

# Biora Therapeutics Progresses Research Collaboration for the BioJet™ Systemic Oral Delivery Platform

January 2, 2024

## BioJet device exceeded performance targets for delivery of collaborator AstraZeneca's molecule in preclinical study

SAN DIEGO, Jan. 02, 2024 (GLOBE NEWSWIRE) -- <u>Biora Therapeutics, Inc</u>. (Nasdaq: BIOR), the biotech company that is reimagining therapeutic delivery, today announced that its BioJet<sup>™</sup> Systemic Oral Delivery Platform has met key performance milestones as part of its research collaboration with AstraZeneca. The BioJet platform is designed to use needle-free, liquid jet injection to deliver drug into the small intestine for systemic absorption.

"In a preclinical study, we assessed the bioavailability of our collaborator's molecule when delivered via our BioJet device in a porcine model, with comparison to subcutaneous administration. BioJet devices were administered endoscopically, which is typical in a porcine model, and released for autonomous activation," said Sharat Singh, PhD, Head of Research at Biora Therapeutics. "We are encouraged by the results, which met our performance targets of greater than 25% bioavailability compared to subcutaneous delivery, and less than 50% coefficient of variation."

"We continue to generate very promising results with our BioJet platform, including those from this study, which are enabling us to progress development under the collaboration," said Adi Mohanty, Chief Executive Officer of Biora Therapeutics. "While many organizations have been working to solve the challenge of oral delivery of large molecules, we believe our approach stands out because of our ability to deliver multi-milligram liquid doses similar to injection, with minimal to no formulation changes."

Biora has previously presented data generated from other molecules with the BioJet platform in animal models, which can be viewed at bioratherapeutics.com/publications.

### About the BioJet<sup>™</sup> Systemic Oral Delivery Platform

Biora's BioJet systemic oral therapeutics platform uses an ingestible capsule for needle-free, oral delivery of large molecules designed to achieve systemic bioavailability and replace injection for better management of chronic diseases.

The BioJet platform uses an ingestible device designed to transit through the digestive system and activate in the small intestine, where liquid jets deliver drug directly into the small intestine for uptake into systemic circulation. The BioJet device is approximately the size of a multivitamin and is designed to autonomously deliver a wide range of large molecules, such as proteins, peptides, and nucleic acids, in liquid formulation at multi-milligram doses, without requiring complex reformulation.

Biora holds a comprehensive patent position for the BioJet systemic oral delivery platform, with approximately 12 issued patents and 29 pending applications that cover its delivery platform and methods for using the platform to treat a disease or condition in a patient using liquid jet delivery of a wide range of drugs.

### **About Biora Therapeutics**

Biora Therapeutics is reimagining therapeutic delivery. By creating innovative smart pills designed for targeted drug delivery to the GI tract, and systemic, needle-free delivery of biotherapeutics, the company is developing therapies to improve patients' lives.

Biora is focused on development of two therapeutics platforms: the <u>NaviCap™ targeted oral delivery platform</u>, which is designed to improve outcomes for patients with inflammatory bowel disease through treatment at the site of disease in the gastrointestinal tract, and the <u>BioJet™ systemic oral</u> <u>delivery platform</u>, which is designed to replace injection for better management of chronic diseases through needle-free, oral delivery of large molecules.

For more information, visit bioratherapeutics.com or follow the company on LinkedIn or Twitter.

#### Safe Harbor Statement or Forward-Looking Statements

This press release contains "forward-looking statements" within the meaning of the "safe harbor" provisions of the Private Securities Litigation Reform Act of 1995, which statements are subject to substantial risks and uncertainties and are based on estimates and assumptions. All statements, other than statements of historical facts included in this press release, including statements concerning the progress and future expectations and goals of our research and development and clinical efforts and research collaboration plans and expectations are forward-looking statements. In some cases, you can identify forward-looking statements by terms such as "may," "might," "will," "objective," "intend," "should," "could," "can," "would," "expect," "believe," "design," "estimate," "predict," "potential," "plan," "target," or the negative of these terms, and similar expressions intended to identify forward-looking statements. These statements reflect our plans, estimates, and expectations, as of the date of this press release. These statements involve known and unknown risks, uncertainties and other factors that could cause our actual results to differ materially from the forward-looking statements expressed or implied in this press release. Such risks, uncertainties, and other factors include, among others, our ability to innovate in the field of therapeutics, our ability to make future filings and initiate clinical trials on expected timelines or at all, our ability to obtain and maintain regulatory approval, clearance, or acceptance of our clinical trials or products on expected timelines or at all, our plans to research, develop, and commercialize new products, the unpredictable relationship between preclinical study results and clinical study results, our expectations regarding allowed patents or intended grants to result in issued or granted patents, our expectations regarding opportunities with current or future pharmaceutical collaborators, our ability to raise sufficient capital to achieve our business objectives, and those risks described in "Risk Factors" and "Management's Discussion and Analysis of Financial Condition and Results of Operations" in our Annual Report on Form 10-K for the year ended December 31, 2022 filed with the SEC and other subsequent documents, including Quarterly Reports, that we file with the SEC.

Biora Therapeutics expressly disclaims any obligation to update any forward-looking statements whether as a result of new information, future events or otherwise, except as required by law.

Investor Contact Chuck Padala Managing Director, LifeSci Advisors IR@bioratherapeutics.com (646) 627-8390

Media Contact media@bioratherapeutics.com