

## What is self-reported data?

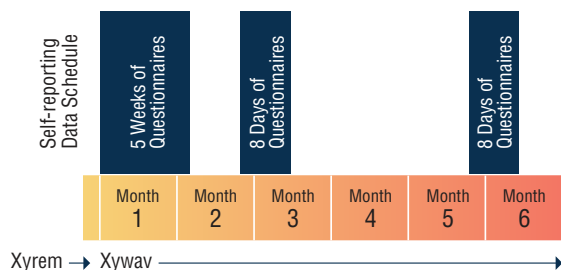
This study is based on self-reported information collected through participant answers to web-based questionnaires.

The purpose of this study is to observe the dosing practices during your transition, learn about the safety and tolerability of the medication, and overall treatment satisfaction.

The information that will be collected will cover the following topics:

- Demographic information
- Brief medical history
- Transition expectations
- Dosing practices
- Narcolepsy symptoms
- Elements of tolerability
- Treatment satisfaction

During different time points of the study, participants will be asked to submit information on a daily or weekly basis, see the visual below that outlines the self-reporting data schedule.



# TENOR

## Want to learn more?

This is a patient-centric, virtual study examining the impact of transitioning from Xyrem to Xywav in a real-world setting. For more details about this study or to enroll, please visit [www.tenorstudy.com](http://www.tenorstudy.com).



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HELP INFORM  
**OTHERS** ABOUT THE  
**TRANSITION** FROM  
**XYREM<sup>®</sup> TO XYWAV<sup>™</sup>**  
BY DOCUMENTING  
YOUR EXPERIENCE



Consider participating in the **TENOR** study for persons with narcolepsy who are transitioning their medication from Xyrem to Xywav

# TENOR

## About the study

The TENOR (Transition Experience of persons with Narcolepsy taking Oxybate in the Real-world) study is a web-based, observational study, for adults (18 years or older) who have been diagnosed with narcolepsy and are transitioning from Xyrem to Xywav.

This study will enroll up to 350 participants. Participation in this study will last approximately 21 weeks, during which participants will be asked to complete web-based study questionnaires.

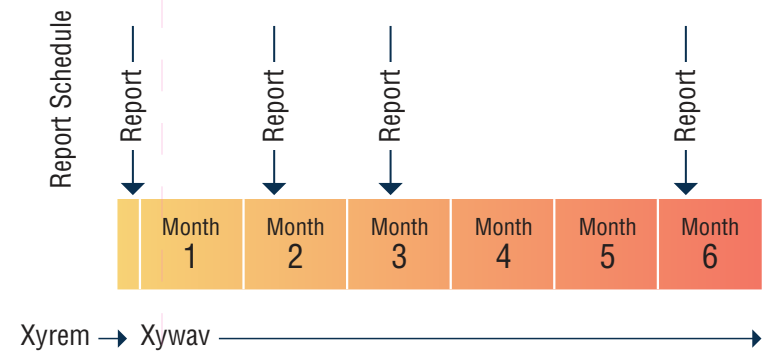
The study was developed with the help of a patient advisory board and input from board certified sleep physicians who specialize in narcolepsy. These questionnaires will collect information on demographics, brief medical history, dosing experience on both Xyrem and Xywav (as well as effectiveness and tolerability of the medication), sleep patterns, and health related quality of life/daily functioning.

## What is an observational study?

An observational study is a research study that collects information from participants to better understand their day to day health experiences. Participants in this study will not receive medication as part of the study or attend in-person study visits. This study will not interfere with the care or treatment your doctor provides.

## Participant reports

As a part of your participation, you will receive reports summarizing your self-reported data throughout the study. At the end of the study, you will receive a consolidated report of your personal data. These reports can be shared with your doctor and/or used to help you visualize your treatment experience.



## Who can participate in the study?

You may be eligible to participate, if you:

- Are aged 18 years or older
- Have been diagnosed with type 1 or type 2 narcolepsy by a physician
- Are able to read and understand English
- Have reliable access to a computer or smart phone with internet connection
- Can commit to the time and study procedures
- Are not a current Jazz Pharmaceuticals employee or an immediate family member of a Jazz employee

## Will I be compensated?

If you qualify, you will be compensated for your time during the study period.

## Why is this study important?

If you take part in the study, you may help patients, doctors, and researchers better understand how narcolepsy treatments affect people with narcolepsy type 1 (narcolepsy with cataplexy) or narcolepsy type 2 (narcolepsy without cataplexy type). By learning more about your experience, it may be possible to improve future treatment options.

**TENOR**