

CONFERENCE PROGRAM



University of Nebraska
Medical Center
Nebraska Medicine

2024 PAN PACIFIC **LYMPHOMA** CONFERENCE

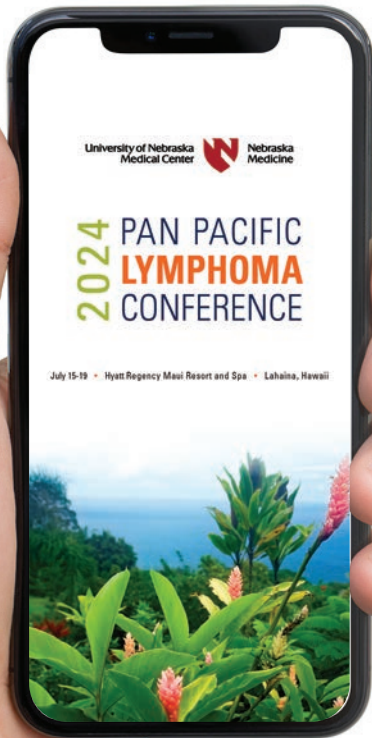
July 15-19 • Hyatt Regency Maui



Connect with us on  and  using #PPLC24 #IamUNMC

DOWNLOAD THE EVENT APP

Make the most of your conference experience by downloading the **FREE** mobile app!



1. Download and install the eventScribe app

- Scan the QR code or search for **eventScribe** in the Apple App Store or in Google Play.
- Enter **PPLC 2024** in the Search for an Event field.



2. Login to the conference app

- To start using the conference app, select **“Create Account”** and type in **PPLC24** as the **event code**.
- Enter **your name** and the **email address** that you used to register for this conference.

3. Check your email for your access code

- Your access code is your password to login to the conference app.

Laptop and Desktop Users

1. Visit the conference portal

- Go to www.panpacificlymphoma.eventscribe.net

2. Login/Create an account

- If this is the first time you are accessing the portal, click **Sign Up** to create an account.
- Enter **your name** and the **email address** that you used to register for this conference. Use **PPLC24** as the **event code**.

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30 Years of Educational Excellence

Welcome to the 2024 Pan Pacific Lymphoma Conference!

Since its inception in 1994, PPLC has grown and become a premier international conference on the latest advances in lymphoma. This year's conference is a hybrid conference with a live in-person activity at the Hyatt Regency Maui Resort and Spa, Hawaii and a virtual option for attendees who are not able to travel. We are very happy to welcome over 670 attendees!

For your easy reference and convenience, we encourage you to download the conference mobile app or visit www.panpacificlymphoma.eventscribe.net. Please note that all reference to conference days and times follow Hawaii Standard Time (HST).

Here's what you can expect for the week:

- **Day 1 (Monday):** We will feature an Interprofessional Education Day, with various presentations for the interprofessional lymphoma team members with sessions covering The Day-to-Day Treating of Lymphoma Patients, How to Increase Your Pl'nness and Practitioner Wellness.
- **Days 2-5 (Tuesday through Friday):** Expert faculty will present in scientific sessions covering Future Assessment of Lymphoma, Rock the Vote: Choosing Initial Therapy Wisely, Lymphoplasmacytic Lymphoma and Plasma Cell Dyscrasias, Just in "Case" You Asked Why I Did That, The Masked Lymphoma Doctor, Royal Lymphoma Rumble, Who Knows, Management of Orphan Diseases: When and How, and The Class of the Future is Here. Caron A. Jacobson, MD, MMSc will present the James O. Armitage Lymphoma Clinical Investigator Award Lecture and Elaine S. Jaffe, MD will deliver the Oliver Press Memorial Lecture, in memory of our esteemed colleague, Oliver W. Press, MD, PhD.
- **Days 2-4 (Tuesday through Thursday) afternoons:** Join us for the Ask the Experts sessions where we will discuss Indolent Non-Hodgkin Lymphoma (iNHL), DLBCL, cHL, PTCL and Waldenström Macroglobulinemia/Multiple Myeloma. Also, on Day 2/Tuesday, we will have our fast-paced Bierman's Brain Trivia – ERAS Edition with four teams competing for Day 3/Wednesday's Bierman's Brain Trivia Pineapple Cup Finals.
- **Days 1-5 (Monday through Friday):** If you are joining us in-person, please come visit the exhibits in the Grand Promenade.
- **Days 1-5 (Monday through Friday):** The conference e-poster gallery detailing the latest research in lymphoma and transplantation is available via the conference mobile app and online.
- **Day 2 (Tuesday) evening:** If you are attending in-person, please join us for the Welcome Reception from 7-9 p.m. in the Napili Gardens/Lawn where you can meet and network with other conference attendees. Light hors d'oeuvres and beverages will be served. Paid/registered guests are welcome to attend.
- **Day 4 (Thursday) evening:** In-person attendees are encouraged to join the Conference Dinner from 7-9 p.m. in Halona Kai. Dinner and beverages will be served. Paid/registered guests are welcome to attend.

Friendly reminders if you are attending in-person:

- Please be sure that you and your paid guests wear your conference badge at all times.
- The attire for this conference is casual; layered clothing is recommended as temperature in meeting rooms can fluctuate. A light jacket or sweater for outdoor evening activities is suggested. Leave your ties in your room!

If you are on X or Instagram, we encourage you to tag your photos and posts [#PPLC24](https://twitter.com/PPLC24) and [#lamUNMC](https://twitter.com/lamUNMC).

Once again, thank you for joining the 2024 Pan Pacific Lymphoma Conference. **Aloha!**

Conference Directors



James O. Armitage, MD
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Division of Oncology and Hematology
Department of Internal Medicine
University of Nebraska Medical Center



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Conference Sponsors

University of Nebraska Medical Center (Accredited Provider)

As Nebraska's only public academic health sciences center, the University of Nebraska Medical Center (UNMC) is committed to the education of a 21st century health care work force, to finding cures and treatments for devastating diseases, to providing the best care for patients, and to serving our state and its communities through award-winning outreach. UNMC is also committed to embracing the richness of diversity and is a major economic engine for the state of Nebraska. Led by Chancellor Jeffrey P. Gold, MD, UNMC has six colleges, two institutes, and a graduate studies program, serving nearly 4,300 students in more than two dozen programs. As an academic health science center, UNMC offers patients world-class health care, backed by the latest research innovations and practiced by faculty training the next generation of health care providers. With our hospital partner, Nebraska Medicine, UNMC provides services in about 50 specialties, including cancer, neurosciences, heart disease and others. Through this unique combination of academic, scientific, and health care experience, UNMC transforms the discoveries of the laboratory and theory of the classroom into breakthroughs for health.

