

Abstract	Title	Presentation Timing (PT)
<u>Late-Breaking Presentation</u>		
Mantle Cell Lymphoma		
LBA6	Lack of Benefit of Autologous Hematopoietic Cell Transplantation (auto-HCT) in Mantle Cell Lymphoma (MCL) Patients (pts) in First Complete Remission (CR) with Undetectable Minimal Residual Disease (uMRD): Initial Report from the ECOG-ACRIN EA4151 Phase 3 Randomized Trial	Tuesday, December 10, 2024, 7:30-9 A.M.
<u>Oral Presentations</u>		
Acute Lymphoblastic Leukemia		
1 (Plenary session)	Blinatumomab Added to Chemotherapy Improves Disease-Free Survival in Newly Diagnosed NCI Standard Risk Pediatric B-Acute Lymphoblastic Leukemia: Results from the Randomized Children's Oncology Group Study AALL1731	Sunday, December 8, 2024, 2-4 P.M.
679	Safe and Effective Combination of Donor-Derived, Allogeneic CD19/CD22-CAR T Cells with Myeloablative Graft-Engineered Allo-HCT for High-Risk B-ALL	Sunday, December 8, 2024, 4:30-6 P.M.
727	Early Achievement of Deep Measurable Residual Disease (MRD) Negativity Identifies Patients with B-Cell Acute Lymphoblastic Leukemia (ALL) Who Have Excellent Long-Term Outcomes and Do Not Benefit from Allogeneic Stem Cell Transplant, Irrespective of Baseline High-Risk Cytomolecular Features	Monday, December 9, 2024, 10:30 A.M.-12 P.M.
837	Blinatumomab and Ponatinib for Adults with Newly Diagnosed Philadelphia Chromosome-Positive Acute Lymphoblastic Leukemia: Updated Results and Predictors of Relapse	Monday, December 9, 2024, 3:15-4:45 P.M.
924	TSC-100 and TSC-101 Demonstrate the Potential to Reduce Relapse Rates and Increase Relapse-Free Survival in Patients with AML, ALL, or MDS Undergoing Allogeneic HCT with Reduced Intensity Conditioning (RIC): Preliminary Results from the Phase 1 Alloha Trial	Monday, December 9, 2024, 4-5:30 P.M.
963	Obecabtagene autoleucel (obe-cel) for Adult Relapsed/Refractory B-Cell Acute Lymphoblastic Leukemia (R/R B-ALL): Deep Molecular Remission May Predict Better Outcomes	Monday, December 9, 2024, 5-6:30 P.M.

<u>966</u>	CD19-CAR T Cells As Definitive Consolidation for Older Adults with B-Cell Acute Lymphoblastic Leukemia in First Complete Remission: A Pilot Study	Monday, December 9, 2024, 5:45-7:15 P.M.
Multiple Myeloma		
<u>93</u>	A Phase 1/2 Trial of GLPG5101, a Fresh, Stem-like, Early Memory CD19 CAR T-Cell Therapy with a 7-Day Vein-to-Vein Time, for the Treatment of Relapsed/Refractory Non-Hodgkin Lymphoma	Saturday, December 7, 2024, 10-11:30 A.M.
<u>362</u>	Phase 3 Randomized Study of Daratumumab (DARA) + Bortezomib, Lenalidomide and Dexamethasone (VRd) Versus Alone in Patients with Transplant-Ineligible Newly Diagnosed Multiple Myeloma or for Whom Transplant Is Not Planned As Initial Therapy: Analysis of Minimal Residual Disease in the Cepheus Trial	Saturday, December 7, 2024, 4:15-5:45 P.M.
<u>363</u>	Implications of MRD Progression in Newly Diagnosed Multiple Myeloma (NDMM) Treated with Quadruplet Therapy and Autologous Stem Cell Transplantation	Saturday, December 7, 2024, 4:30-6 P.M.
<u>487</u>	Circulating Tumor Cells As a Biomarker to Identify High-Risk Transplant Eligible Myeloma Patients Treated with Bortezomib, Lenalidomide and Dexamethasone with or without Daratumumab during Induction/Consolidation, and Lenalidomide with or without Daratumumab during Maintenance: Results from the Perseus Study	Sunday, December 8, 2024, 9:30-11 A.M.
