



## Rutgers Cancer Institute of New Jersey and RWJBarnabas Health to Showcase Expansive Portfolio of Hematology/Oncology Data at the 65th American Society of Hematology Annual Meeting and Exposition

36 oral and poster presentations will examine new advancements in the diagnosis, treatment, and management of blood cancers

New Brunswick, N.J., November 28, 2023 – Physician-scientists from Rutgers Cancer Institute of New Jersey and RWJBarnabas Health will present an extensive array of hematology/oncology data from their clinical research program at the 65th American Society of Hematology (ASH) Annual Meeting and Exposition, being held in San Diego, California (and virtually) from December 9-12, 2023. A total of 36 abstracts have been accepted, comprising clinical data and analyses that advance the understanding, treatment, and prognosis of blood cancers such as lymphoma, leukemia, and myeloma. Rutgers Cancer Institute of New Jersey, in partnership with RWJBarnabas Health, is New Jersey's only National Cancer Institute-designated Comprehensive Cancer Center and the leading cancer program in the state.

"Our devoted and highly esteemed team of cancer specialists and researchers at Rutgers Cancer Institute of New Jersey and RWJBarnabas Health remain dedicated to pioneering advancements that transform treatments, patient care, and outcomes by leveraging innovation and evolving novel therapies," said Matthew Matasar, MD, MS, Chief of Blood Disorders at Rutgers Cancer Institute of New Jersey and RWJBarnabas Health. "As the leading cancer program in the state, our commitment to advancing oncology care is unwavering. We strive to make a meaningful difference in the lives of those affected by cancer, and through innovative science and research, we can uncover new insights that may transform the landscape of cancer care in the future."

Highlights of the accepted abstracts include the following oral and poster presentations:

• Data from a randomized phase 3 trial conducted by the National Clinical Trials Network evaluating the tolerability and progression-free survival rate of Nivolumab-AVD compared to Bv-AVD in patients aged ≥60 with newly diagnosed advanced-stage Hodgkin lymphoma (AS-HL). The primary endpoint of the study was progression-free survival (PFS), and secondary endpoints included overall survival (OS), event-free survival (EFS), and detailed toxicity and safety events. Response and progression were assessed by investigators using 2014 Lugano Classification.

- Data from a phase 3 trial evaluating the progression-free survival and toxicity with Nivolumab-AVD compared to Bv-AVD in pediatric patients aged ≥12 years with stage 3-4 Classic Hodgkin Lymphoma (cHL). The trial was led by SWOG and conducted by the National Clinical Trials Network. The primary endpoint was progression-free survival (PFS), and secondary endpoints included overall survival (OS), event-free survival (EFS), and safety.
- Evaluation of the A-HIPI model in generating risk groups with input on strengths and limitations for patients with advanced stage classical Hodgkin lymphoma (AS-HL). This prognostic model, developed and validated by the Hodgkin Lymphoma International Study for Individual Care (HoLISTIC) Consortium, generates the individualized probability of a progression-free survival (PFS) event or death within the first 5 years from diagnosis in patients based on continuous variables.
- Development and validation of the modern-day model, E-HIPI, in the prediction of
  progression-free survival of early-stage Hodgkin lymphoma (E-HL) within the first five years
  since diagnosis. The model incorporates detailed individual patient data from international
  clinical trials and prospective registry data that were standardized, normalized, and harmonized
  as part of the HoLISTIC Consortium. The primary outcome was progression-free survival
  (PFS).
- Data from a clinical trial evaluating the feasibility, safety, and efficacy of home-based intravenous administration of trans-retinoic acid (ATRA) and arsenic trioxide in the treatment of Acute Promyelocytic Leukemia (APL). Patients were given the option of receiving at-home treatment through a partnership with Qualitas Specialty Pharmacy and were evaluated for any adverse effects.

