



Ryan

Age: 42

Occupation:
Physical therapist

Diagnosis: Narcolepsy without cataplexy (narcolepsy type 2; diagnosed 9 years ago)

Reason for visit:
Routine 6-month visit
• *Ongoing EDS*

Ongoing Symptoms

- Feels “exhausted all the time”; lapses into drowsiness and sleep that affect functioning at work and during other daily activities
 - ESS score of 15

Clinical History

- OSA (diagnosed 15 years ago)
 - RDI 19.9, minimum SpO2 87% on PSG
- Narcolepsy without cataplexy (narcolepsy type 2; diagnosed 9 years ago)
 - Mean sleep latency 0.8 min and 4 SOREMPs on MSLT at diagnosis
 - RDI 3.9 on PSG with CPAP
 - No evidence of other primary sleep disorders on PSG or during clinical interview
 - No evidence of cataplexy during clinical interview
 - ESS score of 17 at diagnosis

Previous Medications

- Wake-promoting agents
- Stimulants

Current Therapy

- CPAP (initiated 15 years ago)

Treatment Decision

- Initiated WAKIX to treat ongoing EDS in narcolepsy

CPAP, continuous positive airway pressure; EDS, excessive daytime sleepiness; ESS, Epworth Sleepiness Scale; MSLT, Multiple Sleep Latency Test; OSA, obstructive sleep apnea; PSG, polysomnogram; RDI, respiratory disturbance index; SOREMP, sleep-onset REM period; SpO2, oxygen saturation.

Based on an actual patient case provided by Gerard Meskill, MD, FAASM.

Indications and Usage

- WAKIX is indicated for the treatment of excessive daytime sleepiness (EDS) or cataplexy in adult patients with narcolepsy.

Important Safety Information

Contraindications

- WAKIX is contraindicated in patients with known hypersensitivity to pitolisant or any component of the formulation. Anaphylaxis has been reported. WAKIX is also contraindicated in patients with severe hepatic impairment.

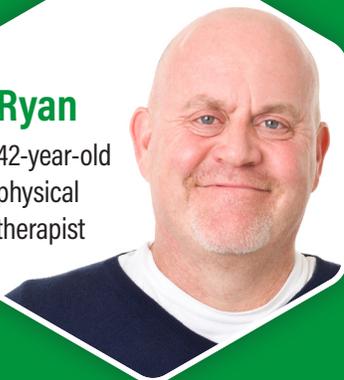
Warnings and Precautions

- WAKIX prolongs the QT interval; avoid use of WAKIX in patients with known QT prolongation or in combination with other drugs known to prolong the QT interval. Avoid use in patients with a history of cardiac arrhythmias, as well as other circumstances that may increase the risk of the occurrence of torsade de pointes or sudden death, including symptomatic bradycardia, hypokalemia or hypomagnesemia, and the presence of congenital prolongation of the QT interval.

Wakix[®]
pitolisant tablets

Ryan

42-year-old
physical
therapist

**Why WAKIX?**

- First-in-class histaminergic treatment
- Significantly reduced EDS vs placebo in clinical studies
- Convenient once-daily morning dosing

Setting Patient Expectations**Ryan was advised:**

- Symptoms may improve at different times or rates when taking WAKIX; it may take up to 8 weeks for some patients to achieve a clinical response
- WAKIX should be taken once daily in the morning upon waking
- WAKIX should be individually titrated to the effective dosage
- WAKIX is *not* a stimulant
- WAKIX is *not* a controlled substance

WAKIX Titration and Administration

- WAKIX was initiated at a dosage of 8.9 mg once daily and titrated weekly to the maximum recommended dosage of 35.6 mg once daily by Week 3
 - Administered once daily in the morning upon waking

Clinical Outcome

- At his 6-week follow-up appointment, Ryan reported a reduction in EDS at a stable dosage of 35.6 mg once daily
 - ESS score of 12
 - Reported feeling less likely to doze or fall asleep during afternoon activities



After initiating treatment with WAKIX, it's important to regularly assess patients for symptom improvement and tolerability

Not all patients respond equally to WAKIX. Individual results may vary.

