Genmab to Highlight Advances Across Its Oncology Portfolio at the 2026 American Society of Clinical Oncology (ASCO) Annual Meeting and the European Hematology Association (EHA) 2026 Congress

Media Release

COPENHAGEN, Denmark; May 21, 2026

- **Twenty-three abstracts will highlight the versatility and strength of Genmab's portfolio and pipeline, including data from the comprehensive epcoritamab development program, including 4 oral presentations**
- **Three abstracts will show the ongoing clinical trial evaluations of the safety and efficacy of Genmab's investigational late-stage medicines, rinatabart sesutecan and petosemtamab**
- **Data evaluating epcoritamab will demonstrate outcomes across monotherapy, combination regimens and fixed-duration use, including outpatient administration and use in earlier lines of treatment**

Genmab A/S (Nasdaq: GMAB) announced today that 23 abstracts, including 20 abstracts evaluating epcoritamab, a subcutaneous T-cell engaging bispecific antibody, will be presented or published at the 2026 American Society of Clinical Oncology (ASCO) Annual Meeting in Chicago, IL, from May 29-June 2, and at the European Hematology Association (EHA) 2026 Congress in Stockholm, Sweden, from June 11-14.

Key presentations at ASCO and EHA will highlight data evaluating the potential utility of epcoritamab across multiple settings, including as a monotherapy, in combination regimens, in fixed-duration use and in earlier lines of therapy. Oral sessions will feature the first presentation of the full results from the Phase 3 EPCORE® DLBCL-1 trial comparing epcoritamab monotherapy to investigator's choice chemotherapy in patients with relapsed/refractory (R/R) large B-cell lymphoma (LBCL), as well as additional data from the Phase 3 EPCORE® FL-1 trial evaluating epcoritamab in combination with rituximab and lenalidomide (R^2) versus R^2 alone in patients with R/R follicular lymphoma (FL). Additional presentations will include real-world evidence and health economic and outcomes research data, as well as overviews of trials-in-progress evaluating late-stage medicines.

"This year at ASCO and EHA, Genmab will once again present data highlighting the depth and breadth of the epcoritamab development program, including encouraging results across multiple treatment settings for patients with B-cell malignancies. These findings underscore the versatility of epcoritamab as a monotherapy, in combination regimens and as a potential core therapy across the spectrum of B-cell malignancies," said Dr. Judith Klimovsky, Executive Vice President and Chief Development Officer of Genmab. "Additional presentations and publications at ASCO will further reflect our commitment to advancing other antibody-based therapeutics."

All abstracts accepted for presentation and publication have been published and may be accessed online via the ASCO Meeting Library and EHA Open Access Library.

Abstracts accepted for presentation at ASCO:

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Epcoritamab:

Abstract Number	Abstract Title	Type of Presentation	Date/Time of Presentation
7002*	Phase 2 trial of epcoritamab in combination with rituximab-mini CVP for older unfit/frail or anthracycline-ineligible adult patients with newly diagnosed diffuse large B-cell lymphoma: Interim futility analysis	Oral presentation	Saturday, May 30, 3:00 PM-6:00 PM CDT
7061	Epcoritamab (epcor) + chemoimmunotherapy (CIT) in patients (pts) with relapsed/refractory large B cell lymphoma (R/R LBCL) eligible for autologous stem cell transplant (ASCT): Pooled results from Arms 4 and 10 of EPCORE NHL-2	Poster	Monday, June 1, 9:00 AM-12:00 PM CDT
e19003	NHL-6: Phase 2 study of subcutaneous (SC) epcoritamab as outpatient treatment for 2L+ relapsed/refractory diffuse large B-cell lymphoma (R/R DLBCL)	Publication Only	NA

***Investigator-led trial**

Rinatabart sesutecan (Rina-S®):

Abstract Number	Abstract Title	Type of Presentation	Date/Time of Presentation
TPS5646	RAINFOL-03 (ENGOT-EN-31/GOG-3128): A phase 3, open-label, randomized study of Rinatabart sesutecan vs investigator's choice of chemotherapy in patients with endometrial cancer after platinum-based chemotherapy and programmed death ligand 1 inhibition	Poster	Monday, June 1, 9:00 AM-12:00 PM CDT
TPS5641	RAINFOL-04 (ENGOT-OV96/GOG-3134): A phase 3, open-label, randomized study of Rinatabart sesutecan plus standard of care (SOC) vs SOC as maintenance treatment after second-line platinum-based chemotherapy in patients with recurrent platinum-sensitive ovarian cancer	Poster	Monday, June 1, 9:00 AM-12:00 PM CDT

Petosemtamab:

Abstract Number	Abstract Title	Type of Presentation	Date/Time of Presentation
TPS8662	Petosemtamab plus pembrolizumab as first-line (1L) treatment of PD-L1 high metastatic non-small cell lung cancer (NSCLC): Global phase 2 trial	Poster	Sunday, May 31, 9:00 AM-12:00 PM CDT

