SLEEP 2022, the 36th Annual Meeting of the Associated Professional Sleep Societies (APSS)
June 4-8, 2022 • Charlotte, NC

Introduction

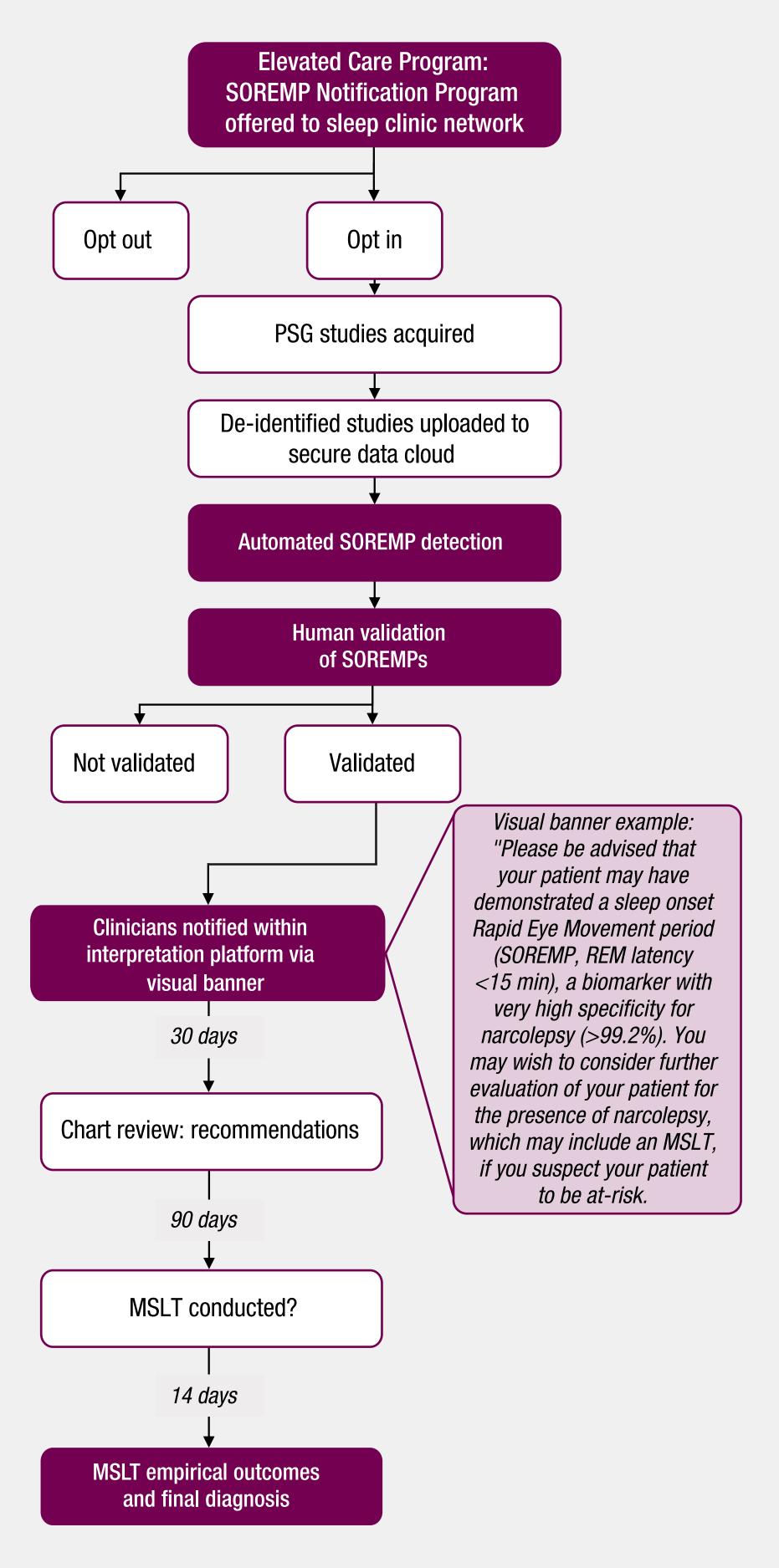
- Narcolepsy frequently remains undiagnosed for many years following symptom onset, likely due, in part, to a combination of limited narcolepsy-specific education for clinicians and few opportunities to gain experience, as well as substantial medical comorbidity (eg, sleep apnea) that may be associated with similar symptoms¹
- Rapid eye movement (REM) sleep detected by polysomnography (PSG) occurring within 15 minutes of nocturnal sleep (sleep onset REM period; SOREMP) is a known biomarker for hypocretin-deficient narcolepsy²
- However, among patients who undergo routine diagnostic sleep testing, as few as 4% with a SOREMP receive further evaluation for narcolepsy, suggesting that SOREMPs are often underappreciated in these individuals³
- To enhance identification of SOREMP episodes, an automated process was developed to detect and advise sleep clinicians of SOREMPs occurring during overnight PSG

Objective

• To evaluate the impact of automated SOREMP notification on clinician recommendations for narcolepsy diagnostic evaluation and multiple sleep latency test (MSLT) outcomes

Methods

Figure 1. Study Design



MSLT, multiple sleep latency test; PSG, polysomnography; REM, rapid eye movement; SOREMP, sleep onset rapid eye movement period.

