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DexTech Medical's Myeloma Study, Dose Group 2, DMC Gives OK for Dose Group 3

The study is being conducted at Karolinska University Hospital Huddinge and at Uddevalla Hospital. The treatment lasts for a total of 14 weeks with 2 doses per month. Three dose levels of ODX are studied, 3mg/kg body weight, 6mg/kg, and 9mg/kg. The Principal Investigator (PI) is Dr Katarina Uttervall, MD, PhD, Department of Hematology/HERM, Karolinska University Hospital Huddinge. Dr Dorota Knut is the principal investigator at the Department of Hematology at Uddevalla Hospital. Analysis of biomarkers takes place at the Central Laboratory, Karolinska University Hospital Solna, NKS. Adult myeloma patients with progressive *treatment-resistant disease*, who have previously received 1–5 prior lines of therapy, are included in the study. The primary objective is to confirm ODX safety and tolerability and with a secondary objective to demonstrate indications of treatment response.

Dose group 2 (6mg/kg) has been fully recruited. Two patients in dose group 2 have progressive disease after completion of treatment. The other 2 patients will finish the treatment in early December. The DMC (Independent Data Monitoring Committee) approves the continuation to dose group 3. No ODX related serious side effects have been noted. A new patient for dose group 3 has been screened.
Follow-up of all patients who have achieved stable disease is done to determine how long the disease-slowing effect persists after the ODX treatment has been discontinued, i.e., until new progress.

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This information is information that DexTech Medical AB is obliged to make public pursuant to the EU Market Abuse Regulation. The information was submitted for publication, through the agency of the contact person set out above, on October 16, 2025.

***DexTech Medical AB** is a Swedish research company that, based on its technology platform, has developed four drug candidates that are protected by patents. The lead candidate is OsteoDex for the treatment of castration-resistant prostate cancer (CRPC) with bone metastases. A successful clinical phase II study has been conducted with OsteoDex where the results show high tolerability with mild side effects and treatment effect on patients who fail existing drugs. DexTech's goal is to out-license each drug candidate no later than after completion of the phase II study. DexTech Medical AB is listed on the Spotlight Stock Market.*