<u>497</u>	Phase I Study of Belantamab Mafodotin in Combination with Standard of Care in Transplant-Ineligible Newly Diagnosed Multiple Myeloma: Dreamm-9 Updated Interim Analysis	Sunday, December 8, 2024, 10:30-12 P.M.
<u>675</u>	Daratumumab Plus Lenalidomide (D-R) Versus Lenalidomide (R) Alone As Maintenance Therapy in Newly Diagnosed Multiple Myeloma (NDMM) after Transplant: Analysis of the Phase 3 Auriga Study Among Clinically Relevant Subgroups	Sunday, December 8, 2024, 5-6:30 P.M.
<u>770</u>	Isatuximab, Bortezomib, Lenalidomide, and Dexamethasone (Isa-VRd) in Patients with Newly Diagnosed Multiple Myeloma (NDMM): Analyses of Minimal Residual Disease (MRD) Negativity Dynamics in the Phase 3 Imroz Study	Monday, December 9, 2024, 10:45 A.M.-12:15 P.M.
<u>772</u>	Belantamab Mafodotin, Bortezomib, and Dexamethasone Vs Daratumumab, Bortezomib, and Dexamethasone in Relapsed/Refractory Multiple Myeloma: Overall Survival Analysis and Updated Efficacy Outcomes of the Phase 3 Dreamm-7 Trial	Monday, December 9, 2024, 11:15 A.M.-12:45 P.M.

<u>1031</u>	Phase 2 Registrational Study of Anitocabtagene Autoleucel for the Treatment of Patients with Relapsed and/or Refractory Multiple Myeloma: Preliminary Results from the IMMagine-1 Trial	Monday, December 9, 2024, 5:30-7 P.M.
<u>1032</u>	Ciltacabtagene Autoleucel (Cilta-cel) Vs Standard of Care (SoC) in Patients with Lenalidomide (Len)-Refractory Multiple Myeloma (MM) after 1-3 Lines of Therapy: Minimal Residual Disease (MRD) Negativity in the Phase 3 Cartitude-4 Trial	Monday, December 9, 2024, 5:45-7:15 P.M.
Mantle Cell Lymphoma		
<u>236</u>	Addition or Substitution of Acalabrutinib in Intensive Frontline Chemoimmunotherapy for Patients ≤70 Years Old with Mantle Cell Lymphoma: Outcomes of the 3-Arm Randomized Phase II Intergroup Trial ECOG-ACRIN EA4181	Saturday, December 7, 2024, 2:15-3:45 P.M.
<u>746</u>	MRD-Driven Time-Limited Therapy of Acalabrutinib and Lenalidomide Plus Rituximab (ALR) or Obinutuzumab (ALO) in Patients with Treatment-Naive Mantle Cell Lymphoma: Phase 2 Trial Outcomes with MRD and cfDNA Analyses	Monday, December 9, 2024, 10:45 A.M.-12:15 P.M.
<u>750</u>	Phase 1b/2 Study of Venetoclax, Ibrutinib, Prednisone, Obinutuzumab, and Lenalidomide (ViPOR) in Relapsed/Refractory and Treatment-Naive Mantle Cell Lymphoma: Preliminary Analysis of Safety, Efficacy, and Minimal Residual Disease	Monday, December 9, 2024, 11:45 A.M.-1:15 P.M.
Follicular Lymphoma		
<u>342</u>	Fixed-Duration Epcoritamab + R2 Drives Deep and Durable Responses in Patients with Relapsed or Refractory Follicular Lymphoma: 2-Year Follow-up from Arm 2 of the Epcore NHL-2 Trial	Saturday, December 7, 2024, 5:15-6:45 P.M.
Chronic Lymphocytic Leukemia		
<u>883</u>	Epcoritamab Monotherapy in Patients (Pts) with Relapsed or Refractory (R/R) Chronic Lymphocytic Leukemia (CLL): Results from CLL Expansion and Optimization Cohorts of Epcore CLL-1	Monday, December 9, 2024, 2:45-4:15 P.M.
<u>887</u>	Lisocabtagene Maraleucel (liso-cel) Combined with Ibrutinib (ibr) for Patients (pts) with Relapsed or Refractory (R/R) Chronic Lymphocytic Leukemia (CLL)/Small Lymphocytic Lymphoma (SLL): Primary Results from the Open-Label, Phase 1/2 Transcend CLL 004 Study	Monday, December 9, 2024, 3:45-5:15 P.M.

