Genmab to Host Capital Markets Day

October 15, 2020

Media Release

Copenhagen, Denmark, October 15, 2020

- **Genmab will host Capital Markets Day November 13, 2020**
- **Highlights to be covered will include an overview of our pipeline with focus on epcoritamab, tisotumab vedotin and DuoBody®-PD-L1x4-1BB, our DuoBody® technology platform, and insights into our growing capabilities across Genmab**

Genmab A/S (Nasdaq: GMAB) announced today that it will hold a Capital Markets Day for analysts and investors on November 13, 2020 from 8:30 AM EST / 2:30 PM CET to 11:00 AM EST / 5:00 PM CET. The event will be webcast live from Genmab's state-of-the-art R&D Center in Utrecht, the Netherlands, and from its new offices and laboratory facilities in Princeton, NJ, US. The Capital Markets Day will feature Genmab's Executive Management Team and a variety of high-level Genmab speakers who will provide updates on Genmab's business and the ways in which we are delivering on our commitments and evolving into a fully integrated biotech innovation powerhouse.

Topics to be discussed include an overview of our robust product pipeline with a focus on our exciting epcoritamab, tisotumab vedotin and DuoBody-PD-L1x4-1BB programs, a deep dive into our proprietary DuoBody technology and a look at how we are growing our capabilities across the company.

To register for Genmab's 2020 Capital Markets Day, please visit the following website:
https://events.bizzabo.com/GenmabCMD

About Genmab
Genmab is a publicly traded, international biotechnology company specializing in the creation and development of differentiated antibody therapeutics for the treatment of cancer. Founded in 1999, the company is the creator of the following approved antibodies: DARZALEX® (daratumumab, under agreement with Janssen Biotech, Inc.) for the treatment of certain multiple myeloma indications in territories including the U.S., Europe and Japan, Kesimpta® (subcutaneous ofatumumab, under agreement with Novartis AG), for the treatment of adults with relapsing forms of multiple sclerosis in the U.S. and TEPEZZA® (teprotumumab, under agreement with Roche granting sublicense to Horizon Therapeutics plc) for the treatment of thyroid eye disease in the U.S. A subcutaneous formulation of daratumumab, known as DARZALEX FASPRO™ (daratumumab and hyaluronidase-fihj) in the U.S., has been approved in the U.S. and Europe for the treatment of adult patients with certain multiple myeloma indications. The first approved Genmab created therapy, Arzerra® (ofatumumab, under agreement with Novartis AG), approved for the treatment of certain chronic lymphocytic leukemia indications, is available in Japan and is also available in other territories via compassionate use or oncology access programs. Daratumumab is in clinical development by Janssen for the treatment of additional multiple myeloma indications, other blood cancers and amyloidosis. Genmab also has a broad clinical and pre-clinical product pipeline. Genmab's technology base consists of validated and proprietary next generation antibody technologies - the DuoBody® platform for generation of bispecific antibodies, the HexaBody® platform, which creates effector function enhanced antibodies, the HexElect® platform, which combines two co-dependently acting HexaBody molecules to introduce selectivity while maximizing therapeutic potency and the DuoHexaBody® platform, which enhances the potential potency of bispecific antibodies through hexamerization. The company intends to leverage these technologies to create opportunities for full or co-ownership of future products. Genmab has alliances with top tier pharmaceutical and biotechnology companies. Genmab is headquartered in Copenhagen, Denmark with sites in Utrecht, the Netherlands, Princeton, New Jersey, U.S. and Tokyo, Japan.

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This Media 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 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 or technologies 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 and the risk factors included in Genmab's most recent Annual Report on Form 20-F and other filings with the U.S. Securities and Exchange Commission (SEC), which are available at www.sec.gov. Genmab does

not undertake any obligation to update or revise forward looking statements in this Media Release nor to confirm such statements to reflect subsequent events or circumstances after the date made or in relation to actual results, unless required by law.

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