The full list of presentations at this year's ASH Annual Meeting and Exposition follows:

Oral Presentations				
Abstract and	Title	Presentation Date/Time		
Session No.				
Abstract 181	Nivolumab-AVD Is Better Tolerated and	Saturday, December 9, 2023:		
(Session 624)	Improves Progression-Free Survival	2:00 PM PT		
	Compared to Bv-AVD in Older Patients			
	(Aged ≥60 Years) with Advanced Stage			
	Hodgkin Lymphoma Enrolled on SWOG			
	S1826 Clinically Relevant Abstract			
Abstract 308	Improved Survival of R/R Double Hit/Triple	Saturday, December 9, 2023:		
(Session 627)	Hit Lymphoma in the Era of CD19 Chimeric	4:15 PM PT		
	Antigen T Cell (CART) Therapy			
Abstract 382	Treatment Patterns and Outcomes for Patients	Saturday, December 9, 2023:		
(Session 905)	with Classic Hodgkin Lymphoma (cHL) and	4:45 PM PT		
	Cardiomyopathy with Low Ejection Fraction			
	(EF): Real-World Evidence (RWE) from 16			
	US Academic Centers			

Abstract 497	Outcomes of Patients with Richter	Sunday, December 10, 2023:
(Session 905)	Transformation without Prior	10:30 AM PT
	Chemoimmunotherapy for CLL/SLL: An	
	International Multicenter Retrospective Study	
Abstract 603	Mosunetuzumab Monotherapy Continues to	Sunday, December 10, 2023:
(Session 623)	Demonstrate Durable Responses in Patients	5:00 PM PT
	with Relapsed and/or Refractory Follicular	
	Lymphoma after ≥2 Prior Therapies: 3-Year	
	Follow-up from a Pivotal Phase II Study	
Abstract 607	Results from an Intergroup Randomized	Sunday, December 10, 2023:
(Session 624)	Phase II Study of the Combinations of	4:30 PM PT
,	Ipilimumab, Nivolumab and Brentuximab	
	Vedotin in Patients with Relapsed/Refractory	
	Classic Hodgkin Lymphoma: A Trial of the	
	ECOG-ACRIN Research Group (E4412)	
Abstract 610	Progression-Free Survival (PFS) and Toxicity	Sunday, December 10, 2023:
(Session 624)	with Nivolumab-AVD Compared to	5:15 PM PT
	Brentuximab Vedotin-AVD in Pediatric	
	Advanced Stage (AS) Classic Hodgkin	
	Lymphoma (cHL), Results of SWOG S1826	
Abstract 614	Mosunetuzumab Monotherapy Demonstrates	Sunday, December 10, 2023:
(Session 627)	Activity and a Manageable Safety Profile in	4:45 PM PT
	Patients with Relapsed or Refractory	
	Richter's Transformation	
Abstract 981	Pirtobrutinib in Relapsed/Refractory (R/R)	Monday, December 11, 2023:
(Session 623)	Mantle Cell Lymphoma (MCL) Patients with	5:00 PM PT
	Prior cBTKi: Safety and Efficacy Including	
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	High-Risk Subgroup Analyses from the Phase	
Dostor Prosents	High-Risk Subgroup Analyses from the Phase 1/2 BRUIN Study	
Poster Presenta	High-Risk Subgroup Analyses from the Phase 1/2 BRUIN Study	Presentation Date/Time
Abstract and	High-Risk Subgroup Analyses from the Phase 1/2 BRUIN Study	Presentation Date/Time
Abstract and Session No.	High-Risk Subgroup Analyses from the Phase 1/2 BRUIN Study  tions  Title	
Abstract and Session No. Abstract 1079	High-Risk Subgroup Analyses from the Phase 1/2 BRUIN Study  ations  Title  Congenital Dyserythropoietic Anemia Type	Saturday, December 9, 2023,
Abstract and Session No.	High-Risk Subgroup Analyses from the Phase 1/2 BRUIN Study  Itions  Title  Congenital Dyserythropoietic Anemia Type II: An Update from the Congenital	
Abstract and Session No. Abstract 1079	High-Risk Subgroup Analyses from the Phase 1/2 BRUIN Study  tions  Title  Congenital Dyserythropoietic Anemia Type II: An Update from the Congenital Dyseryhtropoietic Anemia Registry of North	Saturday, December 9, 2023,
Abstract and Session No.  Abstract 1079 (Session 101)	High-Risk Subgroup Analyses from the Phase 1/2 BRUIN Study  Itions  Title  Congenital Dyserythropoietic Anemia Type II: An Update from the Congenital Dyseryhtropoietic Anemia Registry of North America (CDAR)	Saturday, December 9, 2023, 5:30 PM-7:30 PM PT