The mission of the UNMC Continuing Education Programs is to provide specialized and inter-professional educational activities and support innovative research that facilitate individual skills and team-based performance that improves outcomes for patients and communities. Accordingly, UNMC's continuing education efforts aim to enhance the knowledge, skills, attitudes, competence and performance of health care professionals, ultimately improving patient outcomes and community health. This mission statement drives and unifies the continuing education efforts of all colleges and departments at UNMC.

The UNMC Center for Continuing Education (UNMC CCE) is one of this country's oldest accredited providers of continuing education to physicians, started in 1958. By 1967, UNMC CCE, was one of the first seven medical school programs to be ACCME accredited. Throughout its history, UNMC CCE's level and number of activities have grown in certified activities offered, variety of topics, delivery modes, and diversity of audience participants. UNMC CCE offers a broad range of activities from primary care issues to the latest developments in medicine and draws attendees throughout the region, nationally and internationally.

The UNMC College of Nursing Continuing interProfessional Development and Innovation (CiPDI) has been serving the profession since 1967. Through the creation of engaging continuing nursing education products using state-of-the-art technologies and innovative educational modalities, we strive to improve patient outcomes in Nebraska and around the globe. Our highly developed technical capabilities include instructional design and module development, video streaming, learning activity hosting, custom web development, visual design, dynamic web pages, and data collection, storage, and analysis. The UNMC CON CiPDI is approved by the California Board of Registered Nursing, and is able to take accredited programs into all 50 states and beyond.

In pursuit of our collective mission, UNMC CCE and UNMC CON CiPDI work collaboratively to sponsor multi-professional educational activities designed to improve the performance of the health care team and thus patient outcomes. To forward an integrated team-based approach and to facilitate multi-disciplinary and specialized training, UNMC achieved Joint Accreditation provider status effective through November 2024.

UNMC Department of Internal Medicine - Division of Oncology and Hematology

Renowned for the diagnosis, therapy, and research of lymphoma, the UNMC Department of Internal Medicine, Division of Oncology and Hematology has treated patients from all over the world. Patients come for consultation, initial diagnosis, second opinions, standard chemotherapy, participation in clinical research trials, stem cell transplants, and chimeric antigen receptor (CAR) T-cell therapies.

The Division of Oncology and Hematology is comprised of 35 faculty and physicians, with five specializing in lymphoma. The Division is interested in the diagnosis and treatment of all malignancies (cancers). Treatment may include chemotherapy, radiation therapy, immunotherapy, pathway targeted agents, or cellular therapies, including CAR T-cell therapy and stem cell transplantation.

The Division's physicians specialize in many types of cancers including lymphomas, leukemia, multiple myeloma, urologic, breast, lung, gastric and pancreatic, neuro-oncology, melanoma, hepatocellular, and other solid tumors.

Nebraska Medicine

Nebraska Medicine is the most esteemed academic health system in the region, consisting of 809 licensed beds at its two hospitals, more than 1,000 physicians, and 40 specialty and primary care clinics in Omaha and surrounding areas.

Nebraska Medicine and UNMC, its research and education partner, share the same mission: to lead the world in transforming lives to create a healthy future for all individuals and communities through premier educational programs, innovative research and extraordinary patient care.

Nebraska Medicine traces its roots back to 1869 with the founding of Omaha's first hospital. In the decades since, it has built an international reputation for breakthroughs in cancer care, organ transplantation and treatment of infectious diseases.

For five straight years, Nebraska Medicine has also been named to Becker's Hospital Review's list of 100 Great Hospitals in America.

Fred & Pamela Buffett Cancer Center

The UNMC Fred & Pamela Buffett Cancer Center is the region's only National Cancer Institute (NCI) designated cancer center and is a member of the National Comprehensive Cancer Network (NCCN). The NCCN is an alliance of the nation's 33 leading cancer centers that develop and institute standards of care for the treatment of cancer and perform outcomes research with the goal of ensuring the delivery of high-quality, cost-effective services to cancer patients nationwide.

The Fred & Pamela Buffett Cancer Center opened in 2017 and continues to lead the world in the battle against cancer. By harnessing the most advanced biomedical and technological tools available, we are increasingly identifying the drivers behind cancer and creating precise therapies that improve outcomes. Through the use of genomics and other new diagnostic tools, we are employing precision medicine to customize therapies and care for each cancer patient.

While all forms of cancer will be treated at the Fred & Pamela Buffett Cancer Center, because of their prevalence in society and the center's potential to have a significant impact in their treatment, the following focus areas have been selected: breast cancer and other women's cancers, head and neck cancers, leukemia and lymphoma, lung cancer, pancreatic and gastrointestinal cancers, and prostate cancer.

The Fred & Pamela Buffett Cancer Center, along with the C.L. Werner Cancer Hospital, aim to provide the side-by-side, rapid development of therapeutics and delivery to patients with cancer.



Conference Information

Conference App


To make the most of your conference experience, we encourage you to use your mobile phone or tablet to access the conference app, or use your laptop or desktop to access the conference portal. See instructions on page 2.

Mobile Device Use

Please remember to turn your mobile device to silent or vibrate mode when attending educational sessions.



Internet Access

 Complimentary WiFi is available in the conference area.
Network: [UNMCPPLC24](#)
FREE Password: [PPLC2024](#)

Complimentary in-room WiFi is available for guests of the Hyatt Regency Maui Resort and Spa.

Photography and Audio Visual Recording Policy

Please be aware that during the 2024 Pan Pacific Lymphoma Conference, attendees, vendors, guests, and exhibitors may be photographed or videotaped by UNMC and/or third parties authorized by UNMC. Some of these photographs or videos may be displayed by UNMC in future publications or other materials. By virtue of your attendance, you agree to allow UNMC to use photographs of you in these promotional materials.

