



Biora Therapeutics to Present Clinical Data on Device Function of the NaviCap™ Oral Delivery Platform at Digestive Disease Week 2024

May 6, 2024

SAN DIEGO, May 06, 2024 (GLOBE NEWSWIRE) -- [Biora Therapeutics, Inc.](https://www.bioratherapeutics.com) (Nasdaq: BIOR), the biotech company that is reimagining therapeutic delivery, today announced that it will present clinical data on device function of the NaviCap™ Targeted Oral Delivery Platform at Digestive Disease Week® (DDW), which will take place both virtually and in-person May 18-21, 2024 in Washington, DC.

Details of the presentations are as follows:

Abstract Title: Results of Human Device Function Studies for the NaviCap™ Targeted Oral Delivery Platform in Healthy Volunteers and Patients with UC
Poster No: Su1778
Session Title: IBD: Controlled Clinical Trials in Humans
Session Date & Time: May 19, 2024 from 12:30 PM to 1:30 PM Eastern time
Presenting Author: Shaoying Nikki Lee, PhD, Director, Clinical and Translational Science, Biora Therapeutics, Inc.

Abstracts selected by the American Gastroenterological Association (AGA) for presentation at DDW will be available in a supplement to *Gastroenterology*. All information presented at DDW, including poster presentations, is embargoed until 12:01 AM Eastern on the day of the presentation. The poster presentation will be made available on the [Biora Therapeutics website](https://www.bioratherapeutics.com) following the meeting.

About Digestive Disease Week

Digestive Disease Week® (DDW) is the largest international gathering of physicians, researchers and academics in the fields of gastroenterology, hepatology, endoscopy and gastrointestinal surgery. Jointly sponsored by the American Association for the Study of Liver Diseases (AASLD), the American Gastroenterological Association (AGA), the American Society for Gastrointestinal Endoscopy (ASGE) and the Society for Surgery of the Alimentary Tract (SSAT), DDW is an in-person and online meeting from May 18-21, 2024. The meeting showcases more than 4,400 abstracts and hundreds of lectures on the latest advances in GI research, medicine and technology. More information can be found at www.ddw.org.

About the NaviCap™ Targeted Oral Delivery Platform

[Biora's NaviCap targeted oral therapeutics platform](https://www.bioratherapeutics.com) utilizes a novel approach that could improve patient outcomes by enabling delivery of therapeutics directly to the site of disease, increasing therapeutic levels in tissue while reducing systemic uptake. For the 1.8 million patients in the United States who suffer from inflammatory bowel disease (IBD), existing therapeutics offer less than ideal efficacy, likely because of the challenges with safely achieving sufficient drug levels in the affected tissues. [Research has shown](#) that direct delivery of therapeutics has the potential to improve patient outcomes in IBD.

The NaviCap platform uses an ingestible device [designed for direct delivery of therapeutics](#) to improve treatment of IBD. Once swallowed, Biora's GItrac™ autolocation technology enables the device to autonomously identify targeted locations in the GI tract and release a therapeutic dose of up to 500µl. Studies in healthy volunteers have demonstrated [accurate localization and delivery in a fasted state](#) and demonstrated the device's [ability to function in both fasted and fed states](#), making it potentially the first ingestible therapeutic delivery device that does not require fasting or other food restriction for use. A device function study in participants with active UC also [demonstrated successful device performance in active UC patients](#).

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 clinical-stage [NaviCap™ targeted oral delivery platform](#) 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 preclinical-stage [BioJet™ systemic oral delivery platform](#) 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](https://www.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, preclinical, and clinical trial efforts including our BT-600 clinical trial execution and data timelines, 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," "forward," "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 and execute clinical trials on expected timelines or at all, our ability to obtain and maintain regulatory approval or clearance of our 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 or partners, 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, 2023 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.

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