medincell.

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Medincell announces the availability of its 2023-24 Universal Registration Document including the Annual Financial and CSR Reports

The 2023-24 Universal Registration Document (URD) filed with the French market authority (Autorités des Marchés Financiers, or AMF) under the reference D.24-0649 includes:

- the Annual Financial Report for the year ending on March 31, 2024
- the management report
- the CSR report
- the report on corporate governance
- the proposed text of the resolutions to be submitted to the Shareholders' Meeting of September 12, 2024

The URD can be consulted on the Company's website (Investors section) and on the AMF's website (www.amf-france.org).

The CSR report alone can also be downloaded from the Company's website (Investors section).

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[®] 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 UZEDY[®] (BEPO[®] technology is licensed to Teva under the name uter options. Based in Montpellier, Medincell currently employs more than 140 people representing more than 25 different nationalities.

UZEDY[®] and SteadyTeq[™] are registered trademarks of Teva Pharmaceuticals.

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This press release may contain forward-looking statements, particularly concerning the progress of the Company's clinical trials. Although the Company considers that its forecasts are based on reasonable assumptions, any statements other than statements of historical fact that may be contained in this press release relating to future events are subject to change without notice, to factors beyond the Company's control and to the Company's financial capabilities.

These statements may include, but are not limited to, any statements beginning with, followed by or including words or expressions such as "objective", "believe", "expect", "aim", "intend", "may", "anticipate", "estimate", "plan", "project", "will", "may", "probably", "should", "could" and other words or expressions of similar meaning or used in the negative. Forward-looking statements are subject to inherent risks and uncertainties beyond the Company's control which may cause actual results, performance or achievements of the Company to differ materially from those anticipated or implied by such statements.

A list and description of such risks, hazards and uncertainties can be found in the documents filed by the Company with the Autorité des Marchés Financiers (AMF) pursuant to its regulatory obligations, including in the Company's document de base, registered with the AMF on September 4, 2018 under number I. 18-062, as well as in documents and reports to be published

subsequently by the Company. Furthermore, these forward-looking statements only apply as of the date of this press release. Readers are cautioned not to place undue reliance on these forward-looking statements. Except as required by law, the Company undertakes no obligation to publicly update these forward-looking statements, nor to update the reasons why actual results may differ materially from those anticipated in the forward-looking statements, even if new information becomes available. The Company's updating of one or more forward-looking statements does not imply that it will or will not update these or any other forward-looking statements.

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