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Polysomnographic features of sleep paralysis (SP) captured in a patient with narcolepsy

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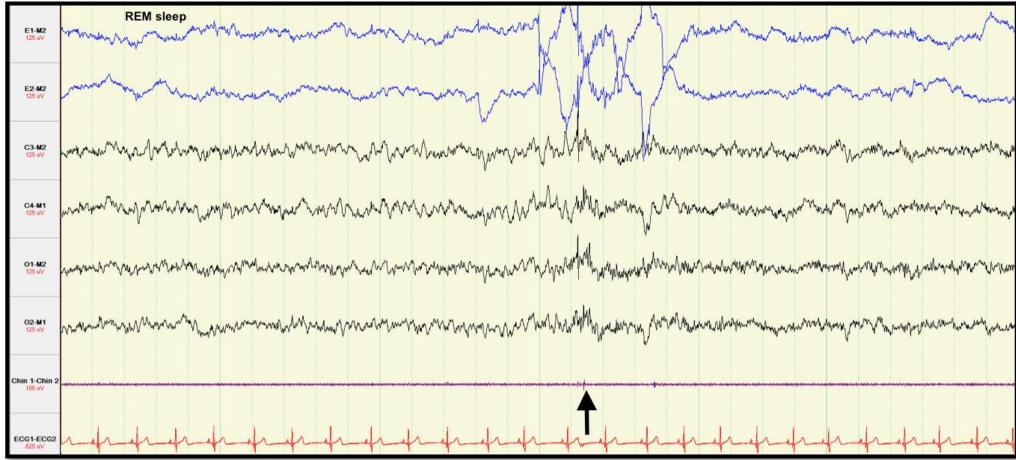
Introduction

Multiple sleep latency test (MSLT) was consistent with Humans rely on a mechanism known as muscle atonia narcolepsy. She reported an episode of SP arising from her during REM sleep in order to protect against injurious final sleep-onset REM period (SOREMP). Figure 1 displays behavior that may occur during emotionally-charged dream enactment [1]. SP is a transient episode of muscle atonia that a 30-second epoch marking the transition from REM sleep to wakefulness. Initially, she is in REM sleep, characterized occurs during the transition between sleep and wakefulness, clinically manifesting as inability to perform voluntary by theta-predominant EEG background and chin EMG movement while consciousness is entirely preserved [2,3]. atonia. Phasic eye movements are not captured in this tracing. The arrow depicts the transition to wakefulness, Isolated SP is known to occur in healthy individuals [1], however, recurrent SP is a well-documented feature of characterized by alpha-predominant EEG, however EMG narcolepsy [4]. There is limited literature describing the tone remains suppressed, indicating SP. polysomnographic features of SP, partly owing to its rarity and unlikelihood of being captured during a single night's mound have been and the property E2-M2 study [3]. We aim to illustrate the key neurophysiological features in a patient experiencing SP on polysomnogram.

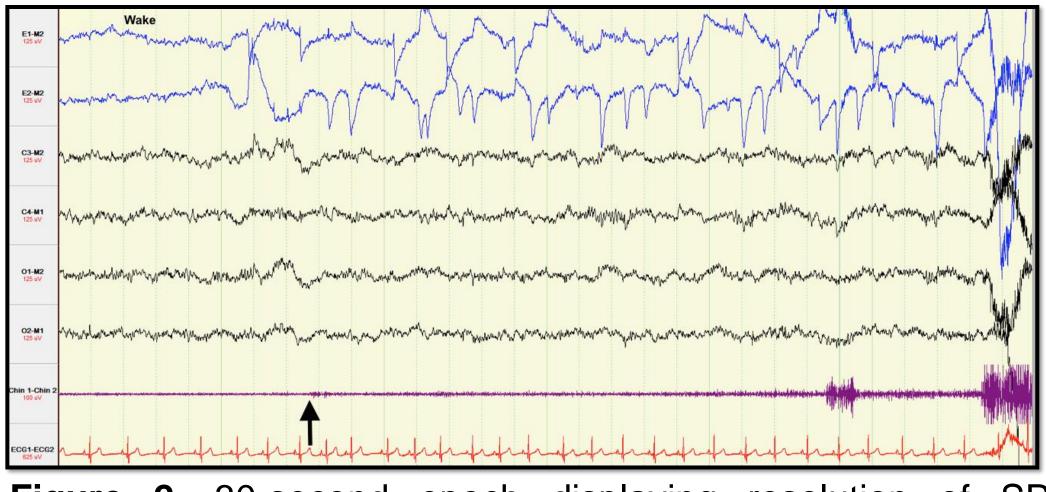
Methods

We report a 24-year-old female who presented to our sleep clinic for excessive daytime somnolence and recurrent SP **Figure 1.** 30-second epoch displaying SP occurring during the transition from REM-sleep to wakefulness depicted by the episodes upon awakening from naps. Despite sleeping 12 arrow. hours per night, she reported an irresistible urge to nap during the day. She also reported experiencing a vivid References "dream-like" state and inability to move her body voluntarily upon awakening from naps. During such times, 1. Olunu E, Kimo R, Onigbinde EO, et al. Sleep paralysis, a medical condition with a diverse cultural interpretation. Appl Basic Med Res 2018;8(3):137-142. she experienced a sensation of fear, causing her to 2. Plazzi G, Serra L and Ferri R. Nocturnal aspects of narcolepsy with cataplexy. Sleep Med Rev 2008;12(2):109-28. hyperventilate. She denied having episodes of cataplexy.

Results



- 3. Denis, D. Relationship between sleep paralysis and sleep quality: current insights. Nature and Science of Sleep 2018;10:355-367.
- 4. Amara, AW, Maddox, MH. Epidemiology of Sleep Medicine. In: Kryger MH, Roth T, Dement C, eds. Principles and Practice of Sleep Medicine, 6th ed. Philadelphia: Elsevier. 2017.
- 5. Dauvilliers Y, Barateau L. Narcolepsy and other central hypersomnias. Continuum (Minneap Minn) 2017;23(4):989-1004.
- 6. Girard TA, Cheyne JA. Timing of spontaneous sleep-paralysis episodes. J Sleep Res 2006;15(2)222-229.





Our patient's MSLT tracing displays mutual existence of REM and wake EEG features occurring in tandem with persistent muscle atonia on chin EMG [3]. This phenomenon underlies the clinical experience of immobility despite transitioning from REM sleep to wakefulness. In narcolepsy, there is loss of hypocretin-mediated suppression of REM-sleep [5], which may explain the inappropriate intrusion of REM-sleep features into wakefulness. SP is most common within the first two hours of sleep-onset or within the hours prior to awakening, corresponding to when timing of SOREMPs and REM-sleep predominate, respectively [2,6].



Figure 2 displays a 30-second epoch marking resolution of SP during wakefulness. The arrow depicts gradual increase in EMG tone, followed by bursts of muscle tone enhancement, indicating resolution of SP.

Figure 2. 30-second epoch displaying resolution of SP episode during wakefulness, depicted by the arrow.

Discussion/Conclusion