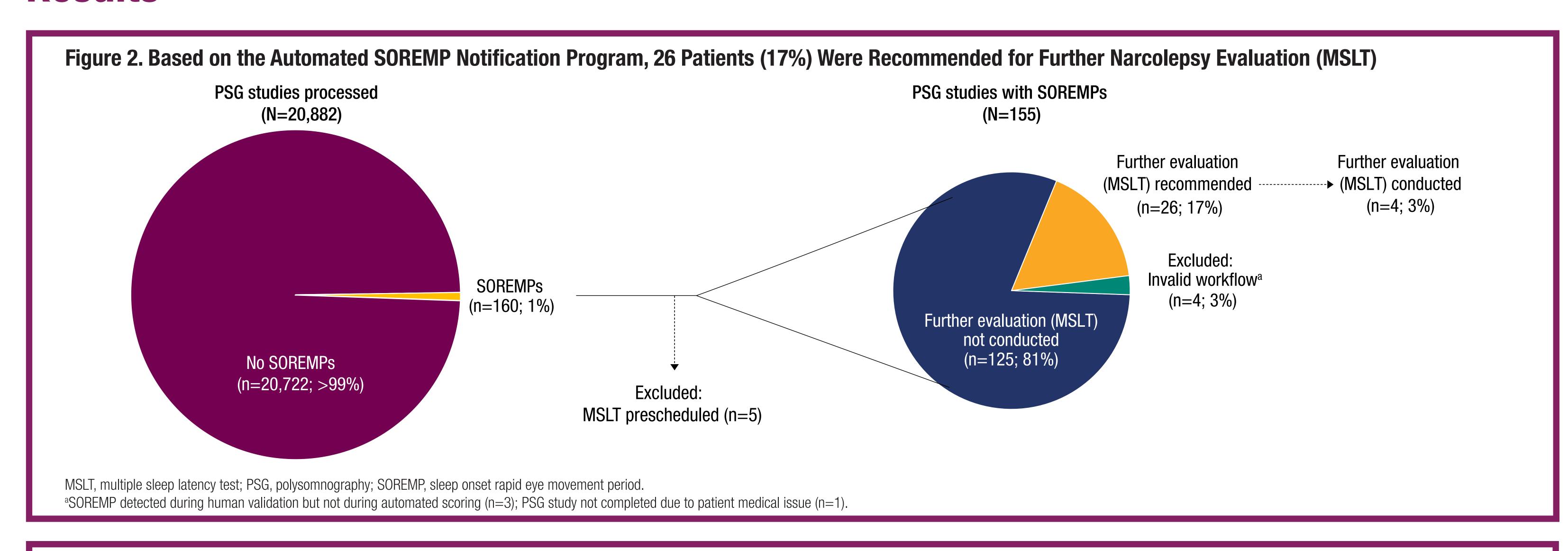
Sleep studies were processed over 3 years

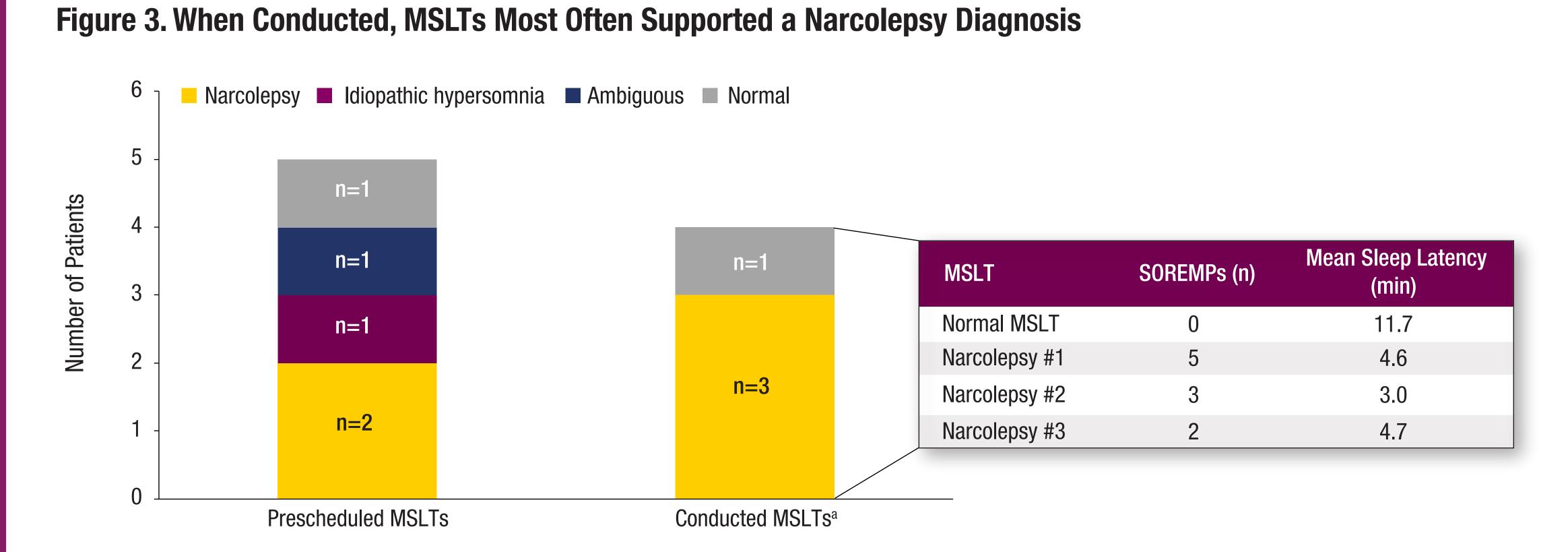
A Narcolepsy Detection Enhancement Paradigm: Automated Nocturnal Detection and Notification of Sleep Onset Rapid Eye Movement Periods

