Azitra to Participate in Dermatologic Rare Disease Panel at Maxim Growth Summit

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BRANFORD, Conn., Oct. 15, 2025 /PRNewswire/ -- Azitra, Inc. (NYSE American: AZTR), a clinical stage biopharmaceutical company focused on developing innovative therapies for precision dermatology, today announced that Travis Whitfill Ph.D., Co-Founder and Chief Operating Officer will participate in a Dermatologic Rare Disease Panel at the Maxim Growth Summit, which is being held at the Hard Rock Hotel in New York City, October 22-23, 2025.

The panel members will discuss Netherton syndrome and Recessive Epidermolysis Bullosa (RDEB), two categories with novel therapeutic approaches spanning early- to late-stages of development. Azitra's lead program, ATR-12, is currently being investigated in a Phase 1b clinical trial in adult Netherton syndrome patients. ATR-12 is composed of an engineered strain of *S. epidermidis* designed to replace and deliver deficient LEKTI protein when applied topically to Netherton syndrome patients.

The details of the panel are as follows:

Event:	Rare Disease Panel at the Maxim Growth Summit
Date & Time:	Wednesday, October 22, 2025 at 8:30 AM EDT
Location:	Hard Rock Hotel in New York City
Registration	https://www.maximgrp.com/2025-growth-summit#day1

During the conference, Dr. Whitfill will also conduct one-on-one meetings with registered investors and potential partners, showcasing Azitra's business and clinical development strategy, recent corporate achievements, and anticipated milestones.

About Azitra

Azitra, Inc. is a clinical stage biopharmaceutical company focused on developing innovative therapies for precision dermatology. The Company's lead program, ATR-12, uses an engineered strain of *S. epidermidis* designed to treat Netherton syndrome, a rare, chronic skin disease with no approved treatment options. Netherton syndrome may be fatal in infancy with those living beyond a year having profound lifelong challenges. The ATR-12 program includes a Phase 1b clinical trial in adult Netherton syndrome patients. ATR-04, Azitra's additional advanced program, utilizes another engineered strain of *S. epidermidis* for the treatment of EGFR inhibitor ("EGFRi") associated rash. Azitra has received Fast Track designation from the FDA for EGFRi associated rash, which impacts approximately 150,000 people in the U.S. Azitra has an open IND for its ATR-04 program in patients with EGFRi associated rash. The ATR-12 and ATR-04 programs were developed from Azitra's proprietary platform of engineered proteins and topical live biotherapeutic products that includes a microbial library comprised of approximately 1,500 bacterial strains. The platform is augmented by artificial intelligence and machine learning technology that analyzes, predicts, and helps screen the library of strains for drug like molecules. For more information, please visit **https://azitrainc.com**.

Forward-looking Statements

This press release contains forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995, as amended. These statements may be identified by words such as "aims," "anticipates," "believes," "could," "estimates," "expects," "forecasts," "goal," "intends," "may," "plans," "possible," "potential," "seeks," "will," and variations of these words or similar expressions that are intended to identify forward-looking statements. Any such statements in this press release that are not statements of historical fact may be deemed to be forward-looking statements. These forward-looking statements include, without limitation, statements regarding Azitra's expectations regarding a period to comply with the Plan and applicable Exchange requirements, and actions of Azitra and/or the Exchange to be taken with respect to matters discussed in the notice.

Any forward-looking statements in this press release are based on current expectations, estimates and projections only as of the date of this release and are subject to a number of risks and uncertainties that could cause actual results to differ materially and adversely from those set forth in or implied by such forward-looking statements. These risks and uncertainties include, but are not limited to that we may experience delays in the dosing the first patient in this Phase 1/2 trial; our product candidates may not be effective; there may be delays in regulatory approval or changes in regulatory framework that are out of our control; our estimation of addressable markets of our product candidates may be inaccurate; we may fail to timely raise additional required funding; our actions and/or the Exchange's actions to be taken with respect to matters discussed in the notice from the Exchange; more efficient competitors or more effective competing treatment may emerge; we may be involved in disputes surrounding the use of our intellectual property crucial to our success; we may not be able to attract and retain key employees and qualified personnel; earlier study results may not be predictive of later stage study outcomes; and we are dependent on third-parties for some or all aspects of our product manufacturing, research and preclinical and clinical testing. Additional risks concerning Azitra's programs and operations are described or incorporated by reference in our annual report on Form 10-K filed with the SEC on February 24, 2025. Azitra explicitly disclaims any obligation to update any forward-looking statements except to the extent required by law.

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