Genmab to Present Results from Comprehensive Epcoritamab Development Program Highlighting Potential Across B-Cell Malignancies at the 2026 American Society of Hematology (ASH) Annual Meeting

Media Release

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- **Twenty-two abstracts underscore the versatility of epcoritamab as monotherapy and in combination across multiple disease subtypes and in earlier lines of therapy**
- **Six oral presentations include first data disclosure from the Phase 3 EPCORE® DLBCL-4 study evaluating epcoritamab in combination with lenalidomide in patients with relapsed or refractory (R/R) diffuse large B-cell lymphoma (DLBCL)**

Genmab A/S (Nasdaq: GMAB) announced today that 22 abstracts evaluating epcoritamab, a subcutaneous T-cell engaging bispecific antibody, will be presented or published at the 68th Annual Meeting and Exposition of the American Society of Hematology (ASH), taking place December 12-15, 2026, in New Orleans, Louisiana, and virtually.

The breadth of data to be shared at ASH reflects continued progress across the comprehensive epcoritamab development program, including its evaluation in earlier lines of therapy and combination regimens across multiple B-cell malignancies. Highlights include the first presentation of the primary results from the Phase 3 EPCORE® DLBCL-4 study evaluating fixed duration epcoritamab plus lenalidomide in patients with relapsed or refractory (R/R) diffuse large B-cell lymphoma (DLBCL). The results from EPCORE DLBCL-4 will also be highlighted in the ASH 2026 Press Program.

Additional presentations include updated subgroup analyses from the Phase 3 EPCORE FL-1 study evaluating epcoritamab in combination with rituximab and lenalidomide in R/R follicular lymphoma (FL), follow-up from various monotherapy and combination trials, and real-world evidence and health economic analyses that further inform the evolving clinical profile of epcoritamab.

“The range of studies accepted for presentation at this year’s American Society of Hematology Annual Meeting reflects our commitment to exploring the full utility of epcoritamab and reinforces our confidence in epcoritamab’s potential to become a core therapy across B-cell malignancies, as a monotherapy and in combination with other medicines across lines of treatment,” said Dr. Judith Klimovsky, Executive Vice President and Chief Development Officer of Genmab. “Together with our partner AbbVie, we look forward to sharing these findings with the global hematology community and further exploring what they could mean for patients.”

2026 R&D Update and ASH Data Review
On December 16 at 10:30 AM EST / 4:30 PM CET, Genmab will host its 2026 R&D Update and ASH Data Review. The event will be virtual and webcast live. Details, including the webcast link and registration will be available on www.genmab.com. This meeting is not an official program of the ASH Annual Meeting.

All regular abstracts accepted for presentation and publication have been published and may be accessed online on the ASH Website. Late-breaking abstracts will be posted online on Monday, December 7, 2026, at 9:00 AM EST.

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The following abstracts have been accepted for presentation at ASH:

Oral Presentations

Abstract Number	Abstract Title	Type of Presentation	Date/Time of Presentation
79*	ctDNA-directed intervention with epcoritamab alone or in combination with lenalidomide and rituximab is feasible in the early post-CAR-T population at high risk of relapse: Interim results from EpLCART	Oral presentation	December 12, 9:30-11:00 AM *(Presentation: 9:30-9:45 AM)*
105	Epcoritamab plus venetoclax leads to high response rates in relapsed or refractory chronic lymphocytic leukemia (R/R CLL): first results from Dose Cohorts 1 and 2 of the EPCORE CLL-1 trial	Oral presentation	December 12, 9:30-11:00 AM *(Presentation: 10:00-10:15 AM)*
166*	A phase II trial of rituximab and epcoritamab as first-line therapy for patients with high tumor burden follicular lymphoma: Results from initial and expansion cohorts	Oral presentation	December 12, 12:00-1:30 PM *(Presentation: 12:45-1:00 PM)*
167*	Phase II investigator-initiated trial of epcoritamab-lenalidomide in treatment naïve follicular lymphoma	Oral presentation	December 12, 12:00-1:30 PM *(Presentation: 1:00-1:15 PM)*
622	Durable benefit with fixed-duration epcoritamab + polatuzumab vedotin, rituximab, cyclophosphamide, doxorubicin, and prednisone (pola-R-CHP) as 1L treatment for patients with diffuse large B-cell lymphoma: 33-month follow-up from EPCORE NHL-5	Oral presentation	December 13, 12:00-1:30 PM (*Presentation: 12:45-1:00 PM*)
919	Epcoritamab plus lenalidomide (E-Len) compared to rituximab plus gemcitabine and oxaliplatin (R-GemOx): Primary results from the phase 3 EPCORE DLBCL-4 study in relapsed or refractory (R/R) diffuse large B-cell lymphoma (DLBCL)	Oral presentation	December 14, 2:45-4:15 PM *(Presentation: 2:45- 3:00 PM)*