Abstract and Session No.  Abstract 1079 (Session 101)  Abstract 1404	High-Risk Subgroup Analyses from the Phase 1/2 BRUIN Study  Itions  Title  Congenital Dyserythropoietic Anemia Type II: An Update from the Congenital Dyseryhtropoietic Anemia Registry of North America (CDAR)  Acetyl Transferase EP300 Deficiency Leads	Saturday, December 9, 2023, 5:30 PM-7:30 PM PT  Saturday, December 9, 2023,
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Abstract and Session No.  Abstract 1079 (Session 101)  Abstract 1404 (Session 603)  Abstract 1551	High-Risk Subgroup Analyses from the Phase 1/2 BRUIN Study  Itions  Title  Congenital Dyserythropoietic Anemia Type II: An Update from the Congenital Dyserythropoietic Anemia Registry of North America (CDAR)  Acetyl Transferase EP300 Deficiency Leads to Chronic Replication Stress in Adult T-Cell Leukemia/Lymphoma	Saturday, December 9, 2023, 5:30 PM-7:30 PM PT  Saturday, December 9, 2023, 5:30 PM-7:30 PM PT  Saturday, December 9, 2023, 5:30 PM-7:30 PM PT
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Abstract and Session No.  Abstract 1079 (Session 101)  Abstract 1404 (Session 603)  Abstract 1551 (Session 616)	High-Risk Subgroup Analyses from the Phase 1/2 BRUIN Study  Itions  Title  Congenital Dyserythropoietic Anemia Type II: An Update from the Congenital Dyserythropoietic Anemia Registry of North America (CDAR)  Acetyl Transferase EP300 Deficiency Leads to Chronic Replication Stress in Adult T-Cell Leukemia/Lymphoma  A Phase Ib/II Study Evaluating Navitoclax after Failure of Hypomethylating Agent and Venetoclax for Treatment of Relapsed or Refractory High-Risk Myelodysplastic Syndrome	Saturday, December 9, 2023, 5:30 PM-7:30 PM PT  Saturday, December 9, 2023, 5:30 PM-7:30 PM PT  Saturday, December 9, 2023, 5:30 PM-7:30 PM PT
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Abstract and Session No.  Abstract 1079 (Session 101)  Abstract 1404 (Session 603)  Abstract 1551 (Session 616)	High-Risk Subgroup Analyses from the Phase 1/2 BRUIN Study  Title  Congenital Dyserythropoietic Anemia Type II: An Update from the Congenital Dyseryhtropoietic Anemia Registry of North America (CDAR)  Acetyl Transferase EP300 Deficiency Leads to Chronic Replication Stress in Adult T-Cell Leukemia/Lymphoma  A Phase Ib/II Study Evaluating Navitoclax after Failure of Hypomethylating Agent and Venetoclax for Treatment of Relapsed or Refractory High-Risk Myelodysplastic Syndrome  Alterations in Immune Cell Composition during First-Line Therapy with Mosunetuzumab for Follicular or Marginal Zone Lymphoma  Pirtobrutinib, a Highly Selective, Non-	Saturday, December 9, 2023, 5:30 PM-7:30 PM PT  Saturday, December 9, 2023, 5:30 PM-7:30 PM PT
Abstract and Session No.  Abstract 1079 (Session 101)  Abstract 1404 (Session 603)  Abstract 1551 (Session 616)  Abstract 1651 (Session 622)	High-Risk Subgroup Analyses from the Phase 1/2 BRUIN Study  Itions  Title  Congenital Dyserythropoietic Anemia Type II: An Update from the Congenital Dyseryhtropoietic Anemia Registry of North America (CDAR)  Acetyl Transferase EP300 Deficiency Leads to Chronic Replication Stress in Adult T-Cell Leukemia/Lymphoma  A Phase Ib/II Study Evaluating Navitoclax after Failure of Hypomethylating Agent and Venetoclax for Treatment of Relapsed or Refractory High-Risk Myelodysplastic Syndrome  Alterations in Immune Cell Composition during First-Line Therapy with Mosunetuzumab for Follicular or Marginal Zone Lymphoma  Pirtobrutinib, a Highly Selective, Non-Covalent (Reversible) BTK Inhibitor in	Saturday, December 9, 2023, 5:30 PM-7:30 PM PT
