

## PREVENTS MIGRAINE DAYS AND KEEPS THEM AWAY OVER TIME

QULIPTA<sup>™</sup> is a pill that significantly reduced monthly migraine days across 12 weeks in clinical studies

## **EXPLORE THE POSSIBILITY AT QULIPTA.COM**

LEARN MORE ABOUT HOW YOU MAY SAVE. TEXT ENROLL TO 785478

#### WHAT IS QULIPTA™?

QULIPTA (atogepant) is a prescription medicine used for the preventive treatment of episodic migraine in adults.

#### **IMPORTANT SAFETY INFORMATION**

Before taking QULIPTA, tell your healthcare provider about all your medical conditions, including if you:

- Have kidney problems or are on dialysis
- Have liver problems
- Are pregnant or plan to become pregnant. It is not known if QULIPTA will harm your unborn baby
- Are breastfeeding or plan to breastfeed. It is not known if QULIPTA passes into your breast milk.
   Talk to your healthcare provider about the best way to feed your baby while taking QULIPTA

#### Please see the Brief Summary of the full Patient Information on the following page.

QULIPTA™ and its design are trademarks of Allergan Pharmaceuticals International Limited, an AbbVie company. © 2021 AbbVie. All rights reserved. US-QULI-210056 12/21 Tell your healthcare provider about all the medicines you take, including prescription and over-the-counter medicines, vitamins, and herbal supplements. QULIPTA may affect the way other medicines work, and other medicines may affect how QULIPTA works. Your healthcare provider may need to change the dose of QULIPTA when taken with certain other medicines.

**The most common side effects of QULIPTA** are nausea, constipation, and fatigue. These are not all the possible side effects of QULIPTA.

You are encouraged to report negative side effects of prescription drugs to the FDA. Visit www.fda.gov/medwatch or call 1-800-FDA-1088.

If you are having difficulty paying for your medicine, AbbVie may be able to help. Visit AbbVie.com/myAbbVieAssist to learn more.



## QULIPTA™ (kew-LIP-tah) (atogepant) tablets, for oral use

#### **CONSUMER BRIEF SUMMARY**

Consult Package Insert for Full Prescribing Information

### **Patient Information**

Read the Patient Information that comes with QULIPTA before you start taking it and each time you get a refill. There may be new information. This brief summary is not comprehensive and does not take the place of talking with your doctor about your medical condition or treatment. For a copy of the full Prescribing Information visit www.QULIPTA.com.

#### What is QULIPTA?

QULIPTA is a prescription medicine used for the preventive treatment of episodic migraine in adults. It is not known if QULIPTA is safe and effective in children.

#### Before you take QULIPTA tell your healthcare provider about all of your medical conditions, including if you:

- have kidney problems or are on dialysis.
- have liver problems.
- are pregnant or plan to become pregnant. It is not known if QULIPTA will harm your unborn baby.
- are breastfeeding or plan to breastfeed. It is not known if QULIPTA passes into your breast milk. Talk to your healthcare provider about the best way to feed your baby while taking QULIPTA.

Tell your healthcare provider about all the medicines you take, including prescription and over-the-counter medicines, vitamins, and herbal supplements. QULIPTA may affect the way other medicines work, and other medicines may affect how QULIPTA works. Your healthcare provider may need to change the dose of QULIPTA when taken with certain other medicines.

Especially tell your healthcare provider if you take any of the following, as your healthcare provider may need to change the dose of QULIPTA:

<ul> <li>ketoconazole or itraconazole</li> </ul>	<ul> <li>rifampin</li> </ul>	<ul> <li>St. John's wort</li> </ul>
<ul> <li>cyclosporine</li> </ul>	<ul> <li>carbamazepine</li> </ul>	<ul> <li>efavirenz</li> </ul>
<ul> <li>clarithromycin</li> </ul>	<ul> <li>phenytoin</li> </ul>	<ul> <li>etravirine</li> </ul>

Keep a list of medicines you take to show to your healthcare provider or pharmacist when you get a new medicine.

#### How should I take QULIPTA?

- Take QULIPTA by mouth 1 time each day with or without food.
- Take QULIPTA exactly as your healthcare provider tells you to take it.

#### What are the possible side effects of QULIPTA?

The most common side effects of QULIPTA include: nausea, constipation, and fatigue.

These are not all of the possible side effects of QULIPTA. Call your doctor for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088.

#### How should I store QULIPTA?

• Store QULIPTA at room temperature between 68°F to 77°F (20°C to 25°C).

Keep QULIPTA and all medicines out of the reach of children.

effective use of QULIPTA. Medicines are sometimes prescribed for purposes other than those listed in a Patient Information leaflet. Do not use QULIPTA for a condition for which it was not prescribed. Do not give QULIPTA to other people, even if they have the same symptoms you have. It may harm them. You can ask your pharmacist or healthcare provider for

General information about the safe and

information about QULIPTA that is written for health professionals.

