



Patient Advocate University

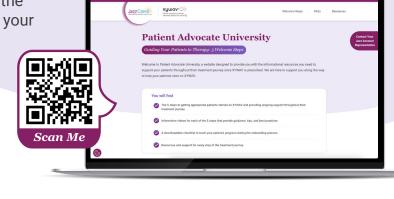
Guiding Your Patients to Therapy: 5 Welcome Steps

Visit Patient Advocate University

This website is designed to provide you with the informational resources you need to support your patients **after XYWAV** is **prescribed**.

On the website, you will find

Five steps to getting appropriate patients started on XYWAV and providing ongoing support throughout their treatment journey













- Informative videos for each of the 5 steps that provide guidance, tips, and best practices
- A downloadable checklist to track your patient's progress during the onboarding process
- Resources and FAQs for every step of the treatment journey

Refer to the next page to learn how to bookmark the Patient Advocate University website for guick and easy access

INDICATIONS AND USAGE

XYWAV is indicated for the treatment of cataplexy or excessive daytime sleepiness (EDS) in patients 7 years of age and older with narcolepsy, and for the treatment of idiopathic hypersomnia (IH) in adults.

Important Safety Information

WARNING: CENTRAL NERVOUS SYSTEM DEPRESSION and ABUSE AND MISUSE.

- Central Nervous System Depression
 XYWAV is a CNS depressant. Clinically significant respiratory depression and obtundation may occur in patients treated with XYWAV at recommended doses. Many patients who received XYWAV during clinical trials in narcolepsy and idiopathic hypersomnia were receiving CNS stimulants.
- Abuse and Misuse
 The active moiety of XYWAV is oxybate or gamma-hydroxybutyrate (GHB). Abuse or misuse of illicit GHB, either alone or in combination with other CNS depressants, is associated with CNS adverse reactions, including seizure, respiratory depression, decreases in the level of consciousness, coma, and death.

Because of the risks of CNS depression and abuse and misuse, XYWAV is available only through a restricted program under a Risk Evaluation and Mitigation Strategy (REMS) called the XYWAV and XYREM REMS.

Please see additional Important Safety Information on the next page and full <u>Prescribing Information</u>, including BOXED Warning.



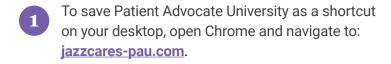


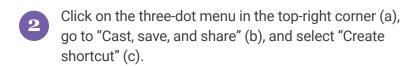
Guidance for Creating a Bookmark

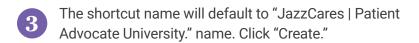
To access Patient Advocate University quickly and easily on a Mac or PC, you can save it as a bookmark. This allows you to reach the site instantly without searching for the link.

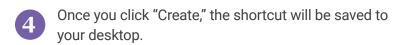
Guidance is provided based on the most used browsers: Google Chrome (PC and Mac), Microsoft Edge (PC), and Safari (Mac).

Google ChromeTM browser - PC/Mac:

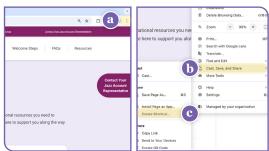


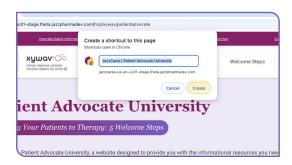














Important Safety Information (cont.)

Contraindications

XYWAV is contraindicated in combination with sedative hypnotics or alcohol and in patients with succinic semialdehyde dehydrogenase deficiency.

Please see additional Important Safety Information on the next page and full <u>Prescribing Information</u>, including BOXED Warning.

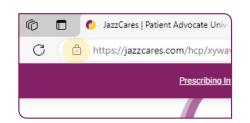




Guidance for Creating a Bookmark (cont.)

Microsoft Edge—PC:

- 1 Open Microsoft Edge and navigate to jazzcares-pau.com.
- Once the website is open, a lock/paddle icon will appear to the left of the address bar.
- Drag and drop the lock/paddle icon to the desktop area to create the desktop shortcut.



Safari-Mac:

- Open Safari and navigate to jazzcares-pau.com.
- Once the website is open, click into the address bar and the Jazz Pharmaceuticals icon will appear.
- Drag and drop the icon to the desktop area to create the desktop shortcut.



Important Safety Information (cont.)