Abstracts accepted for presentation at EHA:

Epcoritamab:

Abstract Number	Abstract Title	Type of Presentation	Date/Time of Presentation
S229	Clinically Relevant Subgroup Analysis from the Randomized Phase 3 EPCORE FL-1 Trial: Treatment (Tx) Effect of Epcoritamab with Lenalidomide and Rituximab (R2) in R/R Follicular Lymphoma (FL)	Oral Presentation	Thursday, June 11, 16:45-18:00 CEST
S235	Results From EPCORE DLBCL-1: Randomized Phase 3 Study of Epcoritamab (Epcor) Vs Investigator's Choice Chemoimmunotherapy (CIT) in Patients with Relapsed/Refractory Large B-cell Lymphoma (R/R LBCL)	Oral Presentation	Friday, June 12, 17:15-18:30 CEST
S153*	Fixed Duration Venetoclax Plus Epcoritamab Shows Favorable Tolerability and High Response Rates with Early Molecular	Oral Presentation	Sunday June 14, 11:00-12:15 CEST

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	Responses in R/R CLL/SLL: Interim Analysis of the Randomized HOVON 165/AETHER Trial		

*Investigator-led trial

PF977	Sustained Remissions Beyond 4 Years with Epcoritamab Monotherapy: Long-term Follow-up Results from the Pivotal EPCORE NHL-1 Trial in Patients with Relapsed or Refractory Large B-cell Lymphoma	Poster	Friday, June 12, 18:45-19:45 CEST
PF1007	Epcoritamab + R-mini-chop Results in 2-year Remissions and High MRD-negativity Rates in Elderly Patients with Newly Diagnosed DLBCL: Results from the EPCORE NHL-2 Trial	Poster	Friday, June 12, 18;45-19:45 CEST
PF1069	Reduced CD20 Expression and Intratumoral CD3+ T Cells Following Epcoritamab Treatment Are Associated with Progressive Disease in a Subset of Diffuse Large B-cell Lymphoma and Follicular Lymphoma	Poster	Friday, June 12, 18:45-19:45 CEST
PF1081	Pharmacodynamic Biomarkers Support the Clinical Benefit of Epcoritamab Plus Rituximab and Lenalidomide (R2) in Patients with Relapsed/Refractory Follicular Lymphoma (R/R FL): Analyses from EPCORE FL-1	Poster	Friday, June 12, 18:45-19:45 CEST
PS2035	Anchored Matching-adjusted Indirect Comparison of Epcoritamab, Lenalidomide, and Rituximab Vs Tafasitamab, Lenalidomide, and Rituximab in Relapsed/Refractory Follicular Lymphoma: EPCORE FL-1 Vs Inmind	Poster	Saturday, June 13, 18:45-19:45 CEST
PS2042	Comparative Effectiveness of Epcoritamab, Lenalidomide, and Rituximab in EPCORE FL-1 Vs Real-world Chemoimmunotherapy in Relapsed/Refractory Lymphoma	Poster	Saturday, June 13, 18:45-19:45 CEST
PS2052	Comparative Analyses of Epcoritamab in Combination with Lenalidomide and Rituximab Vs Obinutuzumab and Bendamustine in Relapsed/Refractory Follicular Lymphoma	Poster	Saturday, June 13, 18:45-19:45 CEST
PS2070	Epcoritamab + Chemoimmunotherapy in Patients with Relapsed/Refractory Large B-cell Lymphoma Eligible for Autologous Stem Cell Transplant: Pooled Results from Arms 4 And 10 of EPCORE NHL-2	Poster	Saturday, June 13, 18:45-19:45 CEST
PS2082	Fixed-duration Epcoritamab Monotherapy Induces High Response and MRD-negativity Rates in Elderly Patients with Newly Diagnosed Large B-cell Lymphoma and Comorbidities: Results from EPCORE DLBCL-3	Poster	Saturday, June 13, 18:45-19:45 CEST
PS2086	Epcoritamab in Relapsed/Refractory Diffuse Large B-cell Lymphoma (R/R DLBCL): Insights from the Real-world Epcoritamab Patient Characteristics and Outcomes Research (Real-epcor) Study	Poster	Saturday, June 13, 18:45-19:45 CEST
PS2497	Epcoritamab Plus Lenalidomide and Rituximab Improves or Preserves Health-related Quality of Life in Patients with Relapsed/Refractory Follicular Lymphoma Who Had High Symptom Burden or Adverse Events	Poster	Saturday, June 13, 18:45-19:45 CEST
NA	Epcoreal: A Prospective Observational Trial-in-progress of Epcoritamab in Patients with Relapsed/Refractory Diffuse Large B-cell Lymphoma and Follicular Lymphoma	Publication Only	NA
NA	Epcoritamab With Lenalidomide and Rituximab in Chinese Patients with Relapsed or Refractory Follicular Lymphoma: A Subgroup Analysis from the Phase 3 Epcore FL-1 Trial	Publication Only	NA

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NA	Cost Per Complete Responder for Epcoritamab + Lenalidomide and Rituximab (R2) Vs Tafasitamab + R2 in Relapsed or Refractory Follicular Lymphoma: A US Medicare Perspective	Publication Only	NA

The safety and efficacy of epcoritamab, Rina-S and petosemtamab have not been established for these investigational uses.