#### What are the ingredients in QULIPTA?

#### Active ingredient: atogepant

Inactive ingredients: colloidal silicon dioxide, croscarmellose sodium, mannitol, microcrystalline cellulose, polyvinylpyrrolidone vinyl acetate copolymer, sodium chloride, sodium stearyl fumarate, and vitamin E polyethylene glycol succinate.

#### Manufactured by:

Forest Laboratories Ireland Ltd. Dublin. Ireland

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# **Doctor on Call**



#### Janet, a 37-year-old female legal assistant and married mother of three who lives in Charlotte writes:

I've been suffering from migraine for most of my life. When I was a kid, sitting with my brothers in the backseat for a summertime family drive to the shore was torture. Aside from enduring their horseplay, almost immediately I would become carsick, and this typically was followed by an everescalating headache that ruined any chance I had of enjoying myself once we finally arrived at the beach.

Now, thanks to the pandemic, I've been working almost entirely from home for most of the past two years, trying to do my job while simultaneously supervising three young children - my oldest is four, very hyperactive and especially demanding of my attention - and that chaotic environment has combined with long hours spent staring at the computer screen to destabilize my migraine. I wake up every day with a headache, and by the time evening rolls around I often am too sick to sit down with my family for dinner and instead just go to bed. On a scale of 1 to 10, the quality of my day-to-day life is subzero.

I asked my physician to assist me in seeking accommodations at work from my employer that would allow me to work part-time, and I even discussed with him the option of using FMLA to stop working entirely for a few months and to my amazed disappointment, he refused to help. He told me flat out that he would not complete any forms or write a specific letter.

Angry with his refusal, I told him precisely what I thought of him and left his office. I'll not be going back. What's the problem with this guy?

Steaming in Carolina

#### The Doctor's Reply:

Janet,

Thank you for identifying a problem which has vexed healthcare providers (HCPs) and their migraine patients for a long, long time.

By its very definition, migraine at least intermittently reduces one's ability to perform routine activities of daily living, and in that sense it is a medical disorder which may convey disability. Unlike a fractured bone or an infarcting heart, however, migraine headache is a *subjective* disorder, and its diagnosis is made not by an x-ray or electrocardiogram but on the basis of history provided by the patient. To measure *objectively* the degree of disability conveyed by a subjective disorder is no easy task...if not impossible. In addition to the history provided by the patient, many instruments have been devised for the purpose of placing a numeric value on migraine burden and migraine-related disability, but in the end, as with the patient's history itself, whatever result is obtained relies solely upon information provided by the patient.

What happens when an HCP agrees to support a disability claim made on the basis of headache? First, such support typically will require time and effort expended by the HCP - letters written, forms filled out, telephone conversations with attorneys, etc. - and rarely do providers charge for that time and effort. Second, the information requested from the provider often requires an objective determination: How long can a patient be expected to stand, how often to lift up to 50 pounds repeatedly or how quickly to press a foot pedal during an eight hour workday? How many hours per week or days per month will the patient be incapacitated by headache to the point that he or she cannot perform at the expected level? If the requested accommodations are granted, by how many hours per month will, say, eliminating fluorescent lighting at the office, providing more frequent breaks during the day or permitting the patient to work exclusively from home restore productivity otherwise lost to acute headache?

No provider is blessed with a third eye, with the inherent ability to gauge precisely how much pain a migraine patient is experiencing and how much disability that pain conveys. Nor do we possess a crystal ball with which we can unerringly predict the migrainous elements of an individual's future. For assessing migraine burden we simply rely upon the words of the patient, and those words we record in the clinical notes of an electronic medical record. As for predicting, we offer treatment, and we hope for the best. Some providers nevertheless acquiesce to the patient's request and agree to fill out the form, write the letter, talk to the attorney, etc. For the provider involved, this choice can serve as a path to frustration, as the realization dawns that this financially uncompensated time and effort could have been spent providing clinical care, teaching or performing research intended to raise the existing standard of care (or perhaps spent attending to one's own family, waiting perhaps not so patiently at home). Others simply say no, a response likely to frustrate the patient and possibly damage the therapeutic alliance. A compromise solution is to advise patients that while your policy is neither to support nor obstruct any type of disability claim made on the basis of headache, you will send copies of your relevant clinic notes to whatever source the patient designates. Sometimes this is wellreceived, sometimes not.

If you as the patient are denied what you request, it is perfectly reasonable to ask for an explanation. If the explanation seems to you insufficient or, worse, if none is offered, it may be time to move on to another provider. There is an art to practicing medicine, and not all providers are equally adept artists.

The issue you raise, Janet, is part and parcel of a much greater question: how do a patient and provider develop and maintain a healthy and mutually acceptable therapeutic alliance? How much should a patient expect from a provider? How does a provider care for a patient with skill and compassion but avoid personal or professional compromise? There is no simple answer, and as health care evolves the "correct" answer is an evermoving target.

