A MIGRAINE PORTFOLIO PRESENTATION INTRODUCING PRE-HEADACHE CLINICAL EVIDENCE FOR UBRELVY®

Tuesday, April 16, 2024 at 6:30 PM CT

Las Brisas
4701 112th Street, Lubbock, TX

Please RSVP using the link or QR code at left:
https://migrainelive.com/register/195839
For questions, contact your AbbVie Representative
Megan Chambless Phone: 806-675-6096 Email: megan.chambless@abbvie.com

PRESENTED BY:

Shiraz Yazdani, MD
Pain Medicine
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INDICATION
UBRELVY® (ubrogepant) is indicated for the acute treatment of migraine with or without aura in adults. UBRELVY is not indicated for the preventive treatment of migraine.

IMPORTANT SAFETY INFORMATION
CONTRAINDICATIONS
Drug Interactions: UBRELVY is contraindicated with concomitant use of strong CYP3A4 inhibitors (eg, ketoconazole, itraconazole, clarithromycin).
Hypersensitivity Reactions: UBRELVY is contraindicated in patients with a history of serious hypersensitivity to ubrogepant or any ingredient of the product. Cases, including anaphylaxis, dyspnea, facial or throat edema, rash, urticaria, and pruritus, have been reported. Hypersensitivity reactions can occur minutes, hours, or days after administration. Most reactions were not serious, and some led to discontinuation. If a serious or severe reaction occurs, discontinue UBRELVY and institute appropriate therapy.
ADVERSE REACTIONS
The most common adverse reactions were nausea (4% vs 2% placebo) and somnolence (3% vs 1% placebo).

Please see accompanying full Prescribing Information or visit https://www.rxabbvie.com/pdf/ubrelvy_pi.pdf.

INDICATION
QULIPTA® (atogepant) is indicated for the preventive treatment of migraine in adults.

IMPORTANT SAFETY INFORMATION
CONTRAINDICATIONS
QULIPTA is contraindicated in patients with a history of hypersensitivity to atogepant or any of the components of QULIPTA.
WARNINGS AND PRECAUTIONS
Cases, including anaphylaxis, dyspnea, rash, pruritus, urticaria, and facial edema, have been reported with use of QULIPTA. Hypersensitivity reactions can occur days after administration. If a hypersensitivity reaction occurs, discontinue QULIPTA and institute appropriate therapy.
ADVERSE REACTIONS
The most common adverse reactions (at least 4% and greater than placebo) are nausea, constipation, and fatigue/somnolence.
Dosage form and strengths: QULIPTA is available in 10 mg, 30 mg, and 60 mg tablets.
Please see accompanying full Prescribing Information or visit https://www.rxabbvie.com/pdf/qulipta_pi.pdf.

The QULIPTA pivotal trials did not allow the use of concomitant medications that act on the CGRP pathway. The recently updated UBRELVY PI indicates that no significant pharmacokinetic interactions were observed when UBRELVY was co-administered with QULIPTA.

This promotional event is brought to you by AbbVie and is not certified for continuing medical education.
The speaker is a paid consultant presenting on behalf of AbbVie and the information being presented is consistent with FDA guidelines. This event is conducted in accordance with the PhRMA Code on Interactions with Healthcare Professionals and is limited to invited healthcare professionals (HCPs). Attendance by guests or spouses is not appropriate. It is AbbVie's policy to include only those healthcare professionals involved in patient care consistent with our product indication(s).
The cost of meals and refreshments provided to U.S. HCPs may be subject to public disclosure. AbbVie's disclosure will allocate the cost of meals and refreshments equally across all attendees regardless of actual consumption. AbbVie abides by applicable federal and state laws, which prohibit or limit the ability of government employees and certain healthcare professionals to accept items of value from AbbVie. Please comply with applicable law.

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