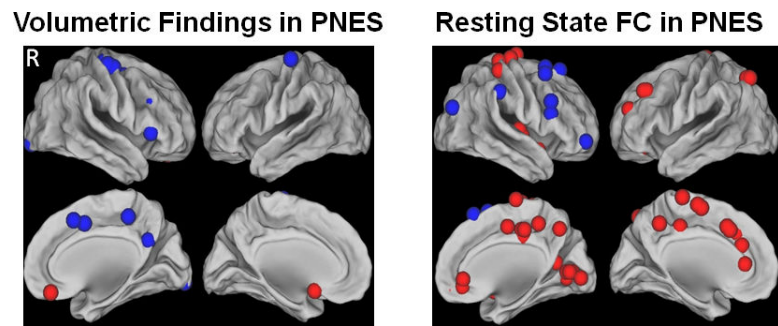


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**Figure 1. Display of volumetric and resting-state functional connectivity (FC) findings in Psychogenic Nonepileptic Seizures (PNES) neuroimaging studies**

Left panel displays peak coordinates of cortical thickness and voxel-based morphometry findings in PNES vs. healthy controls<sup>50,52</sup>. Right panel shows peak coordinates of abnormal resting-state FC in individuals with PNES vs. healthy controls<sup>43,45,47,48</sup>. Blue circles indicate decrease and red circles indicate increase. Evidence from neuroimaging studies, while requiring replication and further investigations, suggests that PNES may potentially develop in the context of alterations within and across brain networks implicated in sensorimotor (e.g. pre- and post-central gyrus, premotor regions), emotion processing/regulation (e.g. anterior cingulate cortex (ACC), middle cingulate cortex (MCC), orbitofrontal cortex, insula), cognitive control (e.g. lateral prefrontal cortex, dorsal ACC, MCC), and multimodal integration functions (e.g. cingulate gyrus, posterior parietal cortex, precuneus). Only main findings are displayed using Caret 5. Cerebellar and basal ganglia foci, and seed region coordinates are not shown.

**Table 1**

Semiologic features that support the diagnosis of Psychogenic Nonepileptic Seizures (PNES) vs. Epileptic Seizures (ES).

<b>Signs Favoring PNES</b>	<b>Signs Favoring ES</b>	<b>Indeterminate Signs</b>
Long Duration	Occurrence From Physiologic Sleep	Gradual Onset
Fluctuating Course	Postictal Confusion	Non-Stereotyped Events
Asynchronous Movements *	Stertorous Breathing	Flailing or Thrashing Movements
Pelvic Thrusting *		Opisthotonus
Side-To-Side Head or Body Movements **		Tongue Biting
Forced Eye Closure		Urinary Incontinence
Ictal Crying		
Memory Recall		

Adopted from Avbersek et al<sup>14</sup> with permission.

\* indicates that sign may not reliably differentiate between PNES and frontal lobe partial epileptic seizures.

\*\* indicates that sign may only be helpful in distinguishing convulsive PNES and ES.

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**Table 2**

Diagnostic levels of certainty for psychogenic nonepileptic seizures

Diagnostic level	History	Witnessed event	EEG
<b>Possible</b>	+	By witness or self-report/description	No epileptiform activity in routine or sleep-deprived interictal EEG
<b>Probable</b>	+	By clinician who reviewed video recording or in person, showing semiology typical or PNES	No epileptiform activity in routine or sleep-deprived interictal EEG
<b>Clinically established</b>	+	By clinician experienced in diagnosis of seizure disorders (on video or in person), showing semiology typical of PNES, while not on EEG	No epileptiform activity in routine EEG or ambulatory ictal EEG, capturing a typical ictus*
<b>Documented</b>	+	By clinician experienced in diagnosis of seizure disorders, showing semiology typical of PNES, while on video EEG	No epileptiform activity immediately before, during or after ictus captured on ictal video EEG with typical PNES semiology

Modified from LaFrance WC Jr, Baker GA, Duncan R, et al. Minimum requirements for the diagnosis of psychogenic nonepileptic seizures: a staged approach: a report from the International League Against Epilepsy Nonepileptic Seizures Task Force. *Epilepsia* 2013;54(1):2005-2018. With permission.

+ history characteristics consistent with PNES; PNES, psychogenic nonepileptic seizures; ES, epileptic seizures; EEG, electroencephalogram.

\* Captured ictus should not resemble types of ES which may not show ictal epileptiform correlate on EEG (e.g., simple partial epileptic seizures).

**Table 3**

Summary of controlled clinical trials in the treatment of Psychogenic Nonepileptic Seizures (PNES).

Study	Subjects	Interventions	Primary End Point	Significant Findings
Goldstein et al, 2010 <sup>63</sup>	66 PNES	CBT + SMC vs. SMC	Sz Frequency	↓ Sz at study end but not at f/up
LaFrance et al, 2010 <sup>64</sup>	38 PNES	Sertraline vs. Placebo	Sz Frequency	↓ Sz with Sertraline
Oto et al, 2010 <sup>69</sup>	25 PNES	Immediate vs. Delayed AED Withdrawal	Sz Frequency	↓ Sz with Immediate Withdrawal
Chen et al, 2013 <sup>68</sup>	64 PNES	Group Psychoeducation vs. SMC	Sz Frequency & Psychosocial Function	Improved Psychosocial Function
LaFrance et al, 2014 <sup>65</sup>	38 PNES	CBT-ip vs. CBT-ip + Sertraline vs. Sertraline vs. SMC	Sz Frequency	↓ Sz with CBT-ip CBT-ip + Sertraline

CBT, cognitive behavioral therapy; CBT-ip, cognitive behavioral therapy-informed psychotherapy; SMC, standard medical care; AED, anti-epileptic drug; Sz, seizures; f/up, follow-up; ↓ - reduction in.