About Epcoritamab
Epcoritamab is an IgG1-bispecific antibody created using Genmab's proprietary DuoBody® technology and administered subcutaneously. Genmab's DuoBody-CD3 technology is designed to direct cytotoxic T cells selectively to elicit an immune response toward target cell types. Epcoritamab is designed to simultaneously bind to CD3 on T cells and CD20 on B cells and induces T-cell-mediated killing of CD20+ cells.[i]

Epcoritamab (approved under the brand name EPKINLY® in the U.S. and Japan, and TEPKINLY® in the EU) has received regulatory approval in certain lymphoma indications in several territories. Epcoritamab is being co-developed by Genmab and AbbVie as part of the companies' oncology collaboration. The companies will share commercial responsibilities in the U.S. and Japan, with AbbVie responsible for further global commercialization. Both companies will pursue additional international regulatory approvals for the investigational R/R FL indication and additional approvals for the R/R DLBCL indication.

Genmab and AbbVie continue to evaluate epcoritamab as a monotherapy, and in combination, across lines of therapy in a range of hematologic malignancies. This includes several ongoing Phase 3, open-label, randomized trials, among them a trial evaluating epcoritamab in combination with R-CHOP in adult patients with newly diagnosed DLBCL (NCT05578976), a trial evaluating epcoritamab in combination with lenalidomide compared to chemotherapy infusion in patients with R/R DLBCL (NCT06508658), and a trial evaluating epcoritamab in combination with lenalidomide and rituximab (R^2) compared to chemoimmunotherapy in patients with previously untreated FL (NCT06191744). The safety and efficacy of epcoritamab has not been established for these investigational uses. Please visit www.clinicaltrials.gov for more information.

Please see local country prescribing information for all labeled indication and safety information.

About Rinatabart Sesutecan (Rina-S; GEN1184)
Rina-S; GEN1184 is an investigational ADC. It is composed of a novel human monoclonal antibody directed at FRα, a hydrophilic protease-cleavable linker, and exatecan, a topoisomerase I inhibitor payload. The clinical development program for Rina-S continues to expand, with multiple ongoing Phase 3 studies in patients with ovarian and endometrial cancer, alongside evaluation in other tumor types with unmet needs. The safety and efficacy of rinatabart sesutecan have not been established. Please visit https://clinicaltrials.gov/ for more information.

About Petosemtamab (GEN1158)
Petosemtamab is an investigational bispecific antibody targeting the epidermal growth factor receptor (EGFR) and the leucine-rich repeat containing G-protein-coupled receptor 5 (LGR5). By engaging both receptors, petosemtamab is designed to inhibit EGFR signaling and trigger EGFR degradation selectively in LGR5+ cancer stem-like cells to support multiple anti-tumor mechanisms, including enhanced immune-mediated activity. It is currently being investigated in head and neck squamous cell carcinoma (HNSCC) and other solid tumors, including colorectal cancer (CRC) and non-small cell lung cancer (NSCLC). The safety and efficacy of petosemtamab have not been established. Please visit https://clinicaltrials.gov/ for more information.

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About Genmab
Genmab is an international biotechnology company dedicated to improving the lives of people with cancer and other serious diseases through innovative antibody medicines. For over 25 years, its passionate, innovative and collaborative team has advanced a broad range of antibody-based therapeutic formats, including bispecific antibodies, antibody–drug conjugates (ADCs), immune-modulating antibodies and other next-generation modalities. Genmab's science powers eight approved antibody medicines, and the company is advancing a strong late-stage clinical pipeline, including wholly owned programs, with the goal of delivering transformative medicines to patients.

Established in 1999, Genmab is headquartered in Copenhagen, Denmark, with international presence across North America, Europe and Asia Pacific. For more information, please visit Genmab.com and follow us on LinkedIn and X.

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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 preclinical 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.

[i] Engelberts PJ, Hiemstra IH, de Jong B, et al. DuoBody-CD3xCD20 induces potent T-cell-mediated killing of malignant B cells in preclinical models and provides opportunities for subcutaneous dosing. EBioMedicine. 2020;52:102625. DOI: 10.1016/j.ebiom.2019.102625.