



HARMONY BIOSCIENCES ANNOUNCES POSITIVE TOPLINE DATA FROM PHASE 2 SIGNAL DETECTION STUDY EVALUATING PITOLISANT IN ADULT PATIENTS WITH MYOTONIC DYSTROPHY TYPE 1

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Clinically meaningful improvements demonstrated in excessive daytime sleepiness and fatigue

PLYMOUTH MEETING, Pa., Dec. 7, 2023 /PRNewswire/ -- Harmony Biosciences Holdings, Inc. ("Harmony" or the "Company") (Nasdaq: HRMY) today announced positive topline results from its Phase 2 signal detection study evaluating the safety and efficacy of pitolisant in adult patients with myotonic dystrophy type 1 (DM1).



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A clinically meaningful improvement was demonstrated on the primary efficacy endpoint, the change from baseline to the end of the double-blind period for excessive daytime sleepiness (EDS) as measured by the Daytime Sleepiness Scale (DSS). A clinically meaningful improvement was also demonstrated for fatigue, a secondary efficacy endpoint measured by the Fatigue Severity Scale (FSS). EDS and fatigue occur in up to 80-90% of patients with DM1 and impact daily functioning as much as muscular symptoms (myotonia and muscle weakness).

"These strong topline results add to the body of evidence supporting the effectiveness of pitolisant for improving EDS," said Jeffrey M. Dayno, M.D., President and CEO of Harmony Biosciences. "In addition, a positive signal for pitolisant has been demonstrated for fatigue, suggesting it could be a potential new treatment option for this symptom as well. We are encouraged by these results, which underscore our commitment to translating innovative science into therapeutic possibilities for patients living with unmet medical needs."

Topline results include:

- Clinically meaningful improvements were demonstrated in EDS as measured by DSS, Epworth Sleepiness Scale (ESS), and Clinical Global Impression of Severity (CGI-S) of EDS.

On DSS, the mean change from baseline to end of double-blind period was -2.5 and -1.0 for the higher dose and lower dose pitolisant treatment groups, respectively, compared to -0.2 for placebo.

On ESS, the mean change from baseline to end of double-blind period was -4.88 for the higher dose pitolisant treatment group compared to -0.10 for placebo.

On CGI-S, the mean change from baseline to end of double-blind period was -0.9 for the higher dose pitolisant treatment group compared to -0.1 for placebo.

- Clinically meaningful improvements were demonstrated for fatigue as measured by the Fatigue Severity Scale (FSS).
On FSS, the mean change from baseline to end of double-blind period was -0.86 and -0.36 for the higher dose and lower dose pitolisant treatment groups, respectively, compared to -0.13 for placebo.
- A clear and consistent dose-response was demonstrated with the higher dose pitolisant group showing a greater response than the lower dose group across the study endpoints.
- Safety and tolerability profile in adult patients with DM1 was consistent with the established safety profile of pitolisant with no new safety signals detected and no serious adverse events reported.

"EDS is the most common non-muscular complaint in patients living with DM1, being present in the majority of patients," said Dr. Jean-Denis Brisson, Centre Intégré Universitaire De Santé Et De Services Sociaux du Saguenay-Lac-Saint-Jean. "The positive signals from this Phase 2 study are very encouraging, and support pitolisant as a potential treatment option for DM1."

This Phase 2 signal detection study was a randomized, double-blind, placebo-controlled study in adults ages 18-65 with DM1. This study was not powered to demonstrate statistical significance. Thirty patients were randomized at baseline to receive lower dose pitolisant, higher dose pitolisant, or placebo in a 1:1:1 treatment ratio titrated over three weeks, followed by eight weeks of stable dosing.

"These positive signals for pitolisant are promising news for the DM1 patient community, where 80-90 percent of the approximately 40,000 diagnosed DM1 patients in the U.S. experience significant EDS and fatigue with no FDA-approved treatments available," said Kumar Budur, M.D., M.S., Chief Medical Officer at Harmony Biosciences. "The strong signals for pitolisant reinforce our confidence in it as a potential treatment option worthy of further study in patients with DM1."

The full dataset from this trial will include findings from other secondary outcomes, including the Myotonic Dystrophy Health Index (MDHI), cognitive functions, and the safety and effectiveness of pitolisant from the ongoing Open-Label Extension (OLE) Phase. Additional results from the trial will be shared early next year. Detailed results will be presented at an upcoming medical meeting and will be submitted for publication.

Pitolisant is marketed as WAKIX® in the U.S. and is FDA approved to treat excessive daytime sleepiness or cataplexy in adults with narcolepsy. Pitolisant is not approved for use in patients with DM1 and is currently being evaluated as an investigational agent in this patient population.

