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# The impact of narcolepsy symptoms and treatment on sex life -Current Evidence and Reports

Wpływ objawów i leczenia narkolepsji na życie seksualne - aktualne dowody i doniesienia naukowe

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## Abstract

**Introduction:** Narcolepsy (NT) can manifest as excessive daytime sleepiness (EDS), cataplexy, hypnagogic (HHG) and hypnopompic (HHP) hallucinations, sleep paralysis, orgasmolepsy – all of which may affect patients' functioning. The aim of the study was to determine the impact of NT, specific sexual dysfunctions and drugs used in NT on sex life.

**Materials and methods:** A narrative review was conducted, using keywords: narcolepsy, sexual dysfunctions, medications in narcolepsy from repository inception to March 17, 2024 searching PubMed/MEDLINE, Google Scholar, Crossref, Cochrane databases. The quality of the reviewed articles was assessed using the Scale for the Assessment of Narrative Review Articles (SANRA).

**Results:** Sexual activity and satisfaction are reduced in up to 81.1% of patients. There are often multiple co-occurring autonomic dysregulations in patients, including the genitourinary system, causing sexual dysfunction (erectile dysfunction in 48% of men; vaginal lubrication in 81% of women). Orgasmolepsy negatively affects sexual and social relationships in more than 1/3 of respondents. Sleep paralysis can be accompanied by HHG, which often present the impression of sexual assault and harassment. They may cause anxiety, post-traumatic stress disorder (PTSD) features, depressed mood, in extreme cases leading to suicide attempts. Of the drugs used in NT, only methylphenidate can increase libido and reduce erectile dysfunction in NT.

**Conclusions:** Orgasmolepsy, sexual and autonomic dysfunction in NT significantly reduce patients' quality of sex life. Sleep paralysis with sexual HHG can reduce psychological well-being. A holistic intervention approach, using behavioural interventions, cognitive-behavioural therapy, education of the patient and their loved ones, is crucial in the treatment of sexual difficulties.

Keywords: narcolepsy, sexual dysfunctions, medications in narcolepsy

## Streszczenie

**Wstęp:** Narkolepsja (NT) może objawiać się nadmierną sennością w ciągu dnia (excessive daytime sleepiness, EDS), katapleksją, halucynacjami hipnagogicznymi (hypnagogic hallucinations, HHG) i hipnopompicznymi (hypnopompic hallucinations, HHP), paraliżem przysennym, orgazmolepsją, które zaburzają funkcjonowanie pacjentów. Celem pracy jest określenie wpływu NT, współwystępujących specyficznych dysfunkcji seksualnych oraz leków stosowanych w terapii NT na życie seksualne.

**Materiały i metody:** Przeprowadzono przegląd narracyjny z użyciem słów kluczowych: narkolepsja, dysfunkcje seksualne, leczenie narkolepsji oraz deskryptorów czasowych do 17 marca 2024 r., przeszukując bazy PubMed/MEDLINE, Google Scholar, Crossref, Cochrane. W celu weryfikacji jakości przeglądanych artykułów posłużono się sześciopunktową skalą SANRA (Scale for the Assessment of Narrative Review Articles).

**Wyniki:** Wykazano, że aktywność i satysfakcja seksualna są obniżone nawet u 81,1% pacjentów. U chorych często współwystępują liczne zaburzenia regulacji autonomicznej, w tym układu moczowo-płciowego, powodując dysfunkcje

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seksualne (erekcji - 48% mężczyzn; nawilżenia pochwy - 81% kobiet). Orgazmolepsja negatywnie wpływa na seksualne i społeczne relacje u ponad ¼ badanych. Paraliżowi przysennemu często towarzyszą nieprzyjemne HHG, które przedstawiają wrażenie napaści i molestowania seksualnego. Mogą powodować lęk, cechy zespołu stresu pourazowego (post-traumatic stress disorder, PTSD), obniżony nastrój, a w skrajnych przypadkach prowadzić do prób samobójczych. Spośród leków stosowanych w NT jedynie metylofenidat może zwiększać libido i zmniejszać zaburzenia erekcji.

**Wnioski:** Orgazmolepsja, dysfunkcje seksualne i autonomiczne w NT znacząco obniżają jakość życia seksualnego pacjentów. Paraliż przysenny z HHG o charakterze seksualnym często negatywnie wpływają na dobrostan psychiczny. Holistyczne podejście z wykorzystaniem interwencji behawioralnych, terapii poznawczo-behawioralnej, edukacji pacjenta i jego bliskich są kluczowe w leczeniu zaburzeń seksualnych w przebiegu NT.

Słowa kluczowe: narkolepsja, dysfunkcje seksualne, leczenie narkolepsji

#### Introduction

Narcolepsy is a chronic rare disease that affects sleep architecture [1,2]. It is distinguished into two types, depending on the presence of cataplexy (Narcolepsy type 1; NT1) and lack of that symptom (Narcolepsy type 2; NT2). It is estimated that the frequency of NT2 is 20.5/10,000 and NT1 is 0.02% - 0.18% in the US and Western European populations and 0.16% - 0.18% in the Japanese populations [3]. Narcolepsy is manifested by excessive daytime sleepiness (EDS), hallucinations and sleep paralysis [4]. Cataplexy is a sudden loss of muscle tension during wakefulness caused by especially intense emotions for example anger, excitement, or those leading to laughter. In this state the patient is conscious, but unable to move. It can last from a few seconds to a few minutes [3,5]. EDS consists of daily periods of uncontrollable urge for sleep, which can lead to real sleep attacks [3]. Hallucinations are multimodal or complex visual sensory misperceptions referring to experiences where individuals perceive visual stimuli in a manner that involves multiple senses or complex sensory distortions [6]. Hallucinations in patients with cataplexy are hypnagogic hallucinations (HHG) and hypnopompic hallucinations (HHP) [3,7]. HGH are dream-like hallucinations that appear at the beginning of sleep and can be very terrifying because the patient is not aware of their hallucinatory nature. HPH occurs upon awakening [3]. Hallucinations can also appear during the day [8]. In some patients the dominance of very real hallucinations limits the possibility of recognizing them as unreal [8-10]. Sleep paralysis is often combined with hallucinations. Normal muscle inhibition or atonia occurs during partial wakefulness, either during the period of falling asleep (HHG) or less frequently, during the period of awakening (HHP). This condition is known as "sleep paralysis" [3,11]. These phenomena - cataplexy, hallucinations and sleep paralysis are categorised as REM dissociation phenomena, characterised by the occurrence of REM sleep dreams without concurrent REM phenomena