<u>1010</u>	Minimal Residual Disease (MRD)-Adapted Duration of Front-Line Venetoclax and Obinutuzumab Treatment for Fit Patients with Chronic Lymphocytic Leukemia (CLL)	Monday, December 9, 2024, 4:45-6:15 P.M.
<u>1011</u>	Combined Pirtobrutinib, Venetoclax, and Obinutuzumab As First-Line Treatment of Patients with Chronic Lymphocytic Leukemia (CLL)	Monday, December 9, 2024, 5-6:30 P.M.
Smoldering Multiple Myeloma		
<u>1027</u>	Early Safety and Efficacy of CAR-T Cell Therapy in Precursor Myeloma: Results of the CAR-PRISM Study Using Ciltacabtagene Autoleucel in High-Risk Smoldering Myeloma	Monday, December 9, 2024, 4:30-6 P.M.
Poster Presentations		
Acute Lymphoblastic Leukemia		
<u>1427</u>	Evaluating the Safety of Tyrosine Kinase Inhibitor Discontinuation in Adult Patients with Ph+ ALL Not Undergoing Allogeneic TransplantClinically Relevant Abstract	Saturday, December 7, 2024, 5:30-7:30 P.M.
<u>1432</u>	Interim Results of a Phase II Study Investigating Dasatinib and Inotuzumab Ozogamicin-Based Induction for Newly-Diagnosed Philadelphia-Chromosome Positive Acute Lymphoblastic Leukemia (Ph+ ALL)	Saturday, December 7, 2024, 5:30-7:30 P.M.
<u>1439</u>	Updated Results from a Phase II Study Hyper-CVAD, with or without Inotuzumab Ozogamicin, and Sequential Blinatumomab in Patients with Newly Diagnosed B-Cell Acute Lymphoblastic Leukemia	Saturday, December 7, 2024, 5:30-7:30 P.M.
<u>1441</u>	Low Intensity Mini-Hypercvd (mHCVD), Inotuzumab Ozogamicin (Ino) with/without Blinatumomab (Blina) in Older Patients with Newly Diagnosed Philadelphia Negative B-Cell Acute Lymphoblastic Leukemia (B-ALL): 10 Years Update	Saturday, December 7, 2024, 5:30-7:30 P.M.
<u>1450</u>	Improved Outcomes of Adult Patients with Philadelphia-like Acute Lymphoblastic Leukemia (Ph-Like ALL) Treated within an Integrated Leukemia/Transplant Program with Incorporation of Pediatric Inspired Regimens and Early Allogeneic Transplant	Saturday, December 7, 2024, 5:30-7:30 P.M.
<u>1465</u>	Clearance of Very Low Levels of Measurable Residual Disease with Blinatumomab Significantly Improves Outcomes in B-Cell Acute Lymphoblastic Leukemia	Saturday, December 7, 2024, 5:30-7:30 P.M.
<u>2837</u>	Enhanced Flow Cytometry and Next Generation Sequencing Assays for Residual B Lymphoblastic Leukemia (B-ALL) Reveal a Subset with Discordant Results Due to Leukemic Changes Post-Therapy	Sunday, December 8, 2024, 6-8 P.M.

<u>3470</u>	Cost-Effective Manufacture and Promising Initial Efficacy of huCART19 Cells Manufactured Using the Clinimacs Prodigy Platform	Sunday, December 8, 2024, 6-8 P.M.
<u>4194</u>	Phase II Study of Inotuzumab Ozogamicin for the Treatment of Measurable Residual Disease-Positive B-Cell Acute Lymphoblastic Leukemia: 3-Year Update	Monday, December 9, 2024, 6-8 P.M.
<u>4200</u>	NGS MRD Negativity on Day 28 after Brexu Cel in Adults with R/R ALL Is Associated with Favorable Progression Free Survival	Monday, December 9, 2024, 6-8 P.M.
<u>4202</u>	Concurrent Blinatumomab and Human Leukocyte Antigen-Mismatched Cellular Therapy in Patients with High-Risk B-Cell Acute Lymphoblastic Leukemia, a Phase I Prospective Cohort Study	Monday, December 9, 2024, 6-8 P.M.
