



NEWS RELEASE

bridgebio shares recurrent event analysis of attribute-cm, demonstrating a 42% reduction by acoramidis on the composite endpoint of all-cause mortality and recurrent cardiovascular-related hospitalization events

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- Acoramidis treatment led to a highly significant reduction in all-cause mortality (ACM) and recurrent cardiovascular-related hospitalizations (CVH) at Month 30 compared with placebo

- Additionally, results from a Phase 3 trial in adults with ATTR-CM conducted in Japan were presented, with no ACM events reported over the 30 month treatment period in participants administered acoramidis

- Greater transthyretin (TTR) stabilization has been associated with improved clinical outcomes for patients, and in ATTRibute-CM, acoramidis, a near-complete stabilizer of TTR, demonstrated a significant impact on mortality, hospitalizations, and quality of life including:

- An early and sustained improvement relative to placebo in time to first event (CVH or ACM) with a separation of event rates starting at Month 3
- A 42% reduction in composite ACM and recurrent CVH events relative to placebo at Month 30
- A 50% reduction in the cumulative frequency of CVH events relative to placebo at Month 30

PALO ALTO, Calif., Sept. 27, 2024 (GLOBE NEWSWIRE) -- BridgeBio Pharma, Inc. (Nasdaq: BBIO) ("BridgeBio" or the "Company"), a commercial-stage biopharmaceutical company focused on genetic diseases, presented a post-hoc



analysis evaluating the effect of acoramidis on the composite endpoint of ACM and recurrent CVH events in its Phase 3 ATTRIBUTE-CM study in ATTR-CM at the Heart Failure Society of America (HFSA) Annual Scientific Meeting 2024. ATTRIBUTE-CM was designed to evaluate the efficacy and safety of acoramidis, an investigational, near-complete, orally-administered, small molecule stabilizer of TTR.

“We are proud to share the results of this post-hoc analysis demonstrating a highly significant reduction in all-cause mortality and the sum of recurrent CV-related hospitalizations in patients with ATTR-CM at 30 Months in the ATTRIBUTE-CM trial,” said Jonathan Fox, M.D., Ph.D., chief medical officer of BridgeBio Cardiorenal. “At BridgeBio, we believe in the importance of both data transparency and continued communication of additional details of this landmark trial, especially to assist physicians choosing ATTR-CM therapies for their patients who rely on their healthcare provider’s recommendations. By conducting and presenting multiple analyses, we hope to both advance the clinical science around ATTR-CM trials and highlight the potential for acoramidis to be a meaningful treatment option for patients.”

The analysis, shared by Daniel Judge, M.D., professor of medicine and cardiology at the Medical University of South Carolina in an oral presentation, included:

- A 42% reduction in composite ACM and recurrent CVH events at 30 months observed with acoramidis treatment compared to placebo by applying a negative binomial regression model (post-hoc) ($p=0.0005$)
- A 42% reduction in the total number of ACM and recurrent CVH events per patient observed over 30 months with acoramidis treatment compared to placebo
- A 30.5% hazard reduction in ACM and recurrent CVH events at 30 months observed with acoramidis treatment compared to placebo by applying the Andersen-Gill model (post-hoc) ($p=0.0008$)

“This post-hoc analysis provides further evidence that near-complete TTR stabilization with acoramidis can improve clinical outcomes for patients with ATTR-CM. The reduction of hospitalizations and all-cause mortality seen in ATTRIBUTE-CM heightens the case for acoramidis as a first-line therapy given its potential to improve the overall quality of life for patients,” said Dr. Judge.

Additionally, during the Annual Scientific Session of the Japanese College of Cardiology (JCC), Jin Endo, M.D., Ph.D., Keio University School of Medicine of Tokyo, Japan presented positive results from the Phase 3 trial of acoramidis in adults with ATTR-CM conducted by Alexion, AstraZeneca Rare Disease in Japan, which has exclusive rights to commercialize acoramidis in Japan. No mortality was reported over the treatment period of 30 months. Results showed consistency with the ATTRIBUTE-CM results, including survival, CVH and other measures of physical function (measured by six-minute walk test) and quality of life (measured by the Kansas City Cardiomyopathy Questionnaire Overall Summary Score) at 30 months.