Photographs taken for the purpose of sharing on social media are allowed, but should avoid showing presentation slides. Please use [#PPLC24](#) and [#IamUNMC](#) when posting on social media.

Children and Other Accompanying Guests

PPLC 2024 does not offer childcare services. Due to limited seating capacity and the highly technical nature of the conference, children are not allowed to attend the educational sessions. For their safety, children must be accompanied by an adult at all times.

Registered accompanying guests may join conference registrants at the following food functions:

- Continental breakfasts (7–9 a.m.) and breaks as part of the main conference from Monday through Friday
- Welcome Reception on Tuesday evening
- Conference Dinner on Thursday evening



Name badges will be required.

Session and Room Assignments

- 1 GRAND PROMENADE**
Conference Registration
Continental Breakfast
Refreshment Break
Exhibits
- 2 MONARCHY BALLROOM 1-4**
General Sessions
- 3 MONARCHY BALLROOM 5-7**
Bierman's Brain Trivia – ERAS Edition
Ask the Experts
- 4 NAPILI GARDENS/LAWN**
Welcome Reception
- 5 HALONA KAI AND POOL DECK**
Conference Dinner

Registration Desk Hours

Sunday, July 14	2–4 p.m.
Monday, July 15	7 a.m. – 12:30 p.m.
Tuesday, July 16	7 a.m. – 12:30 p.m.
Wednesday, July 17	7 a.m. – 12:30 p.m.
Thursday, July 18	7 a.m. – 12:00 p.m.
Friday, July 19	7 a.m. – 12:00 p.m.

Exhibit Hours

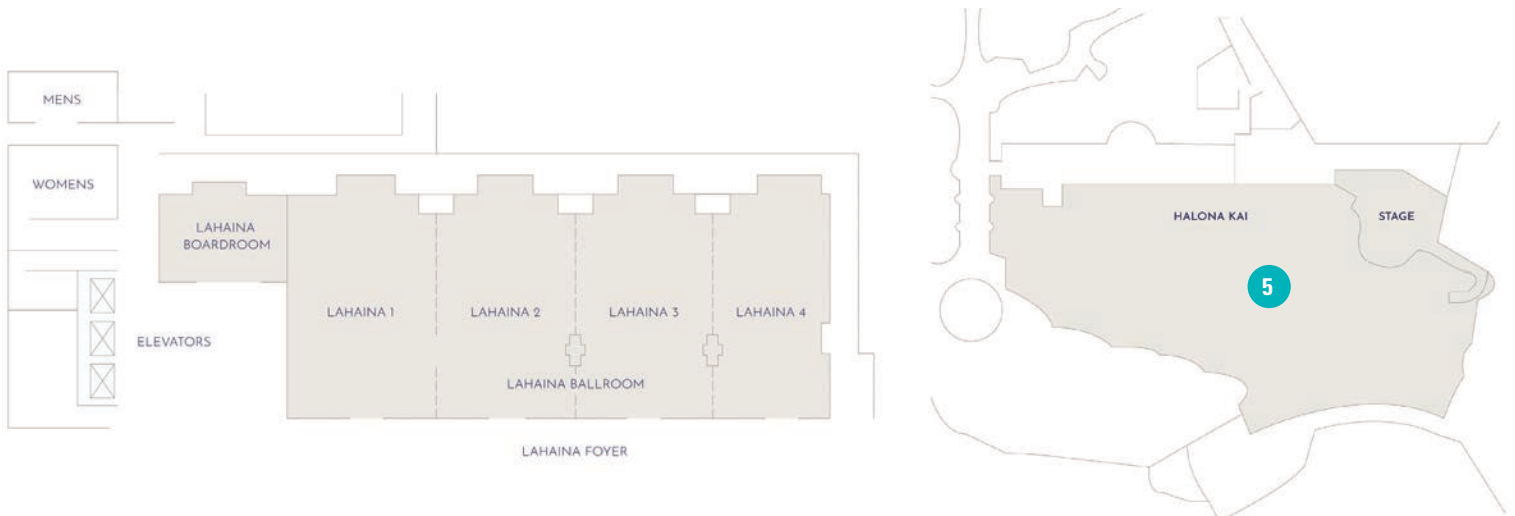
Monday, July 15	7 a.m. – 12:15 p.m.
Tuesday, July 16	7 a.m. – 1 p.m. / 3–5 p.m.
Wednesday, July 17	7 a.m. – 1 p.m. / 3–5 p.m.
Thursday, July 18	7 a.m. – 1 p.m. / 3–5 p.m.
Friday, July 19	7–10 a.m.

Venue Floor Plan

Promenade Level



Lobby Level



Conference Agenda

 All times are displayed in Hawaii Standard Time (HST)

Monday, July 15

INTERPROFESSIONAL EDUCATION DAY

7:00 a.m.	Conference Registration / Continental Breakfast	1
7:50 a.m.	Aloha / Conch Ceremony Matthew A. Lunning, DO, FACP	2
The Day-to-Day Treating of Lymphoma Patients MODERATOR: Ashley L. Schmitt, PA-C		2
8:00 a.m.	Updates on Utilization of Biosimilars and Generics in Lymphomas Jared E. Matya, PharmD, BCOP	
8:20 a.m.	Adjudicating Adverse Events in Lymphoma Patients Ashley L. Schmitt, PA-C	
8:40 a.m.	What the Social Worker Wished You Knew Angela Mortensen, LICSW	
9:00 a.m.	Optimal Team Utilization Round Table Christine Cox, RN Ann S. LaCasce, MD, MMSc Matthew A. Lunning, DO, FACP Caitlin A. Murphy, DNP, APRN, FNP-BC, AOCNP Ashley L. Schmitt, PA-C	
9:30 a.m.	Refreshment Break	1
How to Increase Your PI'ness MODERATOR: Katie Penas, MHA, CNMT, RT(N)		2
9:40 a.m.	How to Increase Your Odds in Obtaining an Investigator-Initiated Trial John P. Leonard, MD	
10:00 a.m.	How to Work with the Food and Drug Administration in Clinical Trials Grzegorz S. Nowakowski, MD	
10:20 a.m.	How to Recognize Clinical Trial Red Flags Katie Penas, MHA, CNMT, RT(N)	
10:40 a.m.	How to Increase Diversity in Clinical Trials Ruemu E. Birhiray, MD	

