



GLAXOSMITHKLINE RECEIVES CONDITIONAL MARKETING AUTHORIZATION IN THE EU FOR ARZERRA™ (OFATUMUMAB)

Summary: GSK has received conditional marketing authorization in the EU for Arzerra™ (ofatumumab) for the treatment of refractory CLL.

Copenhagen, Denmark; April 19, 2010 –GlaxoSmithKline (GSK) and Genmab A/S (OMX: GEN) confirmed today that the European Commission (EC) has granted a conditional marketing authorization for Arzerra™ (ofatumumab) for the treatment of refractory chronic lymphocytic leukemia (CLL). Ofatumumab is indicated for the treatment of CLL in patients who are refractory (have not responded) to fludarabine and alemtuzumab. Fludarabine and alemtuzumab are standard therapies currently used to treat CLL.

About Genmab

Genmab A/S is a leading international biotechnology company focused on developing fully human antibody therapeutics for the potential treatment of cancer. Genmab's world class discovery and development teams are using cutting-edge technology to create and develop products to address unmet medical needs. Our primary goal is to improve the lives of patients who are in urgent need of new treatment options. For more information on Genmab's products and technology, visit www.genmab.com.

About GlaxoSmithKline

GlaxoSmithKline – one of the world's leading research-based pharmaceutical and healthcare companies – is committed to improving the quality of human life by enabling people to do more, feel better, and live longer. For company information, visit GlaxoSmithKline at <http://www.gsk.com>

This Stock Exchange Release 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 product discovery and development, 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 section “Risk Management” in Genmab's Annual Report, which is available on www.genmab.com. Genmab does not undertake any obligation to update or revise forward looking statements in this Stock Exchange Release nor to confirm such statements in relation to actual results, unless required by law.

Genmab®; the Y-shaped Genmab logo®; HuMax®; HuMax-CD20®; HuMax-EGFr™; HuMax-IL8™; HuMax-TAC™; HuMax-HepC™; HuMax-CD38™; HuMax-CD32b™; HuMax-TF™; HuMax-Her2™; HuMax-VEGF™, HuMax-Wnt and UniBody® are all trademarks of Genmab A/S. Arzerra™ is a trademark of GlaxoSmithKline.

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