Mantle Cell Lymphoma		
<u>1626</u>	High-Risk Subgroups and MRD: An Updated Analysis of the Phase 3 ECHO Trial of Acalabrutinib with Bendamustine/Rituximab in Previously Untreated Mantle Cell Lymphoma	Saturday, December 7, 2024, 5:30-7:30 P.M.
<u>1631</u>	Glofitamab Induces High Response Rates and Durable Remissions in Patients (Pts) with Heavily Pretreated Relapsed/Refractory (R/R) Mantle Cell Lymphoma (MCL), Including Those with a Poor Prognosis: Subgroup Results from a Phase I/II Study	Saturday, December 7, 2024, 5:30-7:30 P.M.
<u>1633</u>	Acalabrutinib, Umbralisib and Ublituximab Regimen (AU2) Demonstrates High Response Rate and Undetectable Molecular Minimal Residual Disease (MRD) in Patients (pts) with De Novo Mantle Cell Lymphoma (MCL)	Saturday, December 7, 2024, 5:30-7:30 P.M.
<u>3038</u>	Acalabrutinib with Rituximab Is Highly Effective First Line Treatment for Older Patients with Mantle Cell Lymphoma	Sunday, December 8, 2024, 6-8 P.M.
<u>4408</u>	Preliminary Efficacy and Safety of a Phase 1/2 Study of Acalabrutinib, Venetoclax, and Obinutuzumab in Patients with Relapsed/Refractory and Previously Untreated Mantle Cell Lymphoma (MAVO)	Monday, December 9, 2024, 6-8 P.M.
Diffuse Large B-Cell Lymphoma		
<u>1737</u>	Efficacy and Safety of Epcoritamab Monotherapy in Patients with Relapsed or Refractory LBCL Not Previously Exposed to CAR T: Subanalysis of the Epcore NHL-1 Trial	Saturday, December 7, 2024, 5:30-7:30 P.M.
<u>4480</u>	3-Year Update from the Epcore NHL-1 Trial: Epcoritamab Leads to Deep and Durable Responses in Relapsed or Refractory Large B-Cell Lymphoma	Monday, December 9, 2024, 6-8 P.M.

Chronic Lymphocytic Leukemia		
<u>1865</u>	Primary Endpoint Evaluation of a Multicenter, Phase 2 Study of Acalabrutinib, Venetoclax, Obinutuzumab (AVO) in a Population of Previously Untreated Patients with CLL Enriched for High-Risk Disease	Saturday, December 7, 2024, 5:30-7:30 P.M.
<u>1867</u>	Multicenter Phase II Trial of Zanubrutinib, Obinutuzumab, and Venetoclax (BOVen) in Treatment-Naïve Chronic Lymphocytic Leukemia: 5-Year Follow up, Retreatment Outcomes, and Impact of MRD Kinetics (Δ MRD400)	Saturday, December 7, 2024, 5:30-7:30 P.M.
<u>1871</u>	First-Line Ibrutinib Plus Venetoclax Vs Chlorambucil Plus Obinutuzumab in Elderly or Comorbid Patients (Pts) with Chronic Lymphocytic Leukemia (CLL): Glow Study 64-Month Follow-up (FU) and Adverse Event (AE)-Free Progression-Free Survival (PFS) Analysis	Saturday, December 7, 2024, 5:30-7:30 P.M.
<u>4633</u>	Lisocabtagene Maraleucel (liso-cel) in Patients (pts) with Relapsed or Refractory (R/R) Chronic Lymphocytic Leukemia (CLL)/Small Lymphocytic Lymphoma (SLL): Updated Follow-up of Transcend CLL 004	Monday, December 9, 2024, 6-8 P.M.
Multiple Myeloma		
<u>1976</u>	Isatuximab, Bortezomib, Lenalidomide, and Dexamethasone (Isa-VRd) in Patients with Newly Diagnosed Multiple Myeloma (NDMM) Not Eligible or with No Immediate Intent for Transplant: Long-Term Efficacy and Safety in a Phase 1b Study	Saturday, December 7, 2024, 5:30-7:30 P.M.
<u>3359</u>	Efficacy Outcomes By Minimal Residual Disease (MRD) Negativity in Patients with Relapsed or Refractory Multiple Myeloma Treated with Belantamab Mafodotin Plus Bortezomib and Dexamethasone Vs Daratumumab, Bortezomib, and Dexamethasone: Analysis from the Dreamm-7 Trial	Sunday, December 8, 2024, 6-8 P.M.