Practitioner Wellness		2
MODERATOR: Tara M. Graff, DO, MS		
11:00 a.m.	Taboo Topics in Healthcare David S. Kroll, MD	
11:20 a.m.	Achieving Career Balance: XX Perspective Sarah C. Rutherford, MD	
11:40 a.m.	Achieving Career Balance: XY Perspective Craig H. Moskowitz, MD	
12:00 p.m.	Adjourn	

Tuesday, July 16

7:00 a.m.	Conference Registration / Continental Breakfast	1
7:30 a.m.	Welcome James O. Armitage, MD and Julie M. Vose, MD, MBA	2
Future Assessment of Lymphoma MODERATOR: Kyle Skiermont, PharmD and Joann B. Sweasy, PhD		2
7:40 a.m.	What is the Governor in Lymphomagenesis Michael R. Green, PhD	
8:00 a.m.	Artificial Intelligence in Assessing Lymphoma Swaminathan P. Iyer, MD	
8:20 a.m.	Future of Liquid Biopsies in Lymphoma Davide Rossi, MD, PhD	
8:40 a.m.	Impact of World Health Organization or International Consensus Classification System in Lymphoma Elaine S. Jaffe, MD	
9:00 a.m.	Working Together to Improve Imaging Assessment in Lymphoma Sally F. Barrington, MBBS, MSc, FRCP, FRCR, MD Judith Trotman, MBChB, FRACP, FRCPA	
9:30 a.m.	Incorporating Liquid Biopsies into Clinical Practice in Aggressive Lymphomas Andrew D. Zelenetz, MD, PhD	
9:50 a.m.	Break	1



Rock the Vote: Choosing Initial Therapy Wisely

MODERATORS: Bruce D. Cheson, MD, FACP, FAAAS, FASCO and Matthew A. Lunning DO, FACP

2

- 10:05 a.m.** Advanced Stage Classical Hodgkin Lymphoma
- ABVD-Based – Stephen M. Ansell, MD, PhD
 - BV-Based – Ranjana H. Advani, MD
 - Checkpoint Inhibitors Based – Alex F. Herrera, MD

- 10:35 a.m.** Advanced Stage Large B-Cell Lymphoma
- R-CHOP – Jonathan W. Friedberg, MD, MMSc
 - Pola-R-CHOP – Sonali M. Smith, MD
 - Dose-Adjusted EPOCH-R – Mark Roschewski, MD

- 11:05 a.m.** Advanced Stage Mantle Cell Lymphoma
- Bruton’s Tyrosine Kinase Inhibitors + Chemo-Based – Michael Wang, MD
 - Alternating Regimen – Martin H. Dreyling, MD, PhD
 - BK/R-CHOP + Transplant – Kami J. Maddocks, MD

Lymphoplasmacytic Lymphoma and Plasma Cell Dyscrasias

MODERATOR: Sarah L. Creamer, MD

2

- 11:35 a.m.** When It’s Not Just MGUS
Natalie S. Callander, MD

- 11:55 a.m.** Induction Strategies in Lymphoplasmacytic Lymphoma (LPL)
R. Gregory Bociiek, MD, MSc, FRCP(C)

- 12:15 p.m.** B-Cell Maturation Antigen (BCMA) Directed Therapies in Multiple Myeloma
Krina K. Patel, MD, MSc

- 12:35 p.m.** Non-BCMA Directed Cellular Therapies in Plasma Cell Dyscrasias
Yi Lin, MD, PhD

- 12:55 p.m.** Adjourn

Bierman’s Brain Trivia—ERAS Edition

(ERAS ATTIRE REQUIRED)

MODERATOR: Matthew A. Lunning DO, FACP

3

- 3:00 p.m.** Pineapple Cup Semifinals #1
- | | |
|---|-------------------------------------|
| Team 1: FEARLESS | Team 2: SPEAK NOW |
| • James O. Armitage, MD | • Steven M. Horwitz, MD |
| • Bruce D. Cheson, MD, FACP, FAAAS, FASCO | • Kerry J. Savage, MD, MSc, FRCP(C) |
| • Elaine S. Jaffe, MD | • Jason R. Westin, MD, MS, FACP |

Pineapple Cup Semifinals #2

Team 1: FOLKLORE

- Ranjana H. Advani, MD
- Jonathan W. Friedberg, MD, MMSc
- Andrew D. Zelenetz, MD, PhD

Team 2: REPUTATION

- Alex F. Herrera, MD
- Caron A. Jacobson, MD, MMSc
- Sarah C. Rutherford, MD

Ask the Experts

3

- 4:00 p.m.** Indolent Non-Hodgkin Lymphoma (iNHL)
MODERATOR: R. Gregory Bociiek, MD, MSc, FRCP(C)
Jonathan W. Friedberg, MD, MMSc
Caron A. Jacobson, MD, MMSc
Sarah C. Rutherford, MD

- 5:00 p.m.** Adjourn

- 7:00 p.m.** Welcome Reception

Registered guests welcome.
Name badge required.

