



IMUNON to Present at the Life Sciences Investor Forum on June 25th

June 22, 2026

LAWRENCEVILLE, N.J., June 22, 2026 (GLOBE NEWSWIRE) -- [IMUNON, Inc. \(NASDAQ: IMNN\)](#), a clinical-stage company in Phase 3 development with its DNA-mediated immunotherapy, today announced that Stacy Lindborg, Ph.D., President and Chief Executive Officer of IMUNON, will present at the Life Sciences Investor Forum hosted by [VirtualInvestorConferences.com](#) on Thursday, June 25, 2026, at 12:00 p.m. ET.

DATE: Thursday, June 25th, 2026

TIME: 12:00 p.m. ET

[Webcast Registration Link Here](#)

Available for 1x1 meetings: Schedule 1x1 Meetings [HERE](#)

This will be a live, interactive online event where investors are invited to ask the Company questions in real-time. If attendees are not able to join the event live on the day of the conference, an archived webcast will also be made available after the event under the [News & Investors](#) tab of the IMUNON website at [www.imunon.com](#).

It is recommended that online investors pre-register and run the online system check to expedite participation and receive event updates. Learn more about the event at [www.virtualinvestorconferences.com](#).

About IMUNON

IMUNON is a clinical-stage biotechnology company focused on advancing a portfolio of innovative treatments that harness the body's natural mechanisms to generate safe, effective and durable responses across a broad array of human diseases, constituting a differentiating approach from conventional therapies. IMUNON is developing its non-viral DNA technology across its modalities. The first modality, TheraPlas[®], is developed for the gene-based delivery of cytokines and other therapeutic proteins in the treatment of solid tumors where an immunological approach is deemed promising. The second modality, PlaCCine[®], is developed for the gene delivery of viral antigens that can elicit a strong immunological response.

The Company's lead clinical program, IMNN-001, is a DNA-based immunotherapy for the localized treatment of advanced ovarian cancer that has completed multiple clinical trials including one Phase 2 clinical trial (OVATION 2) and is currently conducting a Phase 3 clinical trial (OVATION 3). IMNN-001 works by instructing the body to produce safe and durable levels of powerful cancer-fighting molecules, such as interleukin-12 and interferon gamma, at the tumor site. Additionally, the Company has completed dosing in a first-in-human study of its COVID-19 booster vaccine (IMNN-101). The Company will continue to leverage these modalities and to advance, either directly or through partnership, the technological frontier of plasmid DNA to better serve patients with difficult-to-treat conditions. For more information, please visit [www.imunon.com](#).

Forward-Looking Statements

IMUNON wishes to inform readers that forward-looking statements in this release are made pursuant to the "safe harbor" provisions of the Private Securities Litigation Reform Act of 1995. All statements, other than statements of historical fact, including, but not limited to, statements regarding the timing and enrollment of the Company's clinical trials, the potential of any therapies developed by the Company to fulfill unmet medical needs, the market potential for the Company's products, if approved, the potential efficacy and safety profile of our product candidates, and the Company's plans and expectations with respect to its development programs more generally, are forward-looking statements. We generally identify forward-looking statements by using words such as "may," "will," "expect," "plan," "anticipate," "estimate," "intend" and similar expressions (as well as other words or expressions referencing future events, conditions or circumstances). Readers are cautioned that such forward-looking statements involve risks and uncertainties including, without limitation, uncertainties relating to unforeseen changes in the course of research and development activities and in clinical trials, including the fact that interim results are not necessarily indicative of final results; the uncertainties of and difficulties in analyzing interim clinical data; the significant expense, time and risk of failure in conducting clinical trials; the need for IMUNON to evaluate its future development plans; possible actions by customers, suppliers, competitors or regulatory authorities; and other risks detailed from time to time in IMUNON's filings with the Securities and Exchange Commission. IMUNON assumes no obligation, except to the extent required by law, to update or supplement forward-looking statements that become untrue because of subsequent events, new information or otherwise.

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