Alyssa Cairns, PhD¹; Richard K. Bogan, MD²; Alex Zheng, PhD³; Shay Bujanover, MD⁴; Prasheel Lillaney, PhD⁵; Andrew Friedberg, PhD⁵; Jed Black, MD⁵,

¹BioSerenity, Columbia, SC, USA; ²University of South Carolina School of Medicine, Columbia, SC, USA; ³Huneo, Fremont, CA, USA; ⁴Formerly Jazz Pharmaceuticals, Palo Alto, CA, USA; ⁵Jazz Pharmaceuticals, Palo Alto, CA, USA; ⁶Stanford University Center for Sleep Sciences and Medicine, Palo Alto, CA, USA

Results





- Of the 26 patients that were recommended for further evaluation, 4 had subsequent MSLTs conducted, 2 had MSLTs scheduled, and 20 had no action at time of printing
- The 4 MSLTs that were conducted based on recommendations from automated PSG screening were mostly consistent with narcolepsy
- Three patients (75%) had numbers of SOREMPs and mean sleep latency values from MSLT that supported the narcolepsy diagnosis
- One patient (25%) had a normal MSLT

Conclusions

- This study implemented enhanced identification and subsequent clinician notification of nocturnal SOREMPs using a novel sensitive SOREMP detection paradigm
- This methodology resulted in the recommendation of further narcolepsy evaluation (MSLT) for 26 patients (17%) who may have otherwise gone undetected
- Most MSLTs (75%) that were scheduled due to the recommendation and for which a diagnosis was listed were consistent with a diagnosis of narcolepsy
- This is a call to action for medical providers to critically evaluate patients who exhibit SOREMPs on nocturnal PSG, as it provides a unique opportunity to identify narcolepsy
- Further research is needed to better understand the low referral rate

References: 1. Thorpy MJ, Krieger AC. Sleep Med. 2014;15:502-7. 2. Andlauer O, et al. JAMA Neurol. 2013;70:891-902. 3. Cairns A, Bogan R. Sleep Med. 2017;32:150-6.

^aBased on recommendation for further evaluation by the automated SOREMP notification program.

Figure 4. SOREMPs Occurred More Frequently Among Patients With Lower AHI

The inverse correlation between apnea hypopnea index and occurrence of SOREMPs aligns

with research demonstrating that SOREMPs during **nocturnal** PSG almost always indicate

MSLT, multiple sleep latency test; SOREMP, sleep onset rapid eye movement period.

AHI, apnea hypopnea index; SOREMP, sleep onset rapid eye movement period.

underlying narcolepsy and are not caused by sleep apnea²

50

Support and Acknowledgments: This study was supported by Jazz Pharmaceuticals. Under the direction of the authors, Hannah K. Ritchie, PhD of Peloton Advantage, LLC, an OPEN Health company, provided medical writing and editorial support for this poster, which was funded by Jazz Pharmaceuticals.

Disclosures: A Cairns is an employee of BioSerenity who during the course of this project received grant funding from Jazz Pharmaceuticals. **R Bogan** has served on the speakers' bureau and participated in advisory boards for Jazz Pharmaceuticals and Harmony Biosciences. **A Zheng** is an employee of Huneo and works with and receives financial support from Jazz Pharmaceuticals on this project. **S Bujanover** is a former full-time employee of Jazz Pharmaceuticals who, in the course of this employment, received stock options exercisable for, and other stock awards of, ordinary shares of Jazz Pharmaceuticals, plc. **P Lillaney and A Friedberg** are full-time employees of Jazz Pharmaceuticals, plc. **J Black** is a part-time employee of Jazz Pharmaceuticals plc.



Scan this code to access this poster online.
This code is not for promotional purposes.