or the loss of muscle tone during wakefulness [4]. One of the first descriptions of narcolepsy with cataplexy originates from German authors Westphal in 1877 [12] and Fisher in 1878 [13]. Theories suggest that narcolepsy is caused by decreased levels of orexin (also known as hypocretin), a neuropeptide involved in regulating alertness [14,15]. In most cases normal concentrations of hypocretin in cerebrospinal fluid (CSF) appear to characterise narcolepsy without cataplexy and narcolepsy with cataplexy is correlated with low concentrations of hypocretin in CSF [14,16,17]. Damage to orexin producing neurons is more limited in NT2 than in NT1 [14,18]. The underlying pathogenesis of NT1 is thought to be due to autoimmune destruction of hypocretin-producing neurons in the hypothalamus; this hypothesis is backed by immune genetic and environmental factors linked to the disease [19-21]. There is evidence that specific genes influence narcolepsy. HLA allele DQB1\*0602 is often associated with cases of autoimmune response in NT1, however there are indications that another background related to methylation is also possible [5,22]. Further areas of active research into the causes of narcolepsy include the study of the histamine system, the search for other immune mediators, and the study of "secondary /symptomatic" narcolepsy attributed to damage to the hypothalamus as a result of traumatic, infectious or demyelinating processes [23-25].

Diagnostic criteria for narcolepsy are based on the presence of EDS, REM phase abnormalities on the Multiple Sleep Latency Test (MSLT), cataplexy and hypocretin levels in cerebrospinal fluid. In addition, in type 2 narcolepsy, the presence of other causes of excessive sleepiness and MSLT abnormalities should be excluded [26]. Previous research over the past four decades and across cultures has shown that narcolepsy can have a serious negative impact on the educational, psychosocial well-being, occupational and health-related quality of life (HRQOL) [27–30], even after medical interventions [31]. Human sexuality is a fundamental universal part of life, and positive sexual relations and sexual function are becoming increasingly recognized as significant markers of quality of life and positive health [32,33]. People with sexual difficulties have lower quality of life compared with healthy people [34,35]. What is the impact of narcolepsy on the sexual life of patients? How does narcolepsy affect that sphere of life?

The aim of the study were:

- 1) to determine if narcolepsy affects patients' sex life
- 2) to identify specific sexual dysfunctions in narcolepsy
- to determine the impact of drugs used in narcolepsy on sex life

#### Materials and methods

A narrative review of available literature in Polish, English and German was conducted, using keywords: narcolepsy, sexual dysfunctions, medications in narcolepsy, from repository inception to March 17, 2024 searching PubMed/MEDLINE, Google Scholar, Crossref, Cochrane databases. The review was based on information from case reports, original papers and reviews. The quality of the reviewed articles was assessed using the Scale for the Assessment of Narrative Review Articles (SANRA) [36].

# Results

For clarity, the review has been divided into the following subsections:

- 1. Impact of narcolepsy on sex life
- 2. Specific sexual dysfunction in narcolepsy
- 3. Impact of drugs used in narcolepsy on sex life

## 1. Impact of narcolepsy on sex life

According to the study conducted by Heloísa Rovere, Sueli Rossini, Rubens Reimão with usage of WHOQOL-BREF on 40 Brazilian narcoleptics, social relations domain, which includes sexual activity, is lower than in general population [37]. In a survey study conducted by Davidson et al. most of 254 narcoleptic participants stated that narcolepsy impacts their sex life (81.1%). Sleepiness may be the most significant cause of psychosocial disability in narcoleptic patients and can lead to sexual problems [38,39]. Sleep attacks, which occur even in 53.2% of narcoleptic patients, can as well impact sexual life in a significant level [40,41]. It is also important to assess the severity of the impact of narcolepsy on sexual dysfunction. Kapella et al. surveyed 120 narcoleptic patients, who rated their sexual life on a 5-point Likert scale with 0=extreme difficulty to 4= little problem. The average score was 3.0+/-0.8 [42]. In a study conducted by Alaia on 95 narcoleptics, 36% reported that their disease was associated with decrease of quality in sexual life. Impotence was the

most common sexual problem in men. When it comes to 15.8% of participants, they described the impact of their symptoms on the quality of their sexual life as significantly adverse. Participants provided ratings for overall sexual satisfaction using a 5-point scale, ranging from 1 (not satisfied) to 5 (very satisfied). The average score was 3.1, with a standard deviation of 1.58. A positive correlation was observed between the interpersonal relationship subscale and the sexual satisfaction scale. This indicates that individuals in supportive intimate relationships tend to report higher levels of sexual satisfaction [43]. The primary neuropathological alteration in narcolepsy involves the specific and permanent degeneration of neurons that produce hypocretin neuropeptide precursor (HCRT) in the lateral hypothalamus [44]. Individuals with NT1 frequently exhibit reduced levels of HCRT in their CSF. These hypocretin neurons project widely to various regions of the brainstem, influencing alterations in the autonomic nervous system [45]. Nevertheless, autonomic symptoms are often overlooked in comparison to the more characteristic symptoms observed in NT1 [46]. Through various assessments, such as microneurography and ambulatory blood pressure monitoring, multiple human studies have identified abnormal sympathetic activation in drug-free narcoleptic patients [47-49]. Therefore, patients with narcolepsy also often have autonomic disorders of the gastrointestinal, cardiovascular, urinary and reproductive systems, as well as dysfunction of thermoregulation and pupil reactivity. Lucie Barateau et al. assessed the severity of the above disorders with the SCOPA-AUT scale in 92 NT1 patients (not taking medication) and 109 healthy people (control group). Results of a study report that sexual dysfunction has been documented in 48% of men, primarily characterised by erection problems, and in 81% of women, primarily marked by issues related to vaginal lubrication. Also, NT1 patients were significantly more likely to report autonomic dysfunctions of other specific body systems than the control group [50]. Joshi et al. demonstrated the involvement of HCRT receptors in sex hormone synthesis by reducing serum testosterone levels in adult mice through the injection of an HCRT receptor antagonist [51]. In another study comparing serum gonadotropin levels in males with NT1, diminished pulsatile luteinizing hormone release was observed compared to controls, suggesting the involvement of HCRT in regulating hypothalamicpituitary-gonadal axis activity. The study group also had reduced levels of orexin and normal gonadal hormone concentrations in plasma [52]. Biobaku et al. presented a case report of a secondary narcolepsy due to a low level of testosterone [24]. Furthermore, inadequate testosterone levels and aberrant hypothalamic-pituitary-gonadal axis functioning may contribute to male sexual dysfunction. Hypocretin is actively involved in the regulation of the

