

# Barinthus Bio Announces Late Breaking Presentation at AASLD 2024 – The Liver Meeting® and Attendance at November Conferences

# Oct 31, 2024

OXFORD, United Kingdom, Oct. 31, 2024 (GLOBE NEWSWIRE) -- Barinthus Biotherapeutics plc (NASDAQ: BRNS), a clinical-stage biopharmaceutical company developing novel immunotherapeutic candidates that guide T cells to control disease, today announced that updated clinical data from its chronic hepatitis B program will be highlighted as a late breaking oral presentation at the American Association for the Study of Liver Diseases (AASLD) – The Liver Meeting® 2024 being held November 15-19, 2024 in San Diego, California and attendance at multiple conferences in November:

#### Medical Congresses

#### AASLD – The Liver Meeting® 2024:

- Late breaking abstract accepted as oral presentation.
- Title: VTP-300 combined with low-dose nivolumab is associated with HBsAg loss in certain chronic hepatitis B participants with HBsAg less than 200 IU/mL: Results from a phase 2b open-label study.

# International Workshop on HBV Cure 2024:

 Formal presentation titled "HBsAg Levels and T Cell Boost in Patients with HBsAg Decline", on November 13, 2024 at 16:50 ET.

# British Society for Immunology (BSI) Immune Therapies Summit:

• Formal presentation titled "SNAP-TI Nanoparticles Induce Antigen-Specific Tolerance", between November 25 and 26, 2024.

#### Investor Banking Conferences

#### Guggenheim's Inaugural Healthcare Innovation Conference:

- Fireside Chat with Bill Enright, Chief Executive Officer, on November 11, 2024 at 14:00 ET.
- Barinthus Bio's management will be available for one-to-one meetings between November 11 and 15, 2024.

#### Jefferies London Healthcare Conference:

• Barinthus Bio's management will be available for one-to-one meetings between November 19 and 21, 2024.

To access recordings and materials associated with the events, please visit: https://investors.barinthusbio.com/events-presentations.

#### About Barinthus Bio

Barinthus Bio is a clinical-stage biopharmaceutical company developing novel immunotherapeutic candidates designed to guide the immune system to overcome chronic infectious diseases and autoimmunity. Helping people living with serious diseases and their families is the guiding principle at the heart of Barinthus Bio. With a focused pipeline built around our proprietary platform technologies, Barinthus Bio is advancing immunotherapeutic product candidates in infectious diseases and autoimmunity, including: VTP-300, that utilizing our ChAdOX/MVA platform designed as a potential component of a functional cure for chronic HBV infection and VTP-1000, utilizing our SNAP-Tolerance Immunotherapy (SNAP-TI) platform and is designed to treat people with celiac disease. Barinthus Bio is also conducting a Phase 1 clinical trial for VTP-850, a second-generation immunotherapeutic candidate designed to treat recurrent prostate cancer. Barinthus Bio's differentiated technology platforms and therapeutic approach, coupled with deep scientific expertise and focus on clinical development, uniquely positions the company to navigate towards delivering treatments that improve the lives of people with chronic infectious diseases and autoimmunity. For more information, visit www.barinthusbio.com.

#### **Forward Looking Statements**

This press release contains forward-looking statements regarding Barinthus Bio within the meaning of the Private Securities Litigation Reform Act of 1995, as amended, which can generally be identified as such by use of the words "may," "will," "plan," "forward," "encouraging," "believe," "potential," "expect," and similar expressions, although not all forward-looking statements contain these identifying words. These forward-looking statements include, without limitation, express or implied statements regarding our future expectations, plans and prospects, including our product development activities and clinical trials, including timing for readouts of any preliminary, interim or final data for any of our programs, our anticipated regulatory filings and approvals, our cash runway, and our ability to develop and advance our current and future product candidates and programs. Any forward-looking statements in this press release are based on our management's current expectations and beliefs and are subject to numerous risks, uncertainties and important factors that may cause actual events or results to differ materially from those expressed or implied by any forward-looking statements releaved, including, without limitation, risks and uncertainties related to the success, cost and timing of our pipeline development activities and planned and ongoing clinical trials, including the risk that the timing for preliminary, interim or final data or initiation of our clinical trials may be delayed, the risk that interim or topline data may not reflect final data or results, our ability to execute on our strategy, regulatory developments, the risk that we may not achieve the anticipated benefits of our pipeline prioritization and corporate restructuring, our ability to fund our operations and access capital, our cash runway, including the risk that our estimate of our cash runway be incorrect, global economic uncertainty, including disruptions in the banking industry, the conflicts in Ukraine, Israel and Gaza, and other ri

Securities and Exchange Commission, including our Annual Report on Form 10-K for the year ended December 31, 2023, our Quarterly Reports on Form 10-Q and Current Reports on Form 8-K. We caution you not to place undue reliance on any forward-looking statements, which speak only as of the date they are made. We expressly disclaim any obligation to publicly update or revise any such statements to reflect any change in expectations or in events, conditions or circumstances on which any such statements may be based, or that may affect the likelihood that actual results will differ from those set forth in the forward-looking statements.

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