NEWS RELEASE

BridgeBio Raises $300 Million Through Partial Capped Monetization of BEYONTTRA® European Royalty

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- BridgeBio has received a $300 million upfront payment, strengthening the Company's balance sheet, and supporting the launch of Attruby® and ongoing late-stage pipeline programs
- Transaction monetizes 60% of BridgeBio's European royalties on the first $500 million of annual BEYONTTRA net sales, with total payments to the investors subject to an initial cap of 1.45x
- In the ATTRibute-CM study, acoramidis demonstrated the most rapid benefit seen in any Phase 3 study of ATTR-CM to date in both ATTRv-CM and ATTRwt-CM patients:
    - In as few as 3 months, the time to first event (ACM or CVH) durably separated relative to placebo
    - 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
- Acoramidis is approved as Attruby by the U.S. FDA and is approved as BEYONTTRA by the European Commission, Japanese Pharmaceuticals and Medical Devices Agency and UK Medicines and Healthcare Products Regulatory Agency

PALO ALTO, Calif., June 30, 2025 (GLOBE NEWSWIRE) -- BridgeBio Pharma, Inc. (Nasdaq: BBIO) ("BridgeBio" or the "Company"), a new type of biopharmaceutical company focused on genetic diseases, today announced it has sold a portion of royalties due to the Company from sales of BEYONTTRA in Europe to HealthCare Royalty ("HCRx") and funds managed by Blue Owl Capital ("Blue Owl") for $300 million. This royalty financing agreement monetizes select anticipated royalties and provides immediate less-dilutive capital to the Company.

"We're excited to partner with HCRx and Blue Owl to strengthen our balance sheet in support of the launch of Attruby and our pipeline of first and best-in-class genetic medicines. This transaction preserves significant upside

for our shareholders, with careful structuring that limits annual as well as total payments made to the royalty investors. This financing highlights the strong early start and large global potential of acoramidis," said Chinmay Shukla, Senior Vice President of Strategic Finance at BridgeBio.

Clarke Futch, Chairman and Chief Executive Officer of HCRx commented:
"We have been following the progress of acoramidis for years and strongly believe in its potential to positively impact the lives of patients living with ATTR-CM. This investment exemplifies HCRx's commitment to supporting innovation in the biopharmaceutical industry and we are pleased to collaborate with BridgeBio on this transaction."

"We are pleased to continue our support of BridgeBio through this royalty monetization transaction," said Sandip Agarwala, Managing Director and Head of Life Sciences at Blue Owl. "This investment reflects our confidence in BEYONTTRA commercial potential and our commitment to providing flexible capital solutions that help advance life-saving therapies."

Under the terms of the agreement, BridgeBio has received $300 million from HCRx and Blue Owl managed funds in exchange for 60% of royalties on the first $500 million of annual BEYONTTRA net sales in Europe. The agreement includes an initial cap of 1.45x. Once the applicable cap is met, no further payments will be owed to the investors.

In March 2024, BridgeBio entered into an exclusive licensing agreement with Bayer Consumer Care AG to commercialize BEYONTTRA in Europe for the treatment for ATTR-CM. To date, BridgeBio has received $210 million in upfront and regulatory milestones, and anticipates receiving a further $75 million in near-term milestone payments, along with tiered royalties starting in the low-30% range on net sales of BEYONTTRA in Europe. Acoramidis is approved in the U.S. as Attruby by the FDA and in Europe as BEYONTTRA by the European Commission. It is also approved as BEYONTTRA by the Japanese Pharmaceuticals and Medical Devices Agency, and UK Medicines and Healthcare Products Regulatory Agency with all labels specifying near-complete stabilization of TTR.

Latham & Watkins LLP served as legal advisor to BridgeBio. Gibson, Dunn & Crutcher LLP served as legal advisor to HCRx and Blue Owl.

About BEYONTTRA

BEYONTTRA is an orally administered near-complete (≥90%) stabilizer of transthyretin (TTR) indicated for the treatment of wild-type or variant transthyretin amyloidosis in adult patients with cardiomyopathy (ATTR-CM). For full prescribing information, please refer to the Summary of Product Characteristics (SmPC) on the Medicines and Healthcare products Regulatory Agency website at **https://products.mhra.gov.uk/**.

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 Pharma, Inc.

BridgeBio Pharma, Inc. (BridgeBio) is a new type of 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** and follow us on **LinkedIn**, **Twitter, Facebook**, and **YouTube**.

About HealthCare Royalty

HealthCare Royalty (HCRx) is a leading royalty acquisition company focused on commercial or near-commercial biopharmaceutical products. With offices in Stamford, Conn., San Francisco, Boston, London and Miami, HCRx has invested $5+ billion in over 90 biopharmaceutical products since inception. For more information, visit **https://www.hcrx.com**. HEALTHCARE ROYALTY® and HCRx® are registered trademarks of HealthCare Royalty Management, LLC.

About Blue Owl

Blue Owl (NYSE: OWL) is a leading asset manager that is redefining alternatives®.

With $273 billion in assets under management as of March 31, 2025, we invest across three multi-strategy platforms: Credit, Real Assets, and GP Strategic Capital. Anchored by a strong permanent capital base, we provide businesses with private capital solutions to drive long-term growth and offer institutional investors, individual investors, and insurance companies differentiated alternative investment opportunities that aim to deliver strong

performance, risk-adjusted returns, and capital preservation. Together with over 1,200 experienced professionals globally, Blue Owl brings the vision and discipline to create the exceptional. To learn more, visit **www.blueowl.com**.

BridgeBio Pharma, Inc. 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," "could," "estimates," "expects," "hopes," "intends," "may," "plans," "potential," "projects," "seeks," "should," "will," and variations of such words or similar expressions. BridgeBio intends 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 regarding the potential commercial performance of BEYONTTRA in Europe and other global markets; the anticipated benefits of the royalty monetization agreement with HCRx and Blue Owl Capital; BridgeBio's expectations for the launch and market uptake of Attruby and BEYONTTRA; the belief in acoramidis' ability to positively impact the lives of patients with ATTR-CM; and the expected use of proceeds to support BridgeBio's pipeline of genetic medicines, reflect BridgeBio's current views about its plans, intentions, expectations, and strategies, which are based on information currently available and assumptions it has made. Although BridgeBio believes that its plans, intentions, expectations, and strategies as reflected in or suggested by these forward-looking statements are reasonable, it can give no assurance that such 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: BridgeBio's dependence on third parties for development or commercialization activities; regulatory authorities requiring additional studies or data to support the continued or expanded commercialization of acoramidis; whether data and results meet applicable regulatory requirements or are sufficient for continued development, review, or approval; and whether other regulatory agencies agree with BridgeBio's strategies or data interpretations. These risks also include impacts from global health emergencies, such as delays in regulatory reviews and other activities, manufacturing and supply chain interruptions, adverse effects on healthcare systems, and disruption of the global economy; and the impacts of macroeconomic and geopolitical events, including changing conditions from hostilities in Ukraine and in Israel and the Gaza Strip, increasing inflation rates, and fluctuating interest rates on BridgeBio's operations and expectations. Additional risks are described in the "Risk Factors" section of BridgeBio's most recent Annual Report on Form 10-K, Quarterly Report on Form 10-Q, and other filings with the U.S. Securities and Exchange Commission. Moreover, BridgeBio operates in a highly competitive and rapidly evolving industry in which new risks may emerge from time to time. These forward-looking statements are based upon the current expectations and beliefs of BridgeBio's 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 these statements. Except as required by

applicable law, BridgeBio assumes no obligation to publicly update any forward-looking statements, whether as a result of new information, future events, or otherwise.

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