dopaminergic system, which is crucial for motor control, but also in promoting wakefulness, in sexual behaviour and reward mechanism [39,53]. Karacan suggests that alterations in the sympathetic-parasympathetic balance linked to sleepiness may impact erectile function mediated by the autonomic nervous system. Other causes could be a greater prevalence of non-insulin-dependent diabetes (NIDD) among patients with narcolepsy. Diabetes could lead to neurogenic and vascular dysfunction that could cause erectile dysfunction. Another theory stated by Karacan is based on evidence that narcoleptic patients have impaired dopamine metabolism, which could lead to erectile dysfunction [41].

## 2. Specific sexual dysfunction in narcolepsy

#### 2.1. Orgasmolepsy

Cataplexy that occurs during orgasm and sexual intercourse was described for the first time and defined "orgasmolepsy" by Jakob Rothfield in 1928 [54]. Davidson et al. reported experiences of catalepsy during sex by almost one third of respondents, while 53.2% of patients fell asleep during intercourse. When it comes to 36.6% of the patients who experienced the above situations, they felt that it negatively affected their sexual activity [40]. Orgasmolepsy might be masked by other sexual disorders, which are commonly observed among NT1 patients as part of a dysautonomic syndrome [50]. However, the onset of an active sexual life may coincide with ongoing effective treatments for cataplexy in individuals with narcolepsy, thus preventing the occurrence of orgasmolepsy [50,55]. Patients experienced orgasmolepsy with their partners within stable relationships, implying that emotional disinhibition and reduced self-control may necessitate a state of intimacy as a precondition for orgasmolepsy. Studies have shown deactivation in the prefrontal cortex, responsible for self-control, during sexual arousal and orgasm, indicating the presence of emotional disinhibition [56,57]. Poryazova et al. propose that orgasmolepsy could be linked to continued amygdala firing during sexual intercourse, leading to the disinhibition of neurons responsible for generating muscle atonia in the pontine and medullary regions [39]. Other theories suggest involvement of hypocretin and dopaminergic systems in pathophysiology of orgasmolepsy [17,58-61]. It should be noted that orgasmolepsy effects on sexual function can have a significant impact on the patient's relationships and emotional well-being [55].

# 2.2. Sleep paralysis and sexual hallucinations

In general, 83% of narcolepsy patients indicated experiencing confusion between dreams and reality, in contrast to a mere 15% of individuals in the control group who reported similar experiences. Dream delusions could potentially represent merely one facet of a broader memory impairment observed in individuals with this disorder. There is a possibility that due to subjective memory difficulties they can confuse other people's stories as their memories. The inability to differentiate memories formed during sleep from those occurring during wakefulness may directly result from the welldocumented neural mechanisms associated with narcolepsy. E. Wemsley et al. speculate that disruption of the orexin system present in narcolepsy may lead to the occurrence of dream delusions, since orexin neurons primarily targets monoaminergic and cholinergic neurons responsible for regulating sleep states. This connection underscores their significance in the control of sleep-wake cycles and it can disrupt encoding of dreams as memories [62]. Sleep paralysis is often accompanied by HHG, which resembles dreams and occurs during the period of falling asleep [11]. Sleep paralysis and HHG are products of "sleeponset REM" (SOREMP), a REM stage that occurs earlier than normal when the patient is still partially conscious [4,11]. The emotional experience of sleep paralysis with hallucinations often involves fear, terror, and panic. Elements such as threatening presences, vulnerability while paralyzed, and uncontrollable visions contribute to intense, predominantly dysphoric, negative emotions. Some aspects of spontaneous thoughts during sleep paralysis can be interpreted as paranoid delusions [63,64]. These disorders profoundly affect social and sexual life and psychological well-being [65,66]. Impression of the presence of malevolent intruders in the bedroom, and physical/sexual assaults are common hallucination themes in sleep paralysis [62,63,67,68]. In a series of case reports by Hays, a 42-year-old woman claimed that she was sexually assaulted multiple times. According to investigators, it turns out to be probably hallucinations due to the onset of narcolepsy, that led the patient to false but sincere belief that she was assaulted. Another patient presented by Hays was 31 years old women with unrecognized narcolepsy, who had hallucinations that reduced her quality of life to such an extent that she attempted suicide. She had hallucinations that she was assaulted sexually. Several other cases similar to that are described by Hays [10]. Female patient claimed that she was sexually assaulted, she had a vivid recollection of this event and could present a detailed description of it. She reported it to the police. Investigation revealed that during the attack, she was seen in public space, sitting quietly. After a few hours, she realised the experience was not real. Treatment with 200 mg of modafinil per day decreased her sleepiness and 50 mg of clomipramine per day reduced her hallucinations to none [38]. Tjokrodipo L, Sneep A, Michielsen presented a case report of a patient with sleep paralysis and sexual hallucinations. The patient was also diagnosed with psychotic disorders, intellectual disability (IQ=74) and complications of cannabis use. During sleep paralysis, the patient experienced hallucinations of sexual intercourse with caregivers and the perception that a woman was sitting on his chest [69]. In a study reported by McNally and Clancy involving people who reported alien encounters, three participants experienced sleep paralysis with perceptions of sexual harassment by aliens. One of them showed some symptoms of past post-traumatic stress disorder (PTSD) after the incident [70]. Also a 24-year-old female patient with narcolepsy without cataplexy presented with visual HHG. She experienced episodes where she felt immobilized in bed, attempting to awaken but encountering dream hallucinations of sexual assault. The patient was treated with modafinil and antidepressants, leading to a reduction in hallucinations and improved daytime alertness [71]. Survivors of sexual assault often encounter various sexual difficulties post-assault, such as sexual dysfunction, diminished sexual satisfaction, fear of sexual intercourse [72]. For patients with narcolepsy nightmares, sexual hallucinations during sleep paralysis can be seen as real incidents. Sexual hallucinations also sometimes led to discontinuation of sexual intercourse. A case of a 46-year-old male patient, with a history of EDS, cataplexy, sleep paralysis, and HHG, was described by A. Moszczynski and F. Coelho. He experienced sporadic HHG episodes throughout the day, with frequent occurrences during the night. He revealed that his sexual HHG had been disrupting his sexual relations with his wife for the past six years. During these sexual HHG episodes, he described observing himself and his wife in action, akin to an out-of-body experience (OBE), accompanied by a sensation of floating. Treatment with methylphenidate and amitriptyline completely resolved the sexual HHG, although residual EDS persisted. Methylphenidate was subsequently substituted with modafinil, resulting in improved daytime alertness [73]. A 45-year-old patient with unrecognised narcolepsy with daytime sleepiness, sleep attacks, cataplexy and hallucinations had a vivid memory of sexual intercourse with his boss's wife. It led to severe conflict with his chief. Reaction of his colleagues convinced him that experience may not be true. He was diagnosed with narcolepsy and treated with 200 mg of modafinil and 50 mg of clomipramine per day, which led to decrease of his sleepiness and disappearance of hallucinations and delusions [38].

