



MedinCell at the 2023 Jefferies Global Healthcare Conference - London, 14-16 November

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- **Christophe Douat, CEO, and Richard Malamut, CMO, will present MedinCell's commercial and advanced-development stage portfolio at the Jefferies London Healthcare Conference on November 16 at 1:30pm GMT**
 - **Investors can access the presentation remotely, either live or on replay, via the following link:**
<https://wsw.com/webcast/jeff287/medcl/1983116>
 - **MedinCell management team will be available for individual interviews with investors throughout the conference**
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MedinCell has recently reached the commercial stage with the market launch by Teva of UZEDY™ for the treatment of schizophrenia. UZEDY is the first innovative product based on MedinCell's long-acting injectable (LAI) BEPO® technology approved by US FDA.

In parallel, MedinCell is developing a portfolio of innovative treatments also using the BEPO technology. This notably includes the following candidate treatments:

2 candidates in clinical Phase 3:

- **mdc-TJK:** Potentially, the first long-acting injectable olanzapine with a favorable safety profile that could enable a large adoption for treatment of patients with more severe forms of schizophrenia
- **mdc-CWM:** An innovative sustained-release formulation of a non-steroidal anti-inflammatory drug celecoxib aiming at facilitating patient recovery after total knee replacement and decreasing the need for potentially addictive opioids

3 candidates ready for Phase 1 studies in 2024:

- **mdc-WWM:** 6-month active subcutaneous contraceptive
- **mdc-GRT:** Monthly subcutaneous tacrolimus LAI to prevent solid organ graft rejection
- **mdc-STM:** Global Health program to fight malaria

Other programs are currently in the evaluation and formulation stage, some as part of the internal program development strategy, others within the framework of partnerships.

About MedinCell

MedinCell is a clinical- and commercial-stage biopharmaceutical 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 already known and used active 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 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™ (BEPO technology is licensed to Teva under the name SteadyTeq™).

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™ 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 needs, capital expenditure plans and ability to obtain funding; and (vii) prospective financial matters 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 capabilities.

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 will make any further updates to such forward-looking statements or other forward-looking statements. Readers are cautioned not to place undue reliance on these forward-looking statements.

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