Genmab Announces Data to be Presented at 24th EHA Annual Congress

May 16, 2019

Media Release

Copenhagen, Denmark, May 16, 2019

15 industry-sponsored abstracts featuring Genmab programs selected for presentation at EHA Annual Congress

Genmab A/S (Nasdaq Copenhagen: GEN) announced today that 15 industry sponsored abstracts regarding Genmab programs were accepted for presentation at the 24th European Hematology Association (EHA) Annual Congress 2019 in Amsterdam, the Netherlands, taking place June 13-16, 2019. A list of accepted Industry-sponsored abstracts featured at the congress includes 14 daratumumab abstracts, four of which were accepted for oral presentations, including a presentation of the Phase III CASSIOPEIA data, which the Scientific Program Committee of the EHA selected for presentation during the Presidential Symposium, which showcases abstracts that represent innovative research in hematology. In addition, one abstract features Genmab’s proprietary DuoBody®-CD3xCD20 product. The abstracts have been published on the EHA website and may be accessed via www.ehaweb.org.

“The presentation of impressive pre-clinical data on our DuoBody-CD3xCD20 program exemplifies how Genmab is advancing its proprietary product pipeline using our strong expertise in antibody drug development to create truly differentiated products to help patients with hematologic malignancies. We are also very pleased that the EHA has selected the CASSIOPEIA data for presentation during the prestigious Presidential Symposium as it reinforces Genmab's impactful contribution to multiple myeloma treatment,” said Jan van de Winkel, Ph.D., Chief Executive Officer of Genmab.

Industry-Sponsored Abstracts are as follows:

DuoBody-CD3xCD20:
Potent Anti-tumor Activity of DuoBody-CD3xCD20 in Pre-clinical Models In Vitro and In Vivo – Poster presentation, Saturday, June 15, 5:30 PM – 7:00 PM CEST

Daratumumab (Submitted by Janssen Biotech, Inc.):
Phase 3 Randomized Study of Daratumumab, Bortezomib, Thalidomide, and Dexamethasone (VTd) Versus VTd in Transplant-eligible Newly Diagnosed Multiple Myeloma: Part 1 CASSIOPEIA Results – Oral presentation, Friday, June 14, 3:45 PM – 4:00 PM CEST
Efficacy of Daratumumab, Bortezomib, Thalidomide, and Dexamethasone in Transplant-eligible Newly Diagnosed Multiple Myeloma Based Minimal Residual Disease Status: Analysis of CASSIOPEIA – Oral presentation, Saturday, June 15, 4:45 PM – 5:00 PM CEST
Randomized, Open-label, Non-inferiority, Phase 3 Study of Subcutaneous Versus Intravenous Daratumumab Administration in Patients with Relapsed or Refractory Multiple Myeloma: COLUMBA – Oral presentation, Saturday, June 15, 11:30 AM – 11:45 AM CEST
Subcutaneous Daratumumab, Cyclophosphamide, Bortezomib, and Dexamethasone in Patients with Newly Diagnosed Amyloid Light Chain Amyloidosis: Updated Safety Run-in Results of ANDROMEDA – Oral presentation, Saturday, June 15, 5:00 PM – 5:15 PM CEST
Stem Cell Yield and Transplantation in Transplant-eligible Newly Diagnosed Multiple Myeloma Patients Receiving Daratumumab, Bortezomib, Thalidomide, and Dexamethasone: Phase 3 CASSIOPEIA Study – Poster presentation, Friday, June 14, 5:30 PM – 7:00 PM CEST
Impact of Age on Efficacy and Safety of Daratumumab in Combination with Lenalidomide and Dexamethasone in Patients with Transplant-ineligible Newly Diagnosed Multiple Myeloma: MAIA – Poster presentation, Friday, June 14, 5:30 PM – 7:00 PM CEST
Faster and Sustained Improvement in Health-related Quality of Life in Transplant-eligible Newly Diagnosed Multiple Myeloma Patients Treated with Daratumumab, Lenalidomide, and Dexamethasone (D-Rd) Versus Rd: MAIA – Poster presentation, Friday, June 14, 5:30 PM – 7:00 PM CEST
Efficacy and Safety of Daratumumab, Lenalidomide, and Dexamethasone in Relapsed or Refractory Multiple Myeloma: Updated Subgroup Analysis of POLLUX Based on Cytogenetic Risk – Poster presentation, Friday, June 14, 5:30 PM – 7:00 PM CEST
Efficacy and Safety of Daratumumab, Bortezomib, and Dexamethasone in Relapsed or Refractory Multiple Myeloma: Updated Subgroup Analysis of CASTOR Based on Cytogenetic Risk – Poster presentation, Friday, June 14, 5:30 PM – 7:00 PM CEST
Characterization of Treatments and Real-life Outcomes in Patients with Newly Diagnosed Multiple Myeloma Who Received Frontline Autologous Stem Cell Transplantation in Sweden – Poster presentation, Friday, June 14, 5:30 PM – 7:00 PM CEST
Characterization of Frontline Treatment Patterns and the Proportion of Patients Reaching Subsequent Lines of Therapy in Transplant-eligible Patients with Newly Diagnosed Multiple Myeloma – Poster presentation, Friday, June 14, 5:30 PM – 7:00 PM CEST
Improvement in Health-related Quality of Life for Newly Diagnosed Multiple Myeloma Transplant-eligible Patients Treated with Daratumumab, Bortezomib, Thalidomide, and Dexamethasone: CASSIOPEIA – Poster presentation, Saturday, June 15, 5:30 PM – 7:00 PM CEST
Results of the Daratumumab Monotherapy Early Access Treatment Protocol in Patients from Europe and Russia with Relapsed or Refractory Multiple...
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 has two approved antibodies, DARZALEX® (daratumumab) for the treatment of certain multiple myeloma indications, and Arzerra® (ofatumumab) for the treatment of certain chronic lymphocytic leukemia indications. Daratumumab is in clinical development for additional multiple myeloma indications and other blood cancers. A subcutaneous formulation of ofatumumab is in development for relapsing multiple sclerosis. 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 and the HexElect® platform, which combines two co-dependently acting HexaBody molecules to introduce selectivity while maximizing therapeutic potency. 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. For more information visit www.genmab.com.

Contact:
Marisol Peron, Corporate Vice President, Communications & Investor Relations
T: +1 609 524 0065; E: mmp@genmab.com

For Investor Relations:
Andrew Carlsen, Senior Director, Investor Relations
T: +45 3377 9558; E: acn@genmab.com

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 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 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.

Genmab A/S and/or its subsidiaries own the following trademarks: Genmab®, the Y-shaped Genmab logo®, Genmab in combination with the Y-shaped Genmab logo®, HuMax®, DuoBody®, DuoBody in combination with the DuoBody logo®, HexaBody®, HexaBody in combination with the HexaBody logo®, DuoHexaBody®, HexElect®, and UniBody®. Arzerra® is a trademark of Novartis AG or its affiliates. DARZALEX® is a trademark of Janssen Pharmaceutica NV.

Media Release no. 07
CVR no. 2102 3884
LEI Code 529900MTJPDE4MHJ122

Genmab A/S
Kalvebod Brygge 43
1560 Copenhagen V
Denmark

Attachment
• 190516_i07MR_EHA Curtain Raiser