# 3. Impact of drugs used in narcolepsy on sex life

A survey study conducted by Teixeira VG, Faccenda JF, Douglas NJ in 2004 on patients with NT1 (n=49) showed that 41% of them had experienced sexual difficulties. Patients were on treatment with either

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stimulant medication, a combination of stimulant and anti-cataplexy agent or using only an anti-cataplexy agent. Three patients were taking no medication [74]. In a survey study conducted by Davidson et al. on 254 narcoleptic participants, among participants with NT1, the usage of drugs known to affect sexual function (selective serotonin reuptake inhibitors / tricyclic drugs) was not correlated with differences in effects on sexual life [40]. The effects of medications on autonomic symptoms in patients with NT1 were assessed. Upon individual analysis of each item of the questionnaire, a deterioration in sexual dysfunction was observed in men exclusively, particularly ejaculation problems. Among those under medication, 42.9% reported such symptoms, compared to 21.9% without treatment [50]. Modafinil, armodafinil, and solriamfetol are medications used to treat EDS [53]. The mechanism of action of modafinil is intricate and unique compared to other wakefulness-promoting medications. It regulates glutamate, GABA, histamine, and hypocretin, and to a lesser extent, the monoaminergic systems [75]. Studies have demonstrated that dopaminergic D1 and D2 receptors play a crucial role in mediating the arousal effects of modafinil [76]. Diminished libido is also a rare adverse reaction linked to modafinil [77]. Methylphenidate can increase libido and reduce erectile dysfunction [78]. It increases norepinephrine and dopamine transmission [79]. In a case report presented by Gregory Bierer, a 20-year-old patient with narcolepsy with cataplexy was firstly treated with 200 mg of modafinil. He reported tremor and agitation and erectile dysfunction, after that he was prescribed with armodafinil, but he experienced the same side effects. His treatment was changed to methylphenidate hydrochloride 10 mg at 8 am and 10 mg at noon if needed. Patient found this treatment to be effective for him. On the methylphenidate regimen he has not experienced cataplexy for 6 months [80]. Antidepressants are mainly treatment for cataplexy [53]. Moreover, almost all antidepressants employed in the management of cataplexy in individuals with NT1 are recognized for their potential side effects, including reduced libido, erectile dysfunction, delayed orgasm or anorgasmia, and delayed ejaculation [81]. The therapeutic effect of antidepressants on cataplexy primarily involves the inhibition of adrenergic uptake, whereas serotonin uptake blockers are comparatively less effective [82]. Pitolisant is histamine H3 receptor antagonist/inverse agonist [83], and is approved by the European Medicines Agency (EMA) for treatment of cataplexy and EDS [84]. Increased and decreased libido are not very common side effects of pitolisant [83]. Sodium oxybate is effective in addressing EDS, cataplexy, and fragmented nocturnal sleep [53]. Sodium oxybate potentially operates through GABA<sup>B</sup> receptors or its specific receptors, alongside

potential modulation of dopaminergic neurotransmission [85].