Important Safety Information**Warnings and Precautions**

- The risk of QT prolongation may be greater in patients with hepatic or renal impairment due to higher concentrations of pitolisant; monitor these patients for increased QTc. Dosage modification is recommended in patients with moderate hepatic impairment and moderate or severe renal impairment (see full prescribing information). WAKIX is not recommended in patients with end-stage renal disease (ESRD).

Adverse Reactions

- In the placebo-controlled clinical trials conducted in patients with narcolepsy with or without cataplexy, the most common adverse reactions ($\geq 5\%$ and twice placebo) for WAKIX were insomnia (6%), nausea (6%), and anxiety (5%). Other adverse reactions that occurred at $\geq 2\%$ and more frequently than in patients treated with placebo included headache, upper respiratory infection, musculoskeletal pain, heart rate increased, hallucinations, irritability, abdominal pain, sleep disturbance, decreased appetite, cataplexy, dry mouth, and rash.

Wakix[®]
pitolisant tablets

Why WAKIX[®] (pitolisant)?



First and only **histaminergic** treatment for EDS or cataplexy in narcolepsy



Not a controlled substance



Significantly reduced EDS and **significantly fewer cataplexy attacks** versus placebo in clinical studies



Established safety and tolerability profile in clinical studies



Convenient **once-daily** morning dosing

Important Safety Information

Drug Interactions

- Concomitant administration of WAKIX with strong CYP2D6 inhibitors increases pitolisant exposure by 2.2-fold. Reduce the dose of WAKIX by half.
- Concomitant use of WAKIX with strong CYP3A4 inducers decreases exposure of pitolisant by 50%. Dosage adjustments may be required (see full prescribing information).
- H₁ receptor antagonists that cross the blood-brain barrier may reduce the effectiveness of WAKIX. Patients should avoid centrally acting H₁ receptor antagonists.
- WAKIX is a borderline/weak inducer of CYP3A4. Therefore, reduced effectiveness of sensitive CYP3A4 substrates may occur when used concomitantly with WAKIX. The effectiveness of hormonal contraceptives may be reduced when used with WAKIX and effectiveness may be reduced for 21 days after discontinuation of therapy.

Use in Specific Populations

- WAKIX may reduce the effectiveness of hormonal contraceptives. Patients using hormonal contraception should be advised to use an alternative non-hormonal contraceptive method during treatment with WAKIX and for at least 21 days after discontinuing treatment.

- There is a pregnancy exposure registry that monitors pregnancy outcomes in women who are exposed to WAKIX during pregnancy. Patients should be encouraged to enroll in the WAKIX pregnancy registry if they become pregnant. To enroll or obtain information from the registry, patients can call [1-800-833-7460](tel:1-800-833-7460).
- The safety and effectiveness of WAKIX have not been established in patients less than 18 years of age.
- WAKIX is extensively metabolized by the liver. WAKIX is contraindicated in patients with severe hepatic impairment. Dosage adjustment is required in patients with moderate hepatic impairment.
- WAKIX is not recommended in patients with end-stage renal disease. Dosage adjustment of WAKIX is recommended in patients with moderate or severe renal impairment.
- Dosage reduction is recommended in patients known to be poor CYP2D6 metabolizers; these patients have higher concentrations of WAKIX than normal CYP2D6 metabolizers.

To report suspected adverse reactions, contact Harmony Biosciences at [1-800-833-7460](tel:1-800-833-7460) or FDA at [1-800-FDA-1088](tel:1-800-FDA-1088) or www.fda.gov/medwatch.

Visit WAKIXhcp.com to learn more and download the Prescription Referral Form



WAKIX is a registered trademark of Bioprojet Europe, Ltd.
Harmony Biosciences name and logo are registered trademarks.
© 2021 Harmony Biosciences. All rights reserved.
US-WAK-2100327/Nov 2021