***Investigator-led trial**

Poster Presentations

Abstract Number	Abstract Title	Type of Presentation	Date/Time of Presentation
1932	Efficacy and safety of epcoritamab + R-ICE in relapsed/refractory diffuse large B-cell lymphoma (R/R DLBCL): Updated results from EPCORE NHL-2	Poster	December 12, 5:30-7:30 PM
1933*	Epcoritamab plus gemcitabine, dexamethasone, and cisplatin (Epco-GDP) demonstrates high complete response rates in relapsed/refractory large B-cell lymphoma eligible for autologous stem cell transplantation or CAR T-cell therapy: Results from a multicenter phase II investigator initiated trial	Poster	December 12, 5:30-7:30 PM
1957*	A pilot study of epcoritamab with dose adjusted etoposide, cyclophosphamide, vincristine, doxorubicin, prednisone and rituximab (EPOCH-R) for treatment of	Poster	December 12, 5:30-7:30 PM

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	newly diagnosed aggressive B-cell non-Hodgkin lymphomas: An investigator initiated trial in progress.		
2109*	Rapid tumor depletion and normalization of myeloid and lymphoid immune landscapes underlie the potent activity of the epcoritamab plus venetoclax combination in relapsed/refractory CLL after targeted therapies	Poster	December 12, 5:30-7:30 PM
2716	Real-world effectiveness of epcoritamab in third-line or later (3L+) settings in patients with relapsed/refractory (R/R) diffuse large B-cell lymphoma (DLBCL): Insights from Real-EPCOR	Poster	December 12, 5:30-7:30 PM
3516	Epcoritamab retains robust cytotoxic activity in the setting of MS4A1 5-UTR splice variant V1 with markedly reduced CD20 protein levels	Poster	December 13, 6:00-8:00 PM
3517	Enhanced NK-cell and T-cell-dependent mechanisms underlie the benefit of epcoritamab combination with rituximab and lenalidomide in relapsed/refractory (R/R) B-cell non-Hodgkin lymphoma (NHL)	Poster	December 13, 6:00-8:00 PM
3519	Sustained minimal residual disease (MRD) negativity and favorable immune kinetics characterize long-term response to epcoritamab treatment in relapsed/refractory (R/R) diffuse large B-cell lymphoma (DLBCL)	Poster	December 13, 6:00-8:00 PM
3550	Epcoritamab plus lenalidomide and rituximab (E+R2) shows a consistent safety profile and efficacy benefit over R2 in patients with relapsed or refractory follicular lymphoma (R/R FL): Updated subgroup analyses from the randomized phase 3 EPCORE FL-1 trial	Poster	December 13, 6:00-8:00 PM
3719*	REPIFIR-AGR-GEL-23: A phase II study of epcoritamab in patients with relapsed or refractory large B-cell lymphoma in first relapse – a response-adapted approach guided by PET-CT and ctDNA-based MRD	Poster	December 13, 6:00-8:00 PM
4406	Cost-effectiveness of fixed-duration epcoritamab+rituximab and lenalidomide in patients with second-line or later relapsed/refractory follicular lymphoma in the United States	Poster	December 13, 6:00-8:00 PM
5441*	A phase II trial of epcoritamab prior to and after Anti-CD19 CAR T for relapsed or refractory large B cell lymphomas	Poster	December 14, 6:00-8:00 PM
5462*	A prospective clinical trial of prophylactic siltuximab to prevent epcoritamab-associated cytokine release syndrome in relapsed/refractory B-cell lymphoma	Poster	December 14, 6:00-8:00 PM
5463	Health-related quality of life (HRQoL) in relapsed or refractory diffuse large B-cell lymphoma (R/R DLBCL) with epcoritamab plus lenalidomide (E-Len) compared with rituximab plus gemcitabine and oxaliplatin (R-GemOx): Results of the EPCORE DLBCL-4 trial	Poster	December 14, 6:00-8:00 PM

***Investigator-led trial**

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e-Publications

Abstract Number	Abstract Title	Type of Presentation	Date/Time of Presentation
NA	Time and cost savings with fixed-duration epcoritamab + lenalidomide versus mosunetuzumab + polatuzumab for relapsed/refractory transplant- and CAR-T ineligible patients with diffuse large B-cell lymphoma	Publication only	NA
NA	Epcoritamab for relapsed or refractory large B-cell lymphoma in Japanese patients: interim analysis results from a real-world multicenter evaluation (J-REALEPCO)	Publication only	NA

The safety and efficacy of epcoritamab 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 more than 65 countries. Where approved, epcoritamab is a readily accessible therapy. Epcoritamab is being co-developed by Genmab and AbbVie as part of the companies' oncology collaboration. The companies 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 relapsed or refractory (R/R) follicular lymphoma (FL) indication and additional approvals for the R/R diffuse large B-cell lymphoma (DLBCL) indication.

Genmab and AbbVie continue to evaluate the use of epcoritamab as a monotherapy, and in combination, across lines of therapy in a range of hematologic malignancies. This includes 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.

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.

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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 or follow us on LinkedIn, *X*, Facebook and Instagram.

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[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.