## Conclusion

Narcolepsy is a rare sleep disorder. According to studies, narcoleptic patients claim that narcolepsy has a negative impact on their quality of life, social life and sexual life [10,34,35,40]. Various studies reported numerous autonomic alterations in patients with narcolepsy, like impotence and problems with vaginal lubrication, which are associated with deficiency of hypocretin [44-47,50]. Other studies suggest that low level of hypocretin can cause sexual dysfunction through alterations in testosterone levels and dopaminergic system [24,39,52,53]. Karacan suggested that greater prevalence of non-insulin-dependent diabetes (NIDD) among patients with narcolepsy may also be the cause of erectile dysfunction as a complication of diabetes [41]. Other possible explanations of narcolepsy impact on sexual life are secondary impact of symptoms of the disease. One of these disorders is orgasmolepsy, which may be underreported due to the embarrassment, effectiveness of treatment, or it might be masked by other sexual disorders [39,40,50,55]. Studies suggest alterations in amygdala, as an explanation for the occurrence of orgasmolepsy [39] or involvement of dopaminergic and orexin systems [58-61,86]. This aspect of narcolepsy's effect on sexual function can have a significant impact on the patient's relationships, emotional well-being and sexual life [41,55]. Honest conversation with a sexual partner may be helpful for easing the distress following this symptom [40]. Pitolisant and sodium oxybate seem to be effective treatments for this symptom in some cases [39,55]. In some patients dreams delusions occur as nightmares and are very terrifying and vivid, with aggressive content like sexual assaults, which some patients interpret as real memories, what can affect psychosocial well-being [62]. Some narcoleptic patients experience hallucinations or sleep paralysis with hallucinations of being sexually assaulted, which can profoundly affect social life and psychological well-being or cause suicide attempts [62-66,73]. Also in some cases it may lead to development of severe stress and occurrence of some symptoms of PTSD [70]. In most cases hallucinations can be treated with stimulants [10,71,73]. Medical providers should ask clear, concrete and direct questions, maintaining nonjudgmental, when asking the social life of a narcoleptic patient. There is an increasing interest in developing behavioural interventions aimed at enhancing the quality of life for people with narcolepsy [40]. One method is to schedule naps during the day. Typically, they last from 15 to 20 minutes. At the same time, it is recommended to extend nocturnal sleep. In addition, physical activity can have a positive impact [79]. Lack of understanding for symptoms of narcolepsy is common. Open communications through meeting with psychologists could lead to building an understanding and solution for sexual problems [40]. The recommended therapy is cognitive behavioural therapy (CBT). The patient learns to notice and modify disturbances in his or her way of thinking and acting. This improves adherence to medication, as well as the patient's functioning and quality of sex life [79]. After a diagnosis of narcolepsy is established, mere prescription of medication is insufficient. Continuous support and counselling for patients are imperative. Educating the patient, their family, and potentially their employer is necessary to alleviate social pressures on these individuals. While support groups for narcoleptics and their families can be beneficial, they have yielded mixed results. Without informed guidance, some groups have disseminated inaccurate information and ineffective therapies. Involvement of sleep professionals is crucial for these support groups to be truly effective [43]. Medications commonly used to manage symptoms of narcolepsy, such as antidepressants and stimulants, including imipramine, clomipramine, fluoxetine, citalopram, dextroamphetamine, phenelzine, and others, have been consistently associated with sexual symptoms. This includes difficulties achieving or maintaining erections, reduced libido, premature ejaculation, and delayed orgasm [50,74,79–81]. Methylphenidate seems to have a positive impact on libido and erectile function [78,80]. There is a speculation that timing sexual activity when drug levels are lower in the morning may reduce interference with sexual function and result in better quality erections. However, formal clinical trials to validate this notion are lacking [41]. A multidisciplinary approach that combines aspects of patient and his family education, as well as behavioural therapy, is important in the treatment of sexual dysfunction in narcolepsy [40,79].

## **Conflict of interest**

The authors have declared no conflict of interest.

#### **References:**

- Nishino S, Mignot E. Narcolepsy and cataplexy. Handb Clin Neurol Ed PJ Vinken GW Bruyn. 31 December 2011;99:783–814. https://doi.org/10.1016/B978-0-444-52007-4.00007-2
- Mahoney CE, Cogswell A, Koralnik IJ, Scammell TE. The neurobiological basis of narcolepsy. Nat Rev Neurosci. February 2019;20(2):83–93. https://doi.org/10.1038/s41583-018-0097-x
- AASM | Clinical Resources | International Classification of Sleep Disorders [Internet]. American Academy of Sleep Medicine

   Association for Sleep Clinicians and Researchers. [cited 9 March 2024]. Available on: https://aasm.org/clinical-resources/ international-classification-sleep-disorders/
- 4. Szûcs A, Mutti C, Papp A, Halász P, Parrino L. REM sleep, REM parasomnias, REM sleep behaviour disorder. Ideggyogyaszati

Szle. 30 May 2022;75(5-06):171-82. HTTPS://DOI. ORG/10.18071/ISZ.75.0171

- Dauvilliers Y, Arnulf I, Mignot E. Narcolepsy with cataplexy. Lancet Lond Engl. 10 February 2007;369(9560):499–511. https://doi.org/10.1016/S0140-6736(07)60237-2
- Ohayon MM. Prevalence of hallucinations and their pathological associations in the general population. Psychiatry Res. 27 December 2000;97(2–3):153–64. https://doi.org/10.1016/ S0165-1781(00)00227-4
- Billiard M. Narcolepsy. Clinical features and aetiology. Ann Clin Res. 1985;17(5):220–6.
- Leu-Semenescu S, De Cock VC, Le Masson VD, Debs R, Lavault S, Roze E, i in. Hallucinations in narcolepsy with and without cataplexy: contrasts with Parkinson's disease. Sleep Med. May 2011;12(5):497–504. https://doi.org/10.1016/j. sleep.2011.03.006
- Matas M, Marriott A. The girl who cried wolf: Pseudologia phantastica and sexual abuse. Can J Psychiatry Rev Can Psychiatr. 1987;32(4):305–9. https://doi. org/10.1177/070674378703200412
- Hays P. False but sincere accusations of sexual assault made by narcoleptic [correction of narcotic] patients,. Med Leg J. 1992;60 (Pt 4):265–71. https://doi.org/10.1177/002581729206000405
- Stefani A, Högl B. Nightmare Disorder and Isolated Sleep Paralysis. Neurotherapeutics. January 2021;18(1):100-6. https://doi.org/10.1007/s13311-020-00966-8
- Schenck CH, Bassetti CL, Arnulf I, Mignot E. English Translations Of The First Clinical Reports On Narcolepsy And Cataplexy By Westphal And Gélineau In The Late 19th Century, With Commentary. J Clin Sleep Med JCSM Off Publ Am Acad Sleep Med. 15 April 2007;3(3):301–11. PMID: 17561602; PMCID: PMC2564780.
- Mignot E. A hundred years of narcolepsy research. Arch Ital Biol. April 2001;139(3):207–20. https://doi.org/10.4449/aib. v139i3.500
- Nishino S, Ripley B, Overeem S, Lammers GJ, Mignot E. Hypocretin (orexin) deficiency in human narcolepsy. Lancet Lond Engl. 1 January 2000;355(9197):39–40. https://doi. org/10.1016/S0140-6736(99)05582-8
- Thannickal TC, Moore RY, Nienhuis R, Ramanathan L, Gulyani S, Aldrich M, i in. Reduced number of hypocretin neurons in human narcolepsy. Neuron. September 2000;27(3):469–74. https://doi. org/10.1016/S0896-6273(00)00058-1
- Mignot E, Lammers GJ, Ripley B, Okun M, Nevsimalova S, Overeem S, i in. The role of cerebrospinal fluid hypocretin measurement in the diagnosis of narcolepsy and other hypersomnias. Arch Neurol. October 2002;59(10):1553–62. https://doi.org/10.1001/archneur.59.10.1553
- Kanbayashi T, Inoue Y, Chiba S, Aizawa R, Saito Y, Tsukamoto H, i in. CSF hypocretin-1 (orexin-A) concentrations in narcolepsy with and without cataplexy and idiopathic hypersomnia. J Sleep Res. March 2002;11(1):91–3. https://doi.org/10.1046/j.1365-2869.2002.00284.x
- Thannickal TC, Nienhuis R, Siegel JM. Localized loss of hypocretin (orexin) cells in narcolepsy without cataplexy. Sleep. August 2009;32(8):993–8. https://doi.org/10.1093/ sleep/32.8.993
- Sarkanen TO, Alakuijala APE, Dauvilliers YA, Partinen MM. Incidence of narcolepsy after H1N1 influenza and vaccinations: Systematic review and meta-analysis. Sleep Med Rev. April 2018;38:177–86. https://doi.org/10.1016/j.smrv.2017.06.006
- 20. Lind A, Ramelius A, Olsson T, Arnheim-Dahlström L, Lamb F, Khademi M, i in. A/H1N1 antibodies and TRIB2 autoantibodies in narcolepsy patients diagnosed in conjunction with the