Based on the positive results from ATTRIBUTE-CM, BridgeBio submitted a New Drug Application to the U.S. Food and Drug Administration, which has been accepted with a PDUFA action date of November 29, 2024, and a Marketing Authorization Application to the European Medicines Agency, with a decision expected in 2025.

About BridgeBio Pharma, Inc.

BridgeBio Pharma Inc. (BridgeBio) is a commercial-stage biopharmaceutical company founded to discover, create, test and deliver transformative medicines to treat patients who suffer from genetic diseases. BridgeBio's pipeline of development programs ranges from early science to advanced clinical trials. BridgeBio was founded in 2015 and its team of experienced drug discoverers, developers, and innovators are committed to applying advances in genetic medicine to help patients as quickly as possible. For more information visit [bridgebio.com](https://www.bridgebio.com) and follow us on [LinkedIn](#), [Twitter](#) and [Facebook](#).

BridgeBio Forward Looking Statements

This press release contains forward-looking statements. Statements in this press release may include statements that are not historical facts and are considered forward-looking within the meaning of Section 27A of the Securities Act of 1933, as amended (the Securities Act), and Section 21E of the Securities Exchange Act of 1934, as amended (the Exchange Act), which are usually identified by the use of words such as "anticipates," "believes," "continues," "estimates," "expects," "hopes," "intends," "may," "plans," "projects," "remains," "seeks," "should," "will," and variations of such words or similar expressions. We intend these forward-looking statements to be covered by the safe harbor provisions for forward-looking statements contained in Section 27A of the Securities Act and Section 21E of the Exchange Act. These forward-looking statements, including statements relating to the impact of acoramidis on clinical outcomes, potential benefits of acoramidis, including its efficacy and potential to improve the quality of life for patients and the potential outcomes and expected timing of regulatory reviews by the U.S. Food and Drug Administration and the European Medicines Agency reflect our current views about our plans, intentions, expectations and strategies, which are based on the information currently available to us and on assumptions we have made. Although we believe that our plans, intentions, expectations and strategies as reflected in or suggested by those forward-looking statements are reasonable, we can give no assurance that the plans, intentions, expectations or strategies will be attained or achieved. Furthermore, actual results may differ materially from those described in the forward-looking statements and will be affected by a number of risks, uncertainties and assumptions, including, but not limited to, initial and ongoing data from our preclinical studies and clinical trials not being indicative of final data, the design and success of ongoing and planned clinical trials, future regulatory filings, approvals and/or sales, the U.S. Food and Drug Administration or such other regulatory agencies not agreeing with our regulatory approval strategies, components of our filings, such as clinical trial designs, conduct and methodologies, or the sufficiency of data submitted, the continuing success of our collaborations, potential volatility in our share price, uncertainty regarding any impacts due to global health emergencies, including delays in regulatory review, manufacturing and supply chain interruptions, adverse effects on healthcare systems and



disruption of the global economy, the impacts of current macroeconomic and geopolitical events, including changing conditions from hostilities in Ukraine and in Israel and the Gaza Strip, increasing rates of inflation and rising interest rates, on our business operations and expectations, as well as those risks set forth in the Risk Factors section of our most recent Annual Report on Form 10-K and our other filings with the U.S. Securities and Exchange Commission. Moreover, we operate in a very competitive and rapidly changing environment in which new risks emerge from time to time. These forward-looking statements are based upon the current expectations and beliefs of our management as of the date of this press release, and are subject to certain risks and uncertainties that could cause actual results to differ materially from those described in the forward-looking statements. Except as required by applicable law, we assume no obligation to update publicly any forward-looking statements, whether as a result of new information, future events or otherwise.

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