Warnings and Precautions

- CNS Depression: Use caution when considering the concurrent use with other CNS depressants. If concurrent use is required, consider dose reduction or discontinuation of one or more CNS depressants (including XYWAV). Consider interrupting XYWAV treatment if short-term opioid use is required. After first initiating treatment and until certain that XYWAV does not affect them adversely, caution patients against hazardous activities requiring complete mental alertness or motor coordination such as operating hazardous machinery, including automobiles or airplanes. Also caution patients against these hazardous activities for at least 6 hours after taking XYWAV. Patients should be queried about CNS depression-related events upon initiation of XYWAV therapy and periodically thereafter.
- Abuse and Misuse: XYWAV is a Schedule III controlled substance. The rapid onset of sedation, coupled with the amnestic
 features of GHB particularly when combined with alcohol, has proven to be dangerous for the voluntary and involuntary
 user (eg, assault victim).
- Respiratory Depression and Sleep-Disordered Breathing: XYWAV may impair respiratory drive, especially in patients
 with compromised respiratory function. In overdoses of oxybate and with illicit use of GHB, life-threatening respiratory
 depression has been reported. Increased apnea and reduced oxygenation may occur with XYWAV administration in adult
 and pediatric patients. A significant increase in the number of central apneas and clinically significant oxygen desaturation
 may occur in patients with obstructive sleep apnea treated with XYWAV. Prescribers should be aware that sleep-related
 breathing disorders tend to be more prevalent in obese patients, in men, in postmenopausal women not on hormone
 replacement therapy, and among patients with narcolepsy.

Please see additional Important Safety Information on the next page and full <u>Prescribing Information</u>, including BOXED Warning.

It is recommended that your patients enroll in <u>JazzCares</u>—a program sponsored by Jazz Pharmaceuticals, Inc.—to help improve access to Jazz medications for appropriate patients.^a Visit <u>www.jazzcares.com/xywav</u> and click "SIGN NOW" to access the online Patient Authorization Form.



^aEligibility requirements may apply.

If you have questions about Patient Advocate University or want to learn more about JazzCares support offerings for your patients



Visit www.jazzcares.com/hcp/xywav



Call 1-833-533-JAZZ (5299), Monday-Friday, 8 AM-8 PM ET



Click here to contact your Jazz Account Representative

Important Safety Information (cont.)

Warnings and Precautions (cont.)

- Depression and Suicidality: In clinical trials in adult patients with narcolepsy and IH, depression and depressed mood
 were reported in patients treated with XYWAV. In most cases, no change in XYWAV treatment was required. Two suicides
 and two attempted suicides occurred in adult clinical trials with oxybate (same active moiety as XYWAV). One patient
 experienced suicidal ideation and two patients reported depression in a pediatric clinical trial with oxybate. Monitor
 patients for the emergence of increased depressive symptoms and/or suicidality while taking XYWAV, which require
 careful and immediate evaluation.
- Other Behavioral or Psychiatric Adverse Reactions: Monitor patients for impaired motor/cognitive function or the
 emergence of or increase in anxiety and/or confusion. The emergence or increase in the occurrence of behavioral or
 psychiatric events in patients taking XYWAV should be carefully monitored.
- Parasomnias: In clinical trials, parasomnias including sleepwalking were reported in adult patients treated with XYWAV.
 Parasomnias, including sleepwalking, also have been reported in a pediatric clinical trial with sodium oxybate (same active moiety as XYWAV) and in postmarketing experience with sodium oxybate. Episodes of sleepwalking should be fully evaluated and appropriate interventions considered.

Most Common Adverse Reactions

The most common adverse reactions (occurring in ≥5% of XYWAV-treated patients in adult clinical trials in either narcolepsy or IH) were nausea, headache, dizziness, anxiety, insomnia, decreased appetite, hyperhidrosis, vomiting, diarrhea, dry mouth, parasomnia, somnolence, fatigue, and tremor.

In the pediatric clinical trial with XYREM (same active moiety as XYWAV) in patients 7 years of age and older with narcolepsy, the most common adverse reactions (\geq 5%) were nausea (20%), enuresis (19%), vomiting (18%), headache (17%), weight decreased (13%), decreased appetite (9%), dizziness (8%), and sleepwalking (6%). The safety profile in pediatric patients with XYWAV is expected to be similar to that of adult patients treated with XYWAV and to that of pediatric patients treated with XYREM.

Please see full <u>Prescribing Information</u>, including BOXED Warning.

For more information about XYWAV, visit www.xywavhcp.com.

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