Pandemrix vaccination campaign in Sweden 2009-2010. J Autoimmun. May 2014;50:99–106. https://doi.org/10.1016/j. jaut.2014.01.031

- 21. Koepsell TD, Longstreth WT, Ton TGN. Medical exposures in youth and the frequency of narcolepsy with cataplexy: a population-based case-control study in genetically predisposed people. J Sleep Res. March 2010;19(1 Pt 1):80–6. https://doi. org/10.1111/j.1365-2869.2009.00756.x
- Seifinejad A, Ramosaj M, Shan L, Li S, Possovre ML, Pfister C, i in. Epigenetic silencing of selected hypothalamic neuropeptides in narcolepsy with cataplexy. Proc Natl Acad Sci U S A. 9 May 2023;120(19):e2220911120. https://doi.org/10.1073/ pnas.2220911120
- Scammell TE, Mochizuki T. Is Low Histamine a Fundamental Cause of Sleepiness in Narcolepsy and Idiopathic Hypersomnia? Sleep. 1 February 2009;32(2):133–4. https://doi.org/10.1093/ sleep/32.2.133
- Biobaku F, Hanna A, Matthews G, Dhindsa S, Dandona P. Narcolepsy, depression, and severe flushing in an obese man. Clin Case Rep. 10 August 2020;8(9):1824–6. https://doi. org/10.1002/ccr3.2873
- Nishino S, Kanbayashi T. Symptomatic narcolepsy, cataplexy and hypersomnia, and their implications in the hypothalamic hypocretin/orexin system. Sleep Med Rev. August 2005;9(4):269–310. https://doi.org/10.1016/j. smrv.2005.03.004
- 26. Sleep Medicine Textbook | ESRS Sleep Book 2nd Edition [Internet]. [cited 17 MArch 2024]. Dostępne na: https://esrs.eu/ sleep-medicine-textbook/
- 27. Ervik S, Abdelnoor M, Heier MS, Ramberg M, Strand G. Healthrelated quality of life in narcolepsy. Acta Neurol Scand. September 2006;114(3):198–204. https://doi.org/10.1111/ j.1600-0404.2006.00594.x
- Raggi A, Plazzi G, Ferri R. Health-Related Quality of Life in Patients With Narcolepsy: A Review of the Literature. J Nerv Ment Dis. February 2019;207(2):84–99. https://doi. org/10.1097/NMD.000000000000918
- David A, Constantino F, dos Santos JM, Paiva T. Health-related quality of life in Portuguese patients with narcolepsy. Sleep Med. March 2012;13(3):273–7. https://doi.org/10.1016/j. sleep.2011.06.021
- Daniels E, King MA, Smith IE, Shneerson JM. Health-related quality of life in narcolepsy. J Sleep Res. March 2001;10(1):75– 81. https://doi.org/10.1046/j.1365-2869.2001.00234.x
- 31. Ozaki A, Inoue Y, Hayashida K, Nakajima T, Honda M, Usui A, i in. Quality of life in patients with narcolepsy with cataplexy, narcolepsy without cataplexy, and idiopathic hypersomnia without long sleep time: comparison between patients on psychostimulants, drug-naïve patients and the general Japanese population. Sleep Med. February 2012;13(2):200–6. https://doi. org/10.1016/j.sleep.2011.07.014
- 32. Lee DM, Vanhoutte B, Nazroo J, Pendleton N. Sexual Health and Positive Subjective Well-Being in Partnered Older Men and Women. J Gerontol B Psychol Sci Soc Sci. July 2016;71(4):698– 710. https://doi.org/10.1093/geronb/gbw018
- Rosen RC, Bachmann GA. Sexual well-being, happiness, and satisfaction, in women: the case for a new conceptual paradigm. J Sex Marital Ther. 2008;34(4):291–7; discussion 298-307. https://doi.org/10.1080/00926230802096234
- Hicham M, Abdellatif B, Nadia O, Hakima A. The effect of sexuality on the quality of life of elderly people in Morocco. Afr J Reprod Health. 31 August 2023;27(8):76–82. https://doi. org/10.29063/ajrh2023/v27i8.8
- 35. Peixoto MM, Lopes J. Quality of life and emotional well-being

during COVID-19 as mediators in the relationship between sexual functioning and satisfaction in Portuguese women. Women Health. April 2022;62(4):358–68. https://doi.org/10.10 80/03630242.2022.2074609

