

PRESS RELEASE - May 13, 2024 - 6:00pm CEST - Montpellier, France - Euronext: MEDCL

# Medincell to Host Videoconference on May 15, 2024, at 10:30 AM CEST to Discuss Company's Latest News, Outlook, and R&D Activities

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Medincell is a clinical- and commercial-stage biopharmaceutical licensing company developing long-acting injectable drugs in many therapeutic areas. The first treatment based on BEPO<sup>®</sup> technology, aimed at treating schizophrenia, was approved by the FDA in April 2023, and has since been distributed in the United States by Teva under the name UZEDY<sup>®</sup> (BEPO<sup>®</sup> technology is licensed to Teva under the name SteadyTeq<sup>™</sup>).

Two other treatments are currently in Phase 3 clinical trials, including mdc-TJK, also developed with Teva, whose positive efficacy results were announced on May 8, 2024 (press release).

Medincell also announced on April 16, 2024, a co-development and licensing agreement with the American pharmaceutical group AbbVie, covering the development of 6 products (press release).

### Product portfolio and R&D pipeline as of May 13, 2024



For strategic reasons, the Company has decided to terminate the mdc-GRT program (tacrolimus, transplantation) which was in the preclinical stage.

# **About Medincell**

Medincell is a clinical- and commercial-stage biopharmaceutical licensing company developing long-acting injectable drugs in many therapeutic areas. Our innovative treatments aim to guarantee compliance with medical prescriptions, to improve the effectiveness and accessibility of medicines, and to reduce their environmental footprint. They combine active pharmaceutical ingredients with our proprietary BEPO® technology which controls

the delivery of a drug at a therapeutic level for several days, weeks or months from the subcutaneous or local injection of a simple deposit of a few millimeters, entirely bioresorbable. The first treatment based on BEPO<sup>®</sup> technology, intended for the treatment of schizophrenia, was approved by the FDA in April 2023, and is now distributed in the United States by Teva under the name UZEDY<sup>®</sup> (BEPO<sup>®</sup> technology is licensed to Teva under the name SteadyTeq<sup>™</sup>). We collaborate with leading pharmaceutical companies and foundations to improve global health through new treatment options. Based in Montpellier, Medincell currently employs more than 140 people representing more than 25 different nationalities.

UZEDY<sup>®</sup> and SteadyTeq™ are trademarks of Teva Pharmaceuticals

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This press release contains forward-looking statements, including statements regarding Company's expectations for (i) the timing, progress and outcome of its clinical trials; (ii) the clinical benefits and competitive positioning of its product candidates; (iii) its ability to obtain regulatory approvals, commence commercial production and achieve market penetration and sales; (iv) its future product portfolio; (v) its future partnering arrangements; (vi) its future capital news, capital expenditure plans and ability to obtain funding; and (viii) prospective financial matter regarding our business. Although the Company believes that its expectations are based on reasonable assumptions, any statements other than statements of historical facts that may be contained in this press release relating to future events are forward-looking statements and subject to change without notice, factors beyond the Company's control and the Company's financial canabilities.

These statements may include, but are not limited to, any statement beginning with, followed by or including words or phrases such as "objective", "believe", "anticipate", "expect", "foresee", "aim", "intend", "may", "anticipate", "estimate", "plan", "project", "will", "may", "probably", "potential", "should", "could" and other words and phrases of the same meaning or used in negative form. Forward-looking statements are subject to inherent risks and uncertainties beyond the Company's control that may, if any, cause actual results, performance, or achievements to differ materially from those anticipated or expressed explicitly or implicitly by such forward-looking statements. A list and description of these risks, contingencies and uncertainties can be found in the documents filed by the Company with the Autorité des Marchés Financiers (the "AMF") pursuant to its regulatory obligations, including the Company's registration document, registered with the AMF on September 4, 2018, under number I. 18-062 (the "Registration Document"), as well as in the documents and reports to be published subsequently by the Company. In particular, readers' attention is drawn to the section entitled "Facteurs de Risques" on page 26 of the Registration Document.

Any forward-looking statements made by or on behalf of the Company speak only as of the date they are made. Except as required by law, the Company does not undertake any obligation to publicly update these forward-looking statements or to update the reasons why actual results could differ materially from those anticipated by the forward-looking statements, including in the event that new information becomes available. The Company's update of one or more forward-looking statements does not imply that the Company imply that the Company imply that the Company imply that the company statements. Readers are cautioned not to place undue reliance on these forward-looking statements.

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