

INDICATION

Qelbree is indicated for the treatment of ADHD in adults and pediatric patients 6 years and older.

IMPORTANT SAFETY INFORMATION**WARNING: SUICIDAL THOUGHTS AND BEHAVIORS**

In clinical studies, higher rates of suicidal thoughts and behaviors were reported in patients with ADHD treated with Qelbree than in patients treated with placebo. Closely monitor all Qelbree-treated patients for clinical worsening and for emergence of suicidal thoughts and behaviors.



Please see full Important Safety Information on page 3.

REAL PATIENT PROFILE—EMMETT*

High School Student

THEN

**Elementary school student—diagnosed with ADHD in 2017**

- Prescribed stimulants
- In 2021, when Emmett was 12 years old, he was prescribed Qelbree

NOW

**2025: High school student—Age 17; has been taking Qelbree for over 3 years...**

Abbreviations: ADHD, attention-deficit/hyperactivity disorder; HCP, healthcare professional.

*Real patient compensated for his time.

CONTRAINDICATIONS

- Concomitant administration of a monoamine oxidase inhibitor (MAOI), or dosing within 14 days after discontinuing an MAOI, because of an increased risk of hypertensive crisis
- Concomitant administration of sensitive CYP1A2 substrates or CYP1A2 substrates with a narrow therapeutic range



Please see full Important Safety Information on page 3.

REAL PATIENT PROFILE—EMMETT*

High School Student



Mom: Emmett consistently stays on task

“The past 3 years, I would say things haven’t changed much, which is great... continuing the medication has been helpful for staying on task...the medication is helping him be consistent...there aren’t any ups and downs, highs and lows with anything...the treatment stays quite consistent...”



Grandmother: Emmett participates more with the family

“One of the biggest changes I’ve noticed is that he’s just part of us. He’s one of us...the distractions have minimized, so, he has been able to participate in different things, where before, he was living in his own little world...”



Emmett: I’m now focusing on my future

“I’m looking forward to graduating high school, going to college...studying architecture and...getting a job that works in that sort of field. I’m glad I did something about my ADHD, because if not, I would have been much less focused in class...I do think that if I wasn’t on Qelbree, it would have been much different.”

SEE EMMETT’S FULL STORY HERE:



THEN: The start of Emmett’s Qelbree journey.



NOW: Where Emmett is today on his Qelbree journey.

INDICATION

Qelbree® (viloxazine extended-release capsules) is indicated for the treatment of ADHD in adults and pediatric patients 6 years and older.

IMPORTANT SAFETY INFORMATION

WARNING: SUICIDAL THOUGHTS AND BEHAVIORS

In clinical studies, higher rates of suicidal thoughts and behaviors were reported in patients with ADHD treated with Qelbree than in patients treated with placebo. Closely monitor all Qelbree-treated patients for clinical worsening and for emergence of suicidal thoughts and behaviors.

CONTRAINdications

- Concomitant administration of a monoamine oxidase inhibitor (MAOI), or dosing within 14 days after discontinuing an MAOI, because of an increased risk of hypertensive crisis
- Concomitant administration of sensitive CYP1A2 substrates or CYP1A2 substrates with a narrow therapeutic range

WARNINGS & PRECAUTIONS

- Suicidal thoughts and behaviors:* Closely monitor all Qelbree-treated patients for clinical worsening and emergence of suicidal thoughts and behaviors, especially during the initial few months of drug therapy, and at times of dosage changes
- Heart rate, blood pressure increases:* Qelbree can cause an increase in diastolic blood pressure and heart rate. Assess these measures prior to starting therapy, following increases in dosage, and periodically during therapy
- Activation of mania or hypomania:* Noradrenergic drugs may induce a manic or mixed episode in patients with bipolar disorder. Prior to initiating treatment with Qelbree, screen patients to determine if they are at risk for bipolar disorder. Screening should include a detailed psychiatric history, including a personal or family history of suicide, bipolar disorder, and depression
- Somnolence and fatigue:* Patients should not perform activities requiring mental alertness, such as operating a motor vehicle or hazardous machinery, due to potential somnolence (including sedation or lethargy) and fatigue, until they know how they will be affected by Qelbree

ADVERSE REACTIONS

The most common adverse reactions ($\geq 5\%$ and at least twice the rate of placebo for any dose) in patients 6 to 17 years were somnolence, decreased appetite, fatigue, nausea, vomiting, insomnia, and irritability, and in adults, insomnia, headache, somnolence, fatigue, nausea, decreased appetite, dry mouth, and constipation.

PREGNANCY

There is a pregnancy exposure registry that monitors pregnancy outcomes in women exposed to Qelbree during pregnancy. Healthcare providers are encouraged to register patients by calling the National Pregnancy Registry for Psychiatric Medications at 1-866-961-2388 or by visiting www.womensmentalhealth.org/preg.

Please see full [Prescribing Information](#), including Boxed Warning.



Qelbree is a registered trademark of Supernus Pharmaceuticals, Inc.
©2025 Supernus Pharmaceuticals, Inc. All rights reserved. QBE.2025-0248