Medicenna Therapeutics to Host a Live Webinar with Q&A to Discuss Updated MDNA11 Clinical Data

December 9, 2025

TORONTO and HOUSTON, Dec. 09, 2025 (GLOBE NEWSWIRE) -- Medicenna Therapeutics Corp. (“Medicenna” or the “Company”) (TSX: MDNA, OTCQX: MDNAF), a clinical-stage immunotherapy company focused on the development of Superkines for the treatment of cancer as well as autoimmune and inflammatory diseases is pleased to announce that it will host a live webinar on December 10, 2025 at 08:30 AM Eastern Time.

As previously announced, Medicenna will present updated clinical data from the ABILITY-1 Phase 1/2 Study, evaluating MDNA11 as a monotherapy and in combination with pembrolizumab, on December 10th at the ESMO Immuno-Oncology Congress 2025. The webinar will have Medicenna’s management team and the presenting Principal Investigator along with commentary from key opinion leaders to discuss the updated data.

Medicenna Therapeutics – KOL Webinar | Live Event Details:

- **Date:** December 10, 2025
- **Time:** 08:30 – 09:30 AM ET
- **Format:** Live Webinar with Q&A
- **Registration:** Participants can register for the webinar through the link: [Registration]. A replay of the webinar will be available on Medicenna’s website following the event.

The webinar will feature presentations from Medicenna's executive and scientific advisory team, including Dr. Fahar Merchant, President and CEO, Dr. Arash Yavari, Director of Clinical Strategy, and the presenting Principal Investigator Dr. André Mansinho, Assistant Professor, Faculty of Medicine, University of Lisbon. Additional commentary from key opinion leaders and live Q&A will follow the presentation.

About Medicenna Therapeutics

Medicenna is a clinical-stage immunotherapy company focused on developing novel, highly selective versions of IL-2, IL-4 and IL-13 Superkines and first-in-class Empowered Superkines. Medicenna’s long-acting IL-2 Superkine, MDNA11, is a next-generation IL-2 with superior affinity toward CD122 (IL-2 receptor beta) and no CD25 (IL-2 receptor alpha) binding, thereby preferentially stimulating cancer-killing effector T cells and NK cells. Medicenna’s first-in-class targeted PD-1 x IL-2 bispecific, MDNA113, is in development for solid tumors and was designed using the Company’s proprietary BiSKITs ™ (Bifunctional SuperKine ImmunoTherapies) and T-MASK™ (Targeted Metalloprotease Activated SuperKine) platforms Medicenna’s IL-4 Empowered Superkine, bizaxofusp (formerly MDNA55), has been studied in 5 clinical trials enrolling over 130 patients, including a Phase 2b trial for recurrent GBM, the most common and uniformly fatal form of brain cancer. Bizaxofusp has obtained FastTrack and Orphan Drug status from the FDA and FDA/EMA, respectively.

For more information, please visit www.medicenna.com, and follow us on X and LinkedIn.

Forward-Looking Statements

This news release contains forward-looking statements within the meaning of applicable securities laws. Forward-looking statements include, but are not limited to, express or implied statements regarding the future operations of the Company, estimates, plans, strategic ambitions, partnership activities and opportunities, objectives, expectations, opinions, forecasts, projections, guidance, outlook or other statements that are not historical facts. Forward-looking statements are often identified by terms such as “will”, “may”, “should”, “anticipate”, “expect”, “believe”, “seek”, “potentially” and similar expressions. and are subject to risks and uncertainties. There can be no assurance that such statements will prove to be accurate and actual results and future events could differ materially from those anticipated in such statements. Important factors that could cause actual results to differ materially from the Company’s expectations include the risks detailed in the latest annual information form of the Company and in other filings made by the Company with the applicable securities regulators from time to time in Canada.

The reader is cautioned that assumptions used in the preparation of any forward-looking information may prove to be incorrect. Events or circumstances may cause actual results to differ materially from those predicted, as a result of numerous known and unknown risks, uncertainties, and other factors, many of which are beyond the control of the Company. The reader is cautioned not to place undue reliance on any forward-looking information. Such information, although considered reasonable by management, may prove to be incorrect and actual results may differ materially from those anticipated. Forward-looking statements contained in this news release are expressly qualified by this cautionary statement. The forward-looking statements contained in this news release are made as of the date hereof and except as required by law, we do not intend and do not assume any obligation to update or revise publicly any of the included forward-looking statements.

This news release contains hyperlinks to information that is not deemed to be incorporated by reference in this new release.

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