Abstract and Session No.  Abstract 1079 (Session 101)  Abstract 1404 (Session 603)  Abstract 1551 (Session 616)  Abstract 1651 (Session 622)	High-Risk Subgroup Analyses from the Phase 1/2 BRUIN Study  Title  Congenital Dyserythropoietic Anemia Type II: An Update from the Congenital Dyseryhtropoietic Anemia Registry of North America (CDAR)  Acetyl Transferase EP300 Deficiency Leads to Chronic Replication Stress in Adult T-Cell Leukemia/Lymphoma  A Phase Ib/II Study Evaluating Navitoclax after Failure of Hypomethylating Agent and Venetoclax for Treatment of Relapsed or Refractory High-Risk Myelodysplastic Syndrome  Alterations in Immune Cell Composition during First-Line Therapy with Mosunetuzumab for Follicular or Marginal Zone Lymphoma  Pirtobrutinib, a Highly Selective, Non-Covalent (Reversible) BTK Inhibitor in Relapsed / Refractory Marginal Zone	Saturday, December 9, 2023, 5:30 PM-7:30 PM PT  Saturday, December 9, 2023, 5:30 PM-7:30 PM PT
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Abstract 1727	Multicenter Pilot Trial of Intrathecal	Saturday, December 9, 2023,
(Session 626)	Liposomal Cytarabine Combined with FAB	5:30 PM-7:30 PM PT
	Chemoimmunotherapy with Reduced	
	Doxorubicin in CAYA with Mature De-Novo	
	B-NHL	
Abstract 1737	Pirtobrutinib in Richter Transformation:	Saturday, December 9, 2023,
(Session 626)	Updated Efficacy and Safety Results with 18-	5:30 PM-7:30 PM PT
,	Month Median Survival Follow-up from the	
	Phase 1/2 BRUIN StudyClinically Relevant	
	Abstract	
Abstract 1738	Immune Reconstitution and Infection Patterns	Saturday, December 9, 2023,
(Session 627)	Following CAR T-Cell Therapy in Patients	5:30 PM-7:30 PM PT
(30381011 027)		3.30 1 WI-7.30 1 WI I I
	with Aggressive LymphomaClinically	
	Relevant Abstract	
Abstract 1908	Treatment Effectiveness with Venetoclax-	Saturday, December 9, 2023,
(Session 642)	Based Therapy after Bruton Tyrosine Kinase	5:30 PM-7:30 PM PT
	Inhibitors in Chronic Lymphocytic Leukemia:	
	An International Real-World Study	
Abstract 2085	CD22 TCR-Engineered T Cells Exert Anti-	Saturday, December 9, 2023,
(Session 703)	Leukemia Cytotoxicity without Causing	5:30 PM-7:30 PM PT
	Inflammatory Responses	
Abstract 2110	Preliminary Results of Nathali-01: A First-in-	Saturday, December 9, 2023,
(Session 704)	Human Phase I/IIa Study of UCART20x22, a	5:30 PM-7:30 PM PT
(36881011 /04)		3.30 FWI-7.30 FWI F I
	Dual Allogeneic CAR-T Cell Product	
	Targeting CD20 and CD22, in Relapsed or	
	Refractory (R/R) Non-Hodgkin Lymphoma	
	(NHL)	
Abstract 2185	Outcomes after RIC and Abatacept-Based	Saturday, December 9, 2023,
(Session 722)	Acute and Chronic Gvhd Prophylaxis in	5:30 PM-7:30 PM PT
	Allogeneic Transplantation for Sickle Cell	
	Disease – Can Calcineurin Inhibitor Use be	
	Curtailed?	
Abstract 2357	No Place like Home: Home-Based	Saturday, December 9, 2023,
(Session 903)	Intravenous Arsenic Trioxide for the	5:30 PM-7:30 PM PT
(Session 703)	Treatment of Acute Promyelocytic Leukemia	3.30 1 WI-7.30 1 WI I I
A1	(APL)	G-41 D 1 0 2022
Abstract 2407	Exploration of Language As a Barrier to the	Saturday, December 9, 2023,
(Session 905)	Assessment and Management of CAR T-Cell	5:30 PM-7:30 PM PT
	Therapy Associated Toxicities	
Abstract 3052	GLOBRYTE: A Phase III, Open-Label,	Sunday, December 10, 2023,
(Session 623)	Multicenter, Randomized Trial Evaluating	6:00 PM-8:00 PM PT
	Glofitamab Monotherapy in Patients with	
	Relapsed or Refractory Mantle Cell	
	Lymphoma	
Abstract 3058	Development and Validation of the Early-	Sunday, December 10, 2023,
(Session 624)	Stage Hodgkin Lymphoma (HL) International	6:00 PM-8:00 PM PT
(50551011 024)	Prognostication Index (E-HIPI): A Report	0.0011111
	, , ,	
	from the Hodgkin Lymphoma International	
	Study for Individual Care (HoLISTIC)	
	Consortium	
Abstract 3067	Identification of Risk Categories from the	Sunday, December 10, 2023,
(Session 624)	Advanced-Stage Hodgkin International	6:00 PM-8:00 PM PT
1	Prognostic Index (A-HIPI) Model: A Detailed	