- Baethge C, Goldbeck-Wood S, Mertens S. SANRA—a scale for the quality assessment of narrative review articles. Res Integr Peer Rev. 26 March 2019;4(1):5. https://doi.org/10.1186/s41073-019-0064-8
- Rovere H, Rossini S, Reimão R. Quality of life in patients with narcolepsy: a WHOQOL-bref study. Arq Neuropsiquiatr. June 2008;66(2A):163-7. https://doi.org/10.1590/s0004-282x2008000200004
- Szűcs A, Janszky J, Holló A, Migléczi G, Halász P. Misleading hallucinations in unrecognized narcolepsy. Acta Psychiatr Scand. 2003;108(4):314–7. https://doi.org/10.1034/j.1600-0447.2003.00114.x
- Poryazova R, Khatami R, Werth E, Bassetti CL. Weak with sex: sexual intercourse as a trigger for cataplexy. J Sex Med. August 2009;6(8):2271–7. https://doi.org/10.1111/j.1743-6109.2009.01328.x
- 40. Davidson RD, Biddle K, Nassan M, Scammell TE, Zhou ES. The impact of narcolepsy on social relationships in young adults. J Clin Sleep Med JCSM Off Publ Am Acad Sleep Med. 1 December 2022;18(12):2751–61. DOI: https://doi.org/10.5664/jcsm.10212
- Karacan I, Gokcebay N, Hirshkowitz M, Ozmen M, Ozmen E, Williams RL. Sexual Dysfunction in Men with Narcolepsy. Loss Grief Care. 10 June 1992;5(3-4):81-90.
- Kapella MC, Berger BE, Vern BA, Vispute S, Prasad B, Carley DW. Health-Related Stigma as a Determinant of Functioning in Young Adults with Narcolepsy. PLOS ONE. 21 April 2015;10(4):e0122478. https://doi.org/10.1371/journal. pone.0122478
- 43. Alaia SL. Life Effects of Narcolepsy: Measures of Negative Impact, Social Support, and Psychological Well-Being. W: Psychosocial Aspects of Narcolepsy. Routledge; 1992.
- 44. Ito E, Inoue Y. [The International Classification of Sleep Disorders, third edition. American Academy of Sleep Medicine. Includes bibliographies and index]. Nihon Rinsho Jpn J Clin Med. June 2015;73(6):916–23.
- 45. Grimaldi D, Silvani A, Benarroch EE, Cortelli P. Orexin/ hypocretin system and autonomic control: new insights and clinical correlations. Neurology. 21 January 2014;82(3):271–8. https://doi.org/10.1212/WNL.000000000000045
- 46. Wang Y, Sun Q, Tang Q, Zhang Y, Tang M, Wang D, i in. Progress of autonomic disturbances in narcolepsy type 1. Front Neurol. 6 March 2023;14:1107632. https://doi.org/10.3389/ fneur.2023.1107632
- Plazzi G, Moghadam KK, Maggi LS, Donadio V, Vetrugno R, Liguori R, i in. Autonomic disturbances in narcolepsy. Sleep Med Rev. June 2011;15(3):187–96. https://doi.org/10.1016/j. smrv.2010.05.002
- Donadio V, Liguori R, Vandi S, Pizza F, Dauvilliers Y, Leta V, i in. Lower wake resting sympathetic and cardiovascular activities in narcolepsy with cataplexy. Neurology. 16 September 2014;83(12):1080–6. https://doi.org/10.1212/ WNL.000000000000793
- 49. Dauvilliers Y, Jaussent I, Krams B, Scholz S, Lado S, Levy P, i in. Non-dipping blood pressure profile in narcolepsy with cataplexy. PloS One. 2012;7(6):e38977. https://doi.org/10.1371/ journal.pone.0038977
- Barateau L, Chenini S, Evangelista E, Jaussent I, Lopez R, Dauvilliers Y. Clinical autonomic dysfunction in narcolepsy type 1. Sleep. 24 December 2019;42(12):zsz187. https://doi. org/10.1093/sleep/zsz187

- Joshi D, Singh SK. The neuropeptide orexin A search for its possible role in regulation of steroidogenesis in adult mice testes. Andrology. May 2018;6(3):465–77. https://doi. org/10.1111/andr.12475
- 52. Kok SW, Roelfsema F, Overeem S, Lammers GJ, Frölich M, Meinders AE, i in. Pulsatile LH release is diminished, whereas FSH secretion is normal, in hypocretin-deficient narcoleptic men. Am J Physiol Endocrinol Metab. October 2004;287(4):E630-636. https://doi.org/10.1152/ajpendo.00060.2004
- Bassetti CLA, Adamantidis A, Burdakov D, Han F, Gay S, Kallweit U, i in. Narcolepsy - clinical spectrum, aetiopathophysiology, diagnosis and treatment. Nat Rev Neurol. September 2019;15(9):519–39. https://doi.org/10.1038/s41582-019-0226-9
- Rothfeld J. Affektiver Tonus- und Bewußtseinsverlust beim Lachen und Orgasmus (Gelo- und Orgasmolepsia). Z Für Gesamte Neurol Psychiatr. 1 December 1928;115(1):516–30.
- Pellitteri G, Dolso P, Valente M, Gigli GL. Orgasmolepsy in Narcolepsy Type 1 Responsive to Pitolisant: A Case Report. Nat Sci Sleep. 31 December 2020;12:1237–40. https://doi. org/10.2147/NSS.S286358
- 56. Georgiadis JR, Kortekaas R, Kuipers R, Nieuwenburg A, Pruim J, Reinders AATS, i in. Regional cerebral blood flow changes associated with clitorally induced orgasm in healthy women. Eur J Neurosci. December 2006;24(11):3305–16 https://doi.org/10.1111/j.1460-9568.2006.05206.x
- Georgiadis JR, Reinders AATS, Van der Graaf FHCE, Paans AMJ, Kortekaas R. Brain activation during human male ejaculation revisited. Neuroreport. 16 April 2007;18(6):553–7. https://doi. org/10.1097/WNR.0b013e3280b10bfe
- Clément P, Pozzato C, Heidbreder C, Alexandre L, Giuliano F, Melotto S. Delay of ejaculation induced by SB-277011, a selective dopamine D3 receptor antagonist, in the rat. J Sex Med. April 2009;6(4):980–8. https://doi.org/10.1111/j.1743-6109.2008.01173.x
- 59. Marcus JN, Aschkenasi CJ, Lee CE, Chemelli RM, Saper CB, Yanagisawa M, i in. Differential expression of orexin receptors 1 and 2 in the rat brain. J Comp Neurol. 18 June 2001;435(1):6–25. https://doi.org/10.1002/cne.1190
- 60. Narita M, Nagumo Y, Miyatake M, Ikegami D, Kurahashi K, Suzuki T. Implication of protein kinase C in the orexin-induced elevation of extracellular dopamine levels and its rewarding effect. Eur J Neurosci. March 2007;25(5):1537–45. https://doi. org/10.1111/j.1460-9568.2007.05403.x
- Muschamp JW, Dominguez JM, Sato SM, Shen RY, Hull EM. A role for hypocretin (orexin) in male sexual behavior. J Neurosci Off J Soc Neurosci. 14 March 2007;27(11):2837–45. https://doi. org/10.1523/JNEUROSCI.4121-06.2007
- Wamsley E, Donjacour CEHM, Scammell TE, Lammers GJ, Stickgold R. Delusional Confusion of Dreaming and Reality in Narcolepsy. Sleep. 1 February 2014;37(2):419–22. https://doi. org/10.5665/sleep.3428
- 63. Cheyne JA, Girard TA. Paranoid delusions and threatening hallucinations: a prospective study of sleep paralysis experiences. Conscious Cogn. December 2007;16(4):959–74. https://doi.org/10.1016/j.concog.2007.01.002
- 64. Nielsen T. Felt presence: paranoid delusion or hallucinatory social imagery? Conscious Cogn. December 2007;16(4):975–83; discussion 984-991. https://doi.org/10.1016/j. concog.2007.02.002
- 65. Solomonova E, Nielsen T, Stenstrom P, Simard V, Frantova E, Donderi D. Sensed presence as a correlate of sleep paralysis distress, social anxiety and waking state social imagery. Conscious Cogn. March 2008;17(1):49–63. https://doi.