4

Wednesday, July 17

- 7:00 a.m.** Continental Breakfast

1

Just in “Case” You Asked Why I Did That

MODERATOR: Philip J. Bierman, MD

2

- 7:00 a.m.** Early Relapsing Follicular Lymphoma
- Auto Transplant – Craig S. Sauter, MD
 - Bispecific Antibody – Gilles Salles, MD, PhD
 - CAR-T – Sarah C. Rutherford, MD

- 7:45 a.m.** Early Relapsing Large B-Cell Lymphoma
- Chemotherapy + Auto Transplant – R. Gregory Bociiek, MD, MSc, FRCP(C)
 - CAR-T – Jason R. Westin, MD, MS, FACP
 - Neither – Nancy L. Bartlett, MD

- 8:30 a.m.** BTK Refractory Mantle Cell Lymphoma
- Non-CAR-T – Tycel Jovelle Phillips, MD
 - CAR-T – Marc S. Hoffmann, MD

- 9:00 a.m.** Relapsed Primary CNS Lymphoma
- BTK-Based – Christopher R. D’Angelo, MD
 - Chemo-Based – Kerry J. Savage, MD, MSc, FRCP(C)

Conference Agenda

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The Masked Lymphoma Doctor 2	
MODERATOR: Matthew A. Lunning, DO, FACP	
9:30 a.m.	A Case of Peripheral T-Cell Lymphoma Sonali M. Smith, MD Julie M. Vose, MD, MBA Kerry J. Savage, MD, MSc, FRCPC
10:00 a.m.	A Case of Mantle Cell Lymphoma Brad S. Kahl, MD John P. Leonard, MD Marc S. Hoffmann, MD
10:30 a.m.	Refreshment Break 1
MODERATORS: Matthew A. Lunning, DO, FACP and Julie M. Vose, MD, MBA 2	
10:50 a.m.	James O. Armitage Lymphoma Clinical Investigator Award Caron A. Jacobson, MD, MMSc
11:20 a.m.	Oliver Press Memorial Lecture Elaine S. Jaffe, MD
11:50 a.m.	Bierman's Brain Trivia Pineapple Cup Finals
12:30 p.m.	Adjourn
Ask the Experts 3	
3:00 p.m.	DLBCL MODERATOR: James O. Armitage, MD Nancy L. Bartlett, MD Jason R. Westin, MD, MS, FACP Andrew D. Zelenetz, MD, PhD
4:00 p.m.	cHL MODERATOR: Julie M. Vose, MD, MBA Ranjana H. Advani, MD Stephen M. Ansell, MD, PhD Craig H. Moskowitz, MD
5:00 p.m.	Adjourn

Thursday, July 18

7:00 a.m.	Continental Breakfast 1
Royal Lymphoma Rumble 2	
MODERATOR: Nitin Jain, MD	
7:30 a.m.	BV-CHP in CD30 Expressing Non-Anaplastic Large Cell Lymphoma - T-Cell Lymphoma For: Kerry J. Savage, MD, MSc, FRCPC Against: Swaminathan P. Iyer, MD

8:00 a.m.	Autologous Transplantation for Mantle Cell Lymphoma For: Andrew D. Zelenetz, MD, PhD Against: Martin H. Dreyling, MD, PhD
8:30 a.m.	BTK Monotherapy for First Line Chronic Lymphocytic Leukemia/Small Lymphocytic Lymphoma For: Danielle M. Brander, MD Against: Matthew S. Davids, MD, MMSc
9:00 a.m.	Optimal CAR-T Construct in 2026 Autologous: Jeremy S. Abramson, MD, MMSc Allogeneic: Frederick L. Locke, MD
9:30 a.m.	Refreshment Break 1
Who Knows 2	
MODERATOR: Vijaya Raj Bhatt, MBBS, MS	
9:50 a.m.	Is There a Right Management of Bispecific Toxicities? Community Perspective: Tara M. Graff, DO, MS Academic Perspective: Sarah C. Rutherford, MD
10:20 a.m.	Is There a Right Management of BTK and BCL-2 Refractory CLL/SLL? Susan M. O'Brien, MD
10:40 a.m.	Is There a Right Management of Extranodal Marginal Zone Lymphoma? Emanuele Zucca, MD
11:00 a.m.	Is There a Right Management of CPI & BV Refractory cHL? John Kuruvilla, MD, FRCP
11:20 a.m.	Is There a Right Management of Primary CNS Lymphoma? Neurologist Perspective: Christian Grommes, MD Oncologist Perspective: Christopher R. D'Angelo, MD
12:00 p.m.	Adjourn
Ask the Experts 3	
3:00 p.m.	PTCL MODERATOR: Matthew A. Lunning, DO, FACP Charles A. Enke, MD, FASTRO, FACR Steven M. Horwitz, MD Swaminathan P. Iyer, MD
4:00 p.m.	Waldenström Macroglobulinemia/Multiple Myeloma Moderator: Christopher R. D'Angelo, MD R. Gregory Bociek, MD, MSc, FRCPC Yi Lin, MD, PhD Krina K. Patel, MD, MSc
5:00 p.m.	Adjourn
7:00 p.m.	Conference Dinner 5

Registered guests welcome.
Name badge required.

Friday, July 19

7:00 a.m.	Continental Breakfast	1
Management of Orphan Diseases: When and How		
MODERATOR: Russell J. McCulloh, MD		
7:30 a.m.	When and How To Treat Gamma-Delta Hepatosplenic T-Cell Lymphoma Steven M. Horwitz, MD	2
7:50 a.m.	When and How To Treat with Total Skin Electron Beam Therapy (TSEBT) in Mycosis Fungoides Charles A. Enke, MD, FASTRO, FACR	
8:10 a.m.	When and How To Treat Autoimmune Complications of CLL/SLL Danielle M. Brander, MD	
8:30 a.m.	When and How To Treat Richter's Disease Matthew S. Davids, MD, MMSc	
8:50 a.m.	When and How To Treat Burkitt Lymphoma Mark Roschewski, MD	
9:10 a.m.	When and How To Treat Checkpoint-Related Adverse Events Alex F. Herrera, MD	
9:30 a.m.	Break	1
The Class of the Future is Here		
MODERATOR: Matthew A. Lunning, DO, FACP		
9:50 a.m.	Future Therapies: Cellular Therapies Nirav N. Shah, MD, MSHP	
10:20 a.m.	Future Therapies: Degradars & Celmods Marc S. Hoffmann, MD	
10:50 a.m.	Future Therapies: Bispecifics or X-Specific Antibodies Michael Dickinson, MD, PhD	
11:20 a.m.	Antibody Drug Conjugates Christopher R. D'Angelo, MD	
11:50 a.m.	Aloha/Conch Ceremony Matthew A. Lunning, DO, FACP	
12:00 p.m.	Adjourn	

Notes

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Conference Faculty

Jeremy S. Abramson, MD, MMSc

Director, Jon and Jo Ann Hagler Center for Lymphoma, Massachusetts General Hospital; Associate Professor of Medicine, Harvard Medical School, Boston, MA