<u>3365</u>	Combination Regimens in MM Post Autologous Hematopoietic Cell Transplantation (AHCT) to Eliminate MRD Utilizing Iberdomide (COMMANDER): Results of Dose-Finding Component of a Phase 1b/2 Trial	Sunday, December 8, 2024, 6-8 P.M.
<u>3373</u>	Belantamab Mafodotin As Pre- and Post-Autologous Stem Cell Transplant (ASCT) Consolidation and Maintenance for Multiple Myeloma (MM) with < Complete Response after Induction: Interim Results of the Ongoing Phase 2 BLAST Study	Sunday, December 8, 2024, 6-8 P.M.

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3388	Idecabtagene Vicleucel (Ide-cel) in Patients (Pts) with Newly Diagnosed Multiple Myeloma (NDMM) with an Inadequate Response to Front-Line Autologous Stem Cell Transplantation (ASCT): KarMMa-2 Cohort 2c Extended Follow-up	Sunday, December 8, 2024, 6-8 P.M.
3784	Single Center Real World Experience of Talquetamab in Patients with Relapsed and Refractory Multiple Myeloma	Sunday, December 8, 2024, 6-8 P.M.
4731	Belantamab Mafodotin Plus Pomalidomide and Dexamethasone Vs Pomalidomide Plus Bortezomib and Dexamethasone in Patients with Relapsed/Refractory Multiple Myeloma: A Subset Analysis in Patients Who Have Received 1 Prior Line of Therapy Including Lenalidomide	Monday, December 9, 2024, 6-8 P.M.
4756	High-Risk Multiple Myeloma in Benefit (IFM 2020-05) Phase 3 Randomized Study of Isatuximab (Isa) Plus Lenalidomide and Dexamethasone (Rd) with Bortezomib Versus IsaRd in Patients with Newly Diagnosed Transplant Ineligible Multiple Myeloma (NDMM TI)	Monday, December 9, 2024, 6-8 P.M.
5028	MRD Testing in Multiple Myeloma: Modeling the Potential Clinical and Economic Outcomes Based on the Master Trial	Monday, December 9, 2024, 6-8 P.M.
Other		
2838	Immunoglobulin Sequencing Biologically Distinguishes B-Lymphoblastic Lymphoma from Acute Lymphoblastic Leukemia and Reveals a Spectrum of Disease Dissemination across Clinical Stages	Sunday, December 8, 2024, 6-8 P.M.
Follicular Lymphoma		
2958	Circulating Tumor DNA Predicts Time to First Treatment in Previously Untreated Follicular Lymphoma: Analysis from a Prospective Clonal Evolution Study	Sunday, December 8, 2024, 6-8 P.M.
3034	Clinical Outcomes of Patients with High-Risk Relapsed/Refractory Follicular Lymphoma Treated with Tisagenlecleucel: Phase 2 ELARA 4-Year Update	Sunday, December 8, 2024, 6-8 P.M.
Chronic Lymphocytic Leukemia		
3237	CRISTALLO: Results from a Phase III Trial of Venetoclax-Obinutuzumab Versus Fludarabine, Cyclophosphamide and Rituximab or Bendamustine-Rituximab in Patients with Untreated Chronic Lymphocytic Leukemia without Del(17p) or TP53 Mutations	Sunday, December 8, 2024, 6-8 P.M.

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<u>3247</u>	Real World Experience with Time Limited Venetoclax and Obinutuzumab (VO) for Frontline Treatment of CLL/SLL with MRD Determination by clonoSEQ®	Sunday, December 8, 2024, 6-8 P.M.
AL Amyloidosis		
<u>3302</u>	Minimal Residual Disease Testing in Relapsed Systemic AL Amyloidosis	Sunday, December 8, 2024, 6-8 P.M.
<u>3304</u>	Elranatamab in Patients with Daratumumab Relapsed and/or Refractory Light Chain Amyloidosis	Sunday, December 8, 2024, 6-8 P.M.
Smoldering Multiple Myeloma		
<u>3360</u>	Phase II Trial of Daratumumab, Bortezomib, Lenalidomide and Dexamethasone in High-Risk Smoldering Multiple Myeloma	Sunday, December 8, 2024, 6-8 P.M.