About Myotonic Dystrophy Type 1

Myotonic dystrophy Type 1 (DM1) is the most common form of adult-onset muscular dystrophy. It is a genetic disorder inherited in an autosomal-dominant pattern. Latest estimates suggest a prevalence of about one per 2,100 people with the genetic defect for DM1. This equates to about 150,000 people in the U.S. with the genetic defect for DM1. Estimates suggest there are 40,000 people currently diagnosed with DM1 in the U.S., with up to 90% of them reporting EDS and fatigue and over 60% of them experiencing cognitive dysfunction.

About WAKIX® (pitolisant) Tablets

WAKIX, a first-in-class medication, is approved by the U.S. Food and Drug Administration for the treatment of excessive daytime sleepiness or cataplexy in adult patients with narcolepsy and has been commercially available in the U.S. since Q4 2019. It was granted orphan drug designation for the treatment of narcolepsy in 2010, and breakthrough therapy designation for the treatment of cataplexy in 2018. WAKIX is a selective histamine 3 (H₃) receptor antagonist/inverse agonist. The mechanism of action of WAKIX is unclear; however, its efficacy could be mediated through its activity at H₃ receptors, thereby increasing the synthesis and release of histamine, a wake promoting neurotransmitter. WAKIX was designed and developed by Bioprojet (France). Harmony has an exclusive license from Bioprojet to develop, manufacture and commercialize pitolisant in the United States.

Indications and Usage

WAKIX is indicated for the treatment of excessive daytime sleepiness 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.

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 at least 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 tract infection, musculoskeletal pain, heart rate increased, hallucinations, irritability, abdominal pain, sleep disturbance, decreased appetite, cataplexy, dry mouth, and rash.

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. 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.

Please see the [Full Prescribing Information](#) for WAKIX for more information.

To report suspected adverse reactions, contact Harmony Biosciences at 1-800-833-7460 or the FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

About Harmony Biosciences

At Harmony Biosciences, we specialize in developing and delivering treatments for rare neurological diseases that others often overlook. We believe that where empathy and innovation meet, a better life can begin for people living with neurological diseases. Established by Paragon Biosciences, LLC, in 2017 and headquartered in Plymouth Meeting, PA, our team of experts from a wide variety of disciplines and experiences is driven by our shared conviction that innovative science translates into therapeutic possibilities for our patients, who are at the heart of everything we do. For more information, please visit www.harmonybiosciences.com.

Forward Looking Statement

This press release contains forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995. All statements contained in this press release that do not relate to matters of historical fact should be considered forward-looking statements, including statements regarding our product WAKIX. These statements are neither promises nor guarantees, but involve known and unknown risks, uncertainties and other important factors that may cause our actual results, performance or achievements to be materially different from any future results, performance or achievements expressed or implied by the forward-looking statements, including, but not limited to, the following: our commercialization efforts and strategy for WAKIX; the rate and degree of market acceptance and clinical utility of WAKIX, pitolisant in additional indications, if approved, and any other product candidates we may develop or acquire, if approved; our research and development plans, including our development activities with Bioprojet, and plans to explore the therapeutic potential of pitolisant in additional indications; our ongoing and planned clinical trials; the availability of favorable insurance coverage and reimbursement for WAKIX; the timing of and our ability to obtain regulatory approvals for pitolisant for other indications as well as any of our product candidates, including those we are developing with Bioprojet; our failure to achieve the potential benefits of the 2022 LCA with Bioprojet; our estimates regarding expenses,

future revenue, capital requirements and needs for additional financing; our ability to identify additional products or product candidates with significant commercial potential that are consistent with our commercial objectives; our commercialization, marketing and manufacturing capabilities and strategy; significant competition in our industry; our intellectual property position; loss or retirement of key members of management; failure to successfully execute our growth strategy, including any delays in our planned future growth; our failure to maintain effective internal controls; the impact of government laws and regulations; volatility and fluctuations in the price of our common stock; the significant costs and required management time as a result of operating as a public company; the fact that the price of Harmony's common stock may be volatile and fluctuate substantially; statements related to our intended share repurchases and repurchase timeframe and the significant costs and required management time as a result of operating as a public company. These and other important factors discussed under the caption "Risk Factors" in our Annual Report on Form 10-K filed with the Securities and Exchange Commission (the "SEC") on February 21, 2023, and our other filings with the SEC could cause actual results to differ materially from those indicated by the forward-looking statements made in this press release. Any such forward-looking statements represent management's estimates as of the date of this press release. While we may elect to update such forward-looking statements at some point in the future, we disclaim any obligation to do so, even if subsequent events cause our views to change.

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