Ranjana H. Advani, MD

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Stephen M. Ansell, MD, PhD

Dorothy W. and Grant L. Sundquist Professor in Hematologic Malignancies Research; Chair, Division of Hematology, Mayo Clinic, Rochester, MN

James O. Armitage, MD★

Joe Shapiro Professor of Medicine, Division of Oncology and Hematology, Department of Internal Medicine, University of Nebraska Medical Center, Omaha, NE

Sally F. Barrington, MBBS, MSc, FRCP, FRCR, MD

Professor of PET Imaging and NIHR Research Professor, School of Biomedical Engineering and Imaging Sciences, King's College London, St. Thomas' Hospital, London, United Kingdom

Nancy L. Bartlett, MD

Professor of Medicine, Koman Chair in Medical Oncology, Washington University School of Medicine, Siteman Cancer Center, St. Louis, MO

Vijaya Raj Bhatt, MBBS, MS

Professor, Division of Oncology and Hematology; Medical Director, Leukemia Program Section Leader, Malignant Hematology, University of Nebraska Medical Center, Omaha, NE

Philip J. Bierman, MD

Professor Emeritus, Division of Oncology and Hematology, Department of Internal Medicine, University of Nebraska Medical Center, Omaha, NE

Ruemu E. Birhiray, MD

Clinical Professor of Medicine, Marian University, College of Osteopathic Medicine; Oncology Specialist, Hematology Oncology of Indiana, A Division of American Oncology Network, PA.; President/CEO, Indy Hematology Education, Inc., Indianapolis, IN

R. Gregory Bociek, MD, MSc, FRCP(C)

Professor, Division of Oncology and Hematology, Department of Internal Medicine, University of Nebraska Medical Center, Omaha, NE

Danielle M. Brander, MD

Assistant Professor of Medicine; Director, CLL & Lymphoma Research Program, Member of the Duke Cancer Institute, Duke Health Integrated Practice, Durham, NC

Natalie S. Callander, MD

Professor of Medicine; Director, Myeloma Clinical Program, University of Wisconsin Carbone Cancer Center, Madison, WI

Bruce D. Cheson, MD, FACP, FAAAS, FASCO

Scientific Advisor to the Lymphoma Research Foundation; Center for Cancer and Blood Disorders, Bethesda, MD

Christine Cox, RN

Dana-Farber/Brigham & Women's Cancer Institute, Boston, MA

Sarah L. Creamer, MD

Hematologist/Medical Oncologist, Nebraska Cancer Specialists, Grand Island, NE

Christopher R. D'Angelo, MD

Assistant Professor, Division of Oncology and Hematology, Department of Internal Medicine, University of Nebraska Medical Center, Omaha, NE

Matthew S. Davids, MD, MMSc

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Education Information

Target Audience

The 2024 Pan Pacific Lymphoma Conference will benefit members of the multidisciplinary lymphoma clinical team including oncologists, hematologists, pathologists, clinical scientists, advanced practice providers, nurses, and pharmacists.

Conference Objectives

At the conclusion of the conference, the participant should be better able to:

1. Provide an update on biosimilar and generic medications utilized in the management of lymphoma.
2. Discuss the recognition of adverse events in new and emerging therapies in lymphoma that may result in dose reductions or dose holds in the management of lymphoma.
3. Identify the best utilization of rehabilitation facilities in the care of patients with lymphoma.
4. Discuss the optimal makeup and strategies for optimal clinic efficiency in a lymphoma-specific clinic.
5. Discuss how to optimally conduct and operationalize clinical trials for all potential subjects.
6. Identify factors related to anxiety, depression, and substance abuse amongst providers and the impact on their well-being.
7. Describe the key alterations that lead to lymphoma, how artificial intelligence may impact the pathological pathway to diagnosis of lymphoma, and how liquid tumor biopsy may supplement the response to therapy analysis.
8. Discuss how the evolving imaging technologies may be integrated into response assessment in the management of lymphoma.
9. Assess the impact of past and future therapies for the initial treatment of classical Hodgkin lymphoma (cHL), large B-cell lymphoma (LBCL), and mantle cell lymphoma (MCL).
10. Evaluate the diverse landscape of monoclonal gammopathies of unknown significance, treatment for lymphoplasmacytic lymphoma (LPL), and novel treatment for plasma cell dyscrasias (PCD).
11. Analyze clinical trial results impacting the management of relapsed and/or refractory follicular lymphoma (FL), peripheral T-cell lymphoma (PTCL), LBCL, MCL, and primary central nervous system lymphoma (PCNSL).
12. Differentiate the treatment options for CD30+ PTCL.
13. Discuss the changing role of consolidative autologous transplantation in MCL.
14. Discuss the optimal initial and relapsed/refractory sequencing for chronic lymphocytic leukemia (CLL)/small lymphocytic lymphoma (SLL).
15. Discuss optimal treatment and management of marginal zone lymphoma (MZL), CLL/SLL, PTCL, and MCL complications.
16. Explain the roles of the interprofessional health care team in the diagnosis, treatment, management, and support of lymphoma and hematologic malignancies.

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CONTRACTED RESEARCH: Pharmacyclics LLC (an AbbVie company)
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GRANT/RESEARCH SUPPORT: Abbvie, Allogene, Arcellx, Bristol Myers Squibb, Celgene, Collectis, Janssen, Nektar, Poseida, Precision Bio, Takeda

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CONTRACTED RESEARCH: Eli Lilly and Company, Miltenyi Biomedicine

GRANT/RESEARCH SUPPORT: Eli Lilly and Company, Miltenyi Biomedicine

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HONORARIUM RECIPIENT: AbbVie, Acerta Pharma, AstraZeneca, BeiGene, Inc., BioInvent, Bristol Myers Squibb, CAHON, Catamount Medical Education, Dava Oncology, Genmab, Janssen, Kite Pharma, MJH Life Sciences, Merck, MSC National Research Institute of Oncology, NIH, Nurix, Pharmacy

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OTHER FINANCIAL OR MATERIAL SUPPORT: AstraZeneca, Janssen, Merck, Roche

