RNS Number : 3679U
Futura Medical PLC
27 March 2023

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Futura Medical plc

("Futura" or the "Company")

Notice of Preliminary Results 2022

Futura Medical plc (AIM: FUM), a pharmaceutical company developing a portfolio of innovative products based on its proprietary, transdermal DermaSys® drug delivery technology and currently focused on sexual health and pain, will announce its preliminary results for the year ended 31 December 2022 on Wednesday, 5 April 2023.

James Barder, Chief Executive Officer, Angela Hildreth, Finance Director and Chief Operating Officer, and Ken James, Executive Director and Head of R&D, will host a webcast for analysts on the day of the results at 10:00am BST, which will be made available within the investor centre section of the Company's website.

Please submit questions ahead of the webcast to investor.relations@futuramedical.com by 10:00am BST on Thursday 30 March 2023.

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About Futura Medical plc

Futura Medical plc (AIM: FUM), is a pharmaceutical company developing a portfolio of innovative products based on its proprietary, transdermal DermaSys® technology. Each DermaSys® formulation is separately patented and specifically tailored for the selected indication and application, as well as being optimised for clinical efficacy, safety, administration and patient convenience. The products are developed for the prescription and consumer healthcare markets as appropriate. Current therapeutic areas are sexual health, including erectile dysfunction, and pain relief. Development and commercialisation strategies are designed to maximise product differentiation and value creation whilst minimising risk.

MED3000 is Futura's topical gel formulation that is a breakthrough treatment for erectile dysfunction ("ED") through a unique evaporative mode of action. Futura has previously conducted a Phase 3 study using MED3000 in ED, referred to as "FM57". This was a 1,000 patient, dose-ranging, multi-centre, randomised, double blind, placebo-controlled, home use, parallel group study, delivering highly statistically significant results compared to pre-treatment baseline, using measures IIEF-EF, SEP2 and SEP3 (internationally accepted clinical trial endpoints in ED) with over 60% of patients experiencing a clinically meaningful improvement in their ED. A second confirmatory Phase 3 clinical study, "FM71" was also conducted to support Futura's regulatory submission to the FDA with 96 ED patients and endpoints at 24 weeks.

also conducted to support Futura's regulatory submission to the FDA with 90 ED patients and endpoints at 24 weeks, demonstrating that MED3000 presents an effective clinically proven treatment for ED with a rapid speed of onset and a favourable benefit versus risk profile, ideally suited for OTC classification.

MED3000 is CE marked inEuropeand CA marked in theUKas a clinically proven topical treatment for adult men with erectile dysfunction under the brand Eroxon® with a key claim of "Helps you get an erection within 10 minutes".

Futura is based in Guildford, Surrey, and its shares trade on the AIM market of the London Stock Exchange. www.futuramedical.com/

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