

Genmab Announces Sale of Manufacturing Facility to Baxter

Company Announcement

- Genmab sells manufacturing facility to Baxter Healthcare Corporation
- Genmab receives USD 10 million

Copenhagen, Denmark; February 28, 2013 – Genmab A/S (OMX: GEN) announced today the signing and closing of an agreement with Baxter Healthcare Corporation for the sale of Genmab’s non-plasma-derived antibody manufacturing facility, located in Brooklyn Park, Minnesota, USA, for USD 10 million. Under the terms of the agreement Genmab receives the USD 10 million (approximately DKK 57 million) in cash and Baxter acquires the facility, land, and equipment at the Brooklyn Park site. Baxter will offer employment to the 23 employees currently supporting the facility.

“We have delivered on our commitment to execute the sale of the antibody manufacturing facility in the first quarter of this year and are very pleased that Baxter acquired the facility, including the 23 employees who have maintained the facility to such a high standard,” said Jan van de Winkel, Ph.D., Chief Executive Officer of Genmab.

About Genmab A/S

Genmab is a publicly traded, international biotechnology company specializing in the creation and development of differentiated human antibody therapeutics for the treatment of cancer. Founded in 1999, the company’s first marketed antibody, ofatumumab (Arzerra®), was approved to treat chronic lymphocytic leukemia in patients who are refractory to fludarabine and alemtuzumab after less than eight years in development. Genmab’s validated and next generation antibody technologies are expected to provide a steady stream of future product candidates. Partnering of innovative product candidates and technologies is a key focus of Genmab’s strategy and the company has alliances with top tier pharmaceutical and biotechnology companies. For more information visit www.genmab.com.

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This Company Announcement contains forward looking statements. The words “believe”, “expect”, “anticipate”, “intend” and “plan” and similar expressions identify forward looking statements. Actual results or performance may differ materially from any future results or performance expressed or implied by such statements. The important factors that could cause our actual results or performance to differ materially include, among others, risks associated with pre-clinical and clinical development of products, uncertainties related to the outcome and conduct of clinical trials including unforeseen safety issues, uncertainties related to product manufacturing, the lack of market acceptance of our products, our inability to manage growth, the competitive environment in relation to our business area and markets, our inability to attract and retain suitably qualified personnel, the unenforceability or lack of protection of our patents and proprietary rights, our relationships with affiliated entities, changes and developments in technology which may render our products obsolete, and other factors. For a further discussion of these risks, please refer to the risk management sections in Genmab’s most recent financial reports, which are available on www.genmab.com. Genmab does not undertake any obligation to update or revise forward looking statements in this Company Announcement nor to confirm such statements in relation to actual results, unless required by law.

Genmab A/S and its subsidiaries own the following trademarks: Genmab®; the Y-shaped Genmab logo®; HuMax®, HuMax-CD20®; DuoBody®, HexaBody™ and UniBody®. Arzerra® is a trademark of GlaxoSmithKline.