Disclosure Information

THE FOLLOWING FACULTY MEMBERS HAVE DISCLOSED OFF-LABEL, EXPERIMENTAL OR INVESTIGATIONAL USE OF DRUGS:

R. Gregory Bociek, MD, MSc, FRCP(C)

Off-label use discussion regarding data from ongoing clinical trials

Natalie S. Callander, MD

Off-label use discussed for smoldering myeloma: Lenalidomide, Daratumumab

Matthew S. Davids, MD, MMSc

Off-label use discussed: Ibrutinib, Acalabrutinib, Zanubrutinib, Obinutuzumab, Venetoclax, Pirtobrutinib

Michael Dickinson, MD, PhD

Off-label use for Lymphoma outside DLBCL or in combination - Glofitamab

Off-label use for Lymphoma outside of DLBCL / FL or in combination - Epcoritamab

Martin H. Dreyling, MD, PhD

Off-label use for MCL (first line) - Ibrutinib

John P. Leonard, MD

Off-label use for investigational lymphoma therapy

Susan M. O'Brien, MD

Off-label use for refractory CLL - Epcoritamab

Tycel Jovelle Phillips, MD

Off-label use for treatment of MCL - Glofitamab, mosunetuzumab/polatuzumab, venetoclax

Nirav N. Shah MD, MSHP

Off-label use of Novel CAR-T cells

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¹Financial relationships noted on pages 16.

²Financial relationships noted on pages 18.

³Financial relationships noted on pages 19.



PRODUCT THEATER KAUAI BALLROOM

Tuesday, July 16, 2024

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**Long-Term Clinical Data From Multiple Phase
3 Trials for CALQUENCE in the Treatment of CLL/SLL**

TARA GRAFF, DO

Hematologist/Oncologist
Mission Blood and Cancer
Des Moines, Iowa

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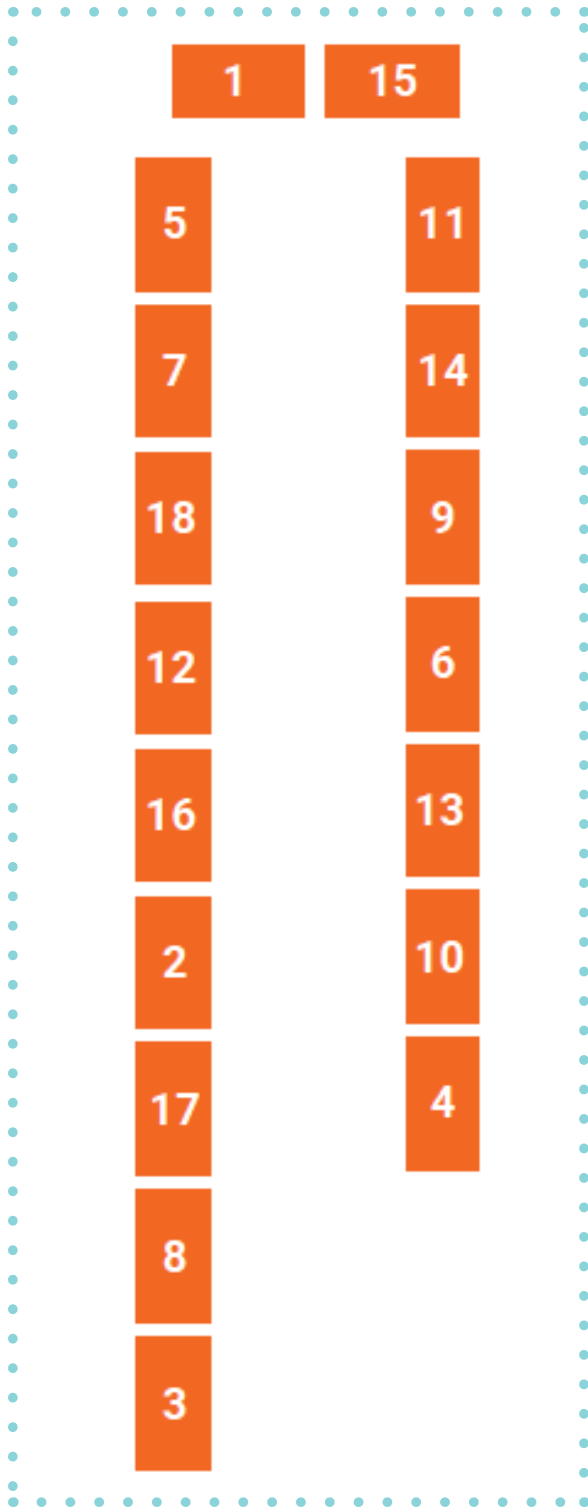
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THIS PRODUCT THEATER PRESENTS CLINICAL EFFICACY AND SAFETY DATA SUPPORTING THE USE OF A SELECTIVE BRUTON TYROSINE KINASE INHIBITOR FOR THE TREATMENT OF PREVIOUSLY UNTREATED AND R/R CLL

Exhibitors

Exhibit Hall

Grand Promenade



- 1 AbbVie, Inc.** is a global, research-driven biopharmaceutical company committed to developing innovative advanced therapies for some of the world's most complex and critical conditions. The company's mission is to use its expertise, dedicated people, and unique approach to innovation to improve treatments across four therapeutic areas: immunology, oncology, virology, and neuroscience. Follow @abbvie on X, Facebook, and LinkedIn, or visit www.abbvie.com.
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- 5 Bristol Myers Squibb Company** is a leading global biopharma company focused on discovering, developing, and delivering innovative medicines for patients with serious diseases in areas including oncology, hematology, immunology, cardiovascular, fibrosis, and neuroscience. Our employees work every day to transform patients' lives through science.
- 6 BTG Pharmaceuticals**, a SERB company, is dedicated to helping healthcare providers treat patients with critical conditions, focusing on emergency care and rare diseases.
- 7 Citius Pharmaceuticals** is a late-stage biopharmaceutical company focused on the development and commercialization of first-in-class critical care products, with a pipeline consisting of candidate therapeutics in Cutaneous T-Cell Lymphoma (CTCL), other oncology and adjunct cancer care, anti-infectives, stem cell therapy, and unique prescription products.

