

Genmab Announces Data to be Presented at 2019 ASCO Annual Meeting

Media Release

Copenhagen, Denmark, April 17, 2019

14 industry sponsored abstracts regarding Genmab programs selected for presentation at ASCO Annual Meeting

Genmab A/S (Nasdaq Copenhagen: GEN) announced today that 14 industry sponsored abstracts regarding Genmab programs were accepted for presentation at the American Society of Clinical Oncology (ASCO) Annual Meeting, taking place in Chicago from May 31 to June 4, 2019. The titles of the abstracts are currently available on the [ASCO iPlanner website](#), with the full abstracts scheduled to be published on May 15, 2019. A list of accepted industry sponsored abstracts is provided below, and includes ten daratumumab abstracts, two of which were accepted for oral presentations, one abstract on enapotamab vedotin, and two abstracts on tisotumab vedotin (Trial in Progress abstracts).

“We are pleased that once again a wide selection of abstracts concerning Genmab proprietary and partnered programs was selected for presentation at the prestigious ASCO Annual Meeting. We are especially looking forward to the first full presentations of data from the COLUMBA and CASSIOPEIA daratumumab Phase III trials,” said Jan van de Winkel, Ph.D., Chief Executive Officer of Genmab. “We are equally excited to begin sharing with the oncology community early data from enapotamab vedotin and ongoing trials in progress for tisotumab vedotin.”

Representatives from Genmab will be in attendance at ASCO, booth #24159.

List of Industry Sponsored Abstracts:

Daratumumab (Submitted by Janssen Biotech, Inc.):

Phase 3 Randomized Study of Daratumumab + Bortezomib/Thalidomide/Dexamethasone (D-VTd) Vs VTd in Transplant-eligible Newly Diagnosed Multiple Myeloma: CASSIOPEIA Part 1 Results – Oral presentation, Sunday, June 2, 10:45 AM – 10:57 PM CDT

Efficacy and Safety of the 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, Sunday, June 2, 11:09 AM – 11:21 AM CDT

Efficacy of Daratumumab + Bortezomib/Thalidomide/Dexamethasone (D-VTd) in Transplant-eligible Newly Diagnosed Multiple Myeloma Based on Minimal Residual Disease Status: Analysis of the CASSIOPEIA Trial –Poster presentation, Monday, June 3, 8:00 AM – 11:00 AM CDT

Stem Cell Yield and Transplantation Results from Transplant-eligible Newly Diagnosed Multiple Myeloma Patients Receiving Daratumumab + Bortezomib/Thalidomide/Dexamethasone (D-VTd) in the Phase 3 CASSIOPEIA Study –Poster presentation, Monday, June 3, 8:00 AM – 11:00 AM CDT

Impact of age on efficacy and safety of daratumumab in combination with lenalidomide and dexamethasone (D-Rd) in patients with transplant-ineligible newly diagnosed multiple myeloma: MAIA – Poster presentation, Monday, June 3, 8:00 AM – 11:00 AM CDT

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Faster and sustained improvement in health-related quality of life for newly diagnosed multiple myeloma patients ineligible for transplant treated with daratumumab, lenalidomide, and dexamethasone (D-Rd) vs Rd alone: MAIA – Poster presentation, Monday, June 3, 8:00 AM – 11:00 AM CDT

Efficacy and safety of daratumumab, bortezomib, and dexamethasone (D-Vd) in relapsed or refractory multiple myeloma based on cytogenetic risk: updated subgroup analysis of CASTOR – Poster presentation, Monday, June 3, 8:00 AM – 11:00 AM CDT

Efficacy and safety of daratumumab, lenalidomide, and dexamethasone (D-Rd) in relapsed or refractory multiple myeloma: updated subgroup analysis of POLLUX based on cytogenetic risk – Poster presentation, Monday, June 3, 8:00 AM – 11:00 AM CDT

Bortezomib, lenalidomide, and dexamethasone (VRd) ± daratumumab in patients with transplant-eligible newly diagnosed multiple myeloma: a multicenter, randomized, phase 3 study (PERSEUS) – Poster, Monday, June 3, 8:00 AM – 11:00 AM CDT

Bortezomib, lenalidomide, and dexamethasone (VRd) ± daratumumab in patients with newly diagnosed multiple myeloma for whom transplant is not planned as initial therapy: a multicenter, randomized, phase 3 study (CEPHEUS) – Poster presentation, Monday, June 3, 8:00 AM – 11:00 AM CDT

Tisotumab vedotin (Submitted by Seattle Genetics):

SGNTV-001: Open Label Phase 2 Study of Tisotumab Vedotin for Locally Advanced or Metastatic Disease in Solid Tumors – Poster presentation, Saturday, June 1, 2019, 8:00 AM – 11:00 AM CDT

Phase 2 Trial of Tisotumab Vedotin in Platinum-Resistant Ovarian Cancer (innovaTV 208) – Poster presentation, Saturday, June 1, 2019, 1:15 PM – 4:15 PM CDT

Enapotamab vedotin:

First-in-human, dose-escalation, phase 1 trial to evaluate safety of anti-Axl antibody–drug conjugate enapotamab vedotin in solid tumors – Poster presentation, Saturday, June 1, 2019, 8:00 AM – 11:00 AM CDT

Ofatumumab (Submitted by Novartis):

Long-term follow-up of previously untreated patients (pts) with chronic lymphocytic leukemia (CLL) treated with ofatumumab and chlorambucil: Final analysis of the phase 3 COMPLEMENT 1 trial –Poster presentation, Monday, June 3, 8:00 AM – 11:00 AM CDT

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



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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.

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