	A 1 ' C 4 TT 11' T 1	T
	Analysis from the Hodgkin Lymphoma	
	International Study for Individual Care	
11 2004	(HoLISTIC) Consortium	G 1 D 1 10 2022
Abstract 3084	AHOD2131: A Randomized Phase 3	Sunday, December 10, 2023,
(Session 624)	Response-Adapted Trial Comparing Standard	6:00 PM-8:00 PM PT
	Therapy with Immuno-Oncology Therapy for	
	Children and Adults with Newly Diagnosed	
	Stage I and II Classic Hodgkin Lymphoma	
Abstract 3359	Health-Related Quality of Life (HRQoL)	Sunday, December 10, 2023,
(Session 652)	Among Patients with Triple-Class Exposed	6:00 PM-8:00 PM PT
	Relapsed/Refractory Multiple Myeloma	
	(RRMM) Treated with Linvoseltamab in	
	Linker-MM1: Interim Assessment up to 36	
	Weeks of TreatmentClinically Relevant	
	Abstract	
Abstract 4455	Age-Based Validation of the Advanced-Stage	Monday, December 11, 2023,
(Session 624)	Hodgkin Lymphoma International Prognostic	6:00 PM-8:00 PM PT
	Index (A-HIPI) in a Real-World Danish	
	Study: Suboptimal Performance in Older	
	Patients	
Abstract 4461	Odronextamab Demonstrates Durable	Monday, December 11, 2023,
(Session 626)	Complete Responses in Patients with Diffuse	6:00 PM-8:00 PM PT
	Large B-Cell Lymphoma (DLBCL)	
	Progressing after CAR-T Therapy: Outcomes	
	from the ELM-1 Study	
Abstract 4746	Patterns of Response to 200 Mg	Monday, December 11, 2023,
(Session 653)	Linvoseltamab in Patients with	6:00 PM-8:00 PM PT
	Relapsed/Refractory Multiple Myeloma:	
	Longer Follow-Up of the Linker-MM1 Study	
Abstract 4901	Abatacept-Prophylaxis Based Haploidentical	Monday, December 11, 2023,
(Session 721)	Transplantation May Allow Sustained	6:00 PM-8:00 PM PT
	Engraftment and Offset Gvhd in Non-	
	Malignant Disorders	
<b>Education Prog</b>		
N/A	MRD Directed Therapy in CLL- Ready for	Saturday, December 9, 2023,
	Primetime?	2:00 PM-3:15 PM PT
N/A	Hodgkin Lymphoma Treatment for Older	Saturday, December 9, 2023,
	Persons in the Modern Era	4:00 PM-5:15 PM PT
N/A	How is the Management Paradigm Evolving	Saturday, December 9, 2023:
	for Hodgkin Lymphoma in 2023?	4:00 PM-5:15 PM PT
Program: Special-Interest Sessions		
Session 2	General Session 2: Making the Most of	Sunday, December 10, 2023,
	<u>Virtual Teaching</u>	7:30 AM-9:30 AM PT

## **About Rutgers Cancer Institute of New Jersey**

As New Jersey's only National Cancer Institute-designated Comprehensive Cancer Center, Rutgers Cancer Institute, together with RWJBarnabas Health, offers the most advanced cancer treatment options, including bone marrow transplantation, proton therapy, CAR T-cell therapy and complex surgical procedures. Along with clinical trials and novel therapeutics such as precision medicine and immunotherapy – many of which are not widely available – patients have access to these cutting-

edge therapies at Rutgers Cancer Institute of New Jersey in New Brunswick, Rutgers Cancer Institute of New Jersey at University Hospital in Newark, as well as through RWJBarnabas Health facilities. To make a tax-deductible gift to support the Cancer Institute of New Jersey, call 848-932-8013 or visit <a href="https://www.cinj.org/giving">www.cinj.org/giving</a>.

## For journalists – contact:

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 $For \ patient \ appointments/inquiries-contact:$ 

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