org/10.1016/j.concog.2007.04.007

- Sharpless BA, Barber JP. Lifetime Prevalence Rates of Sleep Paralysis: A Systematic Review. Sleep Med Rev. October 2011;15(5):311–5. https://doi.org/10.1016/j.smrv.2011.01.007
- Fosse R. REM mentation in narcoleptics and normals: an empirical test of two neurocognitive theories. Conscious Cogn. December 2000;9(4):488–509. https://doi.org/10.1006/ ccog.2000.0466
- Mazzetti M, Bellucci C, Mattarozzi K, Plazzi G, Tuozzi G, Cipolli C. REM-dreams recall in patients with narcolepsy-cataplexy. Brain Res Bull. 15 January 2010;81(1):133–40. https://doi. org/10.1016/j.brainresbull.2009.10.021
- Tjokrodipo L, Sneep A, Michielsen P. A case of incubus phenomenon. Eur Psychiatry. April 2021;64(S1):S818–S818. https://doi.org/10.1192/j.eurpsy.2021.2162.
- McNally RJ, Clancy SA. Sleep Paralysis, Sexual Abuse, and Space Alien Abduction. Transcult Psychiatry. 2005;42(1):113–22. https://doi.org/10.1177/1363461505050715
- Espinoza Lopez DA, Alfaro EC, Booth VA. 60. Hallucinations with sexual context as manifestation of narcolepsy without cataplexy. Case report. Clin Neurophysiol. 1 September 2016;127(9):e316. https://doi.org/10.1016/j.clinph.2016.05.335
- 72. van Berlo W, Ensink B. Problems with sexuality after sexual assault. Annu Rev Sex Res. 2000;11:235–57.
- Moszczynski A, Coelho F. Sexual hypnagogic hallucinations and narcolepsy with cataplexy: A case report. Sleep Sci. 1 January 2011;4:110–2.
- Teixeira VG, Faccenda JF, Douglas NJ. Functional status in patients with narcolepsy. Sleep Med. September 2004;5(5):477– 83. https://doi.org/10.1016/j.sleep.2004.07.001
- Ballon JS, Feifel D. A systematic review of modafinil: Potential clinical uses and mechanisms of action. J Clin Psychiatry. April 2006;67(4):554–66. https://doi.org/10.4088/jcp.v67n0406
- Qu WM, Huang ZL, Xu XH, Matsumoto N, Urade Y. Dopaminergic D1 and D2 receptors are essential for the arousal effect of modafinil. J Neurosci Off J Soc Neurosci. 20 August 2008;28(34):8462–9. https://doi.org/10.1523/ JNEUROSCI.1819-08.2008
- Modafinil referral | European Medicines Agency [Internet]. [cited 9 March 2024]. Available on: https://www.ema.europa. eu/en/medicines/human/referrals/modafinil
- Nazlı ŞB, Sevindik M. Use of Methylphenidate in Coexisting Major Depression, Loss of Libido and Erectile Dysfunction. Alpha Psychiatry. January 2021;22(1):70. https://doi.org/10.5455/ apd.112405
- 79. Franceschini C, Pizza F, Antelmi E, Folli MC, Plazzi G. Narcolepsy treatment: pharmacological and behavioural strategies in adults and children. Sleep Breath. 1 June 2020;24(2):615–27. https://doi.org/10.1007/s11325-019-01894-4
- 80. Bierer G, Banashek LR, Wen E. Narcolepsy with Cataplexy: Treatment Tailored to the Patient. 2013;17.
- Montejo AL, Llorca G, Izquierdo JA, Rico-Villademoros F. Incidence of sexual dysfunction associated with antidepressant agents: a prospective multicenter study of 1022 outpatients. Spanish Working Group for the Study of Psychotropic-Related Sexual Dysfunction. J Clin Psychiatry. 2001;62 Suppl 3:10–21.
- Nishino S. Narcolepsy: pathophysiology and pharmacology. J Clin Psychiatry. 2007;68 Suppl 13:9–15.
- Wakix | European Medicines Agency [Internet]. [cited 9 march 2024]. Available on: https://www.ema.europa.eu/en/ medicines/human/EPAR/wakix
- 84. Guevarra JT, Hiensch R, Varga AW, Rapoport DM. Pitolisant to Treat Excessive Daytime Sleepiness and Cataplexy in Adults with Narcolepsy: Rationale and Clinical Utility. Nat Sci Sleep. 12

October 2020;12:709-19. https://doi.org/10.2147/NSS.S264140

- Bernasconi R, Mathivet P, Bischoff S, Marescaux C. Gammahydroxybutyric acid: an endogenous neuromodulator with abuse potential? Trends Pharmacol Sci. April 1999;20(4):135– 41. https://doi.org/10.1016/s0165-6147(99)01341-3
- 86. Gulia KK, Mallick HN, Kumar VM. Orexin A (hypocretin-1) application at the medial preoptic area potentiates male sexual behavior in rats. Neuroscience. 2003;116(4):921–3. https://doi. org/10.1016/s0306-4522(02)00877-1

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