BRIDGEBIO

NEWS RELEASE

BridgeBio to Present New Acoramidis Data from Open-Label Extension Analyses in ATTR-CM at ESC Congress 2026

2026-08-24

PALO ALTO, Calif., Aug. 24, 2026 (GLOBE NEWSWIRE) -- BridgeBio Pharma, Inc. (Nasdaq: BBIO) (“BridgeBio” or the “Company”), a biopharmaceutical company focused on developing medicines for genetic conditions, announced today that two oral presentations and three posters on new acoramidis data in individuals with transthyretin amyloid cardiomyopathy (ATTR-CM) will be shared at the European Society of Cardiology (ESC) Congress 2026, taking place in Munich, Germany on August 28-31, 2026. The data will further strengthen the differentiated clinical profile of acoramidis, reinforcing it as the first-line treatment of choice for individuals with ATTR-CM. Acoramidis is the only selective small molecule, orally administered, near-complete (≥90%) transthyretin (TTR) stabilizer.

As part of BridgeBio's partnership with Yale's Cardiovascular Data Science (CarDS) Lab to advance AI networks for earlier detection of ATTR-CM, three additional posters will be presented at the ESC Congress 2026.

Acoramidis is approved as Attruby® by the U.S. FDA and is approved as BEYONTTRA® by the European Medicines Agency (EMA), Japanese Pharmaceuticals and Medical Devices Agency, Swissmedic, the Swiss Agency for Therapeutic Products, the UK Medicines and Healthcare Products Regulatory Agency, and the Brazilian Health Regulatory Agency (ANVISA) with all labels specifying near-complete stabilization of TTR.

Oral Presentations:

Acoramidis Reduces Days Lost to Death and/or Cardiovascular-Related Hospitalization, Preserving Time Alive Outside the Hospital in Participants with ATTR-CM: Results from ATTRibute-CM

Presenter: Richard Wright, M.D., Pacific Heart Institute, U.S.

Date: Sunday, August 30 at 8:15 am CEST

Durable Survival Benefits of Acoramidis over 54 Months in Variant Transthyretin Amyloid Cardiomyopathy (ATTR-CM), Including p.V142I: Interim Findings from ATTRibute-CM and its Open-Label Extension
Presenter: Kevin Alexander, M.D., Stanford University School of Medicine, U.S.
Date: Sunday, August 30 at 10:55 am CEST

Moderated ePosters:

Long-term Improvement in Myocardial Structure and Function in Patients with Transthyretin Amyloid Cardiomyopathy (ATTR-CM) Treated with Acoramidis Compared with a Natural History Cohort
Presenter: Awais Sheikh, MBChB, National Amyloidosis Centre, London, UK
Date: Friday, August 28 at 4:15 pm CEST

Acoramidis Improves Health-Related Quality of Life in Wild-Type and Variant Transthyretin Amyloid Cardiomyopathy: An EQ-5D-5L Subgroup Analysis from ATTRibute-CM
Presenter: Emer Joyce, M.D., Ph.D., The Mater Misericordiae University Hospital, IE
Date: Sunday, August 30 at 3:15 pm CEST

Improvement of Health Status with Acoramidis in Patients with Wild-Type and Variant Transthyretin Amyloid Cardiomyopathy: KCCQ Domains Analysis from the ATTRibute-CM Study
Presenter: Nitasha Sarswat, M.D., University of Chicago Medical Center, U.S.
Date: Sunday, August 30 at 3:15 pm CEST

Yale-Partnered Moderated ePosters:

A Novel AI-Derived Digital Biomarker for Monitoring Disease Progression in ATTR-CM: First-In-Trial Use of a Computer Vision AI-ECG Algorithm within a Phase 3 Pivotal Randomized Controlled Trial
Presenter: Rohan Khera, M.D., Yale School of Medicine, U.S.
Date: Monday, August 31 at 11:15 am CEST

A Fully Decentralized, Patient-Led Digital Registry for ATTR-CM Integrating Multisystem EHR and Wearable Data: The DISCOVER-ATTR Study
Presenter: Aline Pedroso, Ph.D., Yale School of Medicine, U.S.
Date: Monday, August 31 at 1:15 pm CEST

Nationwide U.S. Federated Deployment of Artificial Intelligence for Multimodal Screening of ATTR Cardiomyopathy: First Multicenter Analysis from the TRACE-AI Network
Presenter: Bruno Batinica, MBChB, Yale School of Medicine, U.S.
Date: Sunday, August 30 at 1:35 pm CEST

About Attruby™ (acoramidis)

INDICATION

Attruby is a transthyretin stabilizer indicated for the treatment of the cardiomyopathy of wild-type or variant transthyretin-mediated amyloidosis (ATTR-CM) in adults to reduce cardiovascular death and cardiovascular-related hospitalization.

IMPORTANT SAFETY INFORMATION

Adverse Reactions

Diarrhea (11.6% vs 7.6%) and upper abdominal pain (5.5% vs 1.4%) were reported in patients treated with Attruby versus placebo, respectively. The majority of these adverse reactions were mild and resolved without drug discontinuation. Discontinuation rates due to adverse events were similar between patients treated with Attruby versus placebo (9.3% and 8.5%, respectively).

About BridgeBio

BridgeBio exists to develop transformative medicines for genetic conditions. Millions of people worldwide living with genetic conditions lack treatment options, often because drug development for small patient populations can be commercially challenging. We aim to bridge the gap between advancements in genetic science and meaningful medicines for underserved patient populations. Our decentralized, hub-and-spoke model is designed for speed, precision, and scalability. Autonomous and empowered teams focus on individual conditions, while a central hub provides the clinical, regulatory, and commercial capabilities needed to bring innovation to market. For more information, visit **bridgebio.com** and follow us on **LinkedIn**, **X**, **Facebook**, **Instagram**, **YouTube**, and **TikTok**.

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