8

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11

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13

Kite Pharma Inc., a Gilead Company, is a global biopharmaceutical company based in Santa Monica, California, focused on cell therapy to treat and potentially cure cancer. As the global cell therapy leader, Kite has treated more patients with CAR T-cell therapy than any other company. Kite has the largest in-house cell therapy manufacturing network in the world, spanning process development, vector manufacturing, clinical trial production, and commercial product manufacturing. For more information on Kite, please visit www.kitepharma.com.

14

At **Merck & Co., Inc.**, known as MSD outside of the United States and Canada, we are unified around our purpose: We use the power of leading-edge science to save and improve lives around the world. For more than 130 years, we have brought hope to humanity through the development of important medicines and vaccines. We aspire to be the premier research-intensive biopharmaceutical company in the world – and today, we are at the forefront of research to deliver innovative health solutions that advance the prevention and treatment of diseases in people and animals. We foster a diverse and inclusive global workforce and operate responsibly every day to enable a safe, sustainable, and healthy future for all people and communities.

15

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16

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17

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18

Sobi, rare strength, is a specialized international biopharmaceutical company, transforming the lives of people with rare and debilitating diseases. Inspired by caring, powered by science, we are dedicated to ensuring every eligible person living with a rare and debilitating disease within our disease areas is given the opportunity to benefit from our approved medicines, delivering innovative solutions from our pipeline, and maintaining our commitment to patients, our employees, and society. For more information, please visit www.sobi.com.

EPCORITAMAB + R-DHAX/C ELICITS DEEP, DURABLE RESPONSES IN TRANSPLANT-ELIGIBLE PATIENTS WITH RELAPSED OR REFRACTORY DIFFUSE LARGE B-CELL LYMPHOMA, INCLUDING HIGH-RISK DISEASE

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BELLWAVE-011: A PHASE 3 STUDY OF NEMTABRUTINIB VERSUS IBRUTINIB OR ACALABRUTINIB IN PATIENTS WITH UNTREATED CHRONIC LYMPHOCYTIC LEUKEMIA/SMALL LYMPHOCYTIC LYMPHOMA

Razia Akhtar¹ on behalf of Tamar Tadmor, Tamar Tadmor²; Toby A. Eyre³; Ohad Benjamini⁴; Arvind Chaudhry⁵; Juan Shen¹; Siyang Leng¹; Mohammed Farooqui¹; David Lavie⁶

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PRELIMINARY VALIDATION OF A NOVEL PATIENT-REPORTED OUTCOMES (PRO) QUESTIONNAIRE FOR PATIENTS WITH CUTANEOUS T-CELL LYMPHOMA: THE CTCL PRO-18

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TRIAL IN PROGRESS: A PHASE 1-2 STUDY OF ZANUBRUTINIB AND TAFASITAMAB (TFA-ZANU) AS FIRST-LINE TREATMENT OF TRANSPLANT INELIGIBLE MANTLE CELL LYMPHOMA (MCL)

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PRIMARY ANALYSIS OF THE PHASE 2 ELM-2 STUDY: ODRONEXTAMAB IN PATIENTS WITH RELAPSED/REFRACTORY (R/R) DIFFUSE LARGE B-CELL LYMPHOMA (DLBCL)

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ASSESSMENT OF PHYSICIAN TREATMENT PREFERENCES FOR RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA IN THE THIRD-LINE SETTING

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RECENT PATTERNS OF CARE WITH BTK INHIBITORS AND DISTRIBUTION OF SOCIAL DETERMINANTS OF HEALTH AMONG PATIENTS WITH CLL/SLL IN THE US COMMUNITY SETTING

David J. Andorsky,¹ Thomas W. Wilson,² Kathleen M. Aguilar,² Chuck Wentworth,² Yunfei Wang,² Scott D. Goldfarb,³ Keri Yang,³ Erlene K. Seymour,³ Mark Balk,³ Gregory A. Maglinte,³ Ira Zackon²

¹The US Oncology Network/Rocky Mountain Cancer Centers, Boulder, CO, USA; ²Ontada, Boston, MA, USA; ³BeiGene USA, Inc, San Mateo, CA, USA

MOSUNETUZUMAB DEMONSTRATES CLINICALLY MEANINGFUL OUTCOMES IN HIGH-RISK PATIENTS WITH HEAVILY PRE-TREATED R/R FL AFTER ≥3 YEARS OF FOLLOW-UP: SUBGROUP ANALYSIS OF A PIVOTAL PHASE II STUDY

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ORBITAL GERMINAL CENTER DIFFUSE LARGE B-CELL LYMPHOMA

Taras Benzar

The Oncology Institute of Hope And Innovation (TOI), Culver City, California, USA

LENS-SPARING VMAT EBRT FOR THE TREATMENT OF CHOROICAL LYMPHOMA

C. Billena, W Vuong, A Bommireddy, E Murray, A Singh, S, Cherian
Taussig Cancer Center, Cleveland Clinic Foundation, Cleveland, OH, US

MULTICENTER, RANDOMIZED PHASE II STUDY OF EPCORITAMAB FOR PATIENTS WITH LARGE B-CELL LYMPHOMAS ACHIEVING A PARTIAL RESPONSE AFTER CD19-DIRECTED CAR T-CELL THERAPY

Allison M. Bock, MD¹, Sandeep Raj, MD², Sharmila Giri, MS³, Lori A. Leslie, MD⁴, Dilan A. Patel, MD⁵, Madiha Iqbal, MBBS, MD⁶, Natalie Grover, MD⁷, Boyu Hu, MD¹, Miguel-Angel Perales, MD², Grzegorz S. Nowakowski, MD⁸

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TRIAL IN PROGRESS: PHASE 3 STUDY EVALUATING THE EFFICACY AND SAFETY OF ODRONEXTAMAB VERSUS INVESTIGATOR'S CHOICE IN PREVIOUSLY UNTREATED PATIENTS WITH FOLLICULAR LYMPHOMA (OLYMPIA-1)

Elizabeth A. Brem,¹ Ruemu Birhiray,² Stefano Luminari,³ Tae Min Kim,⁴ Münci Yağci,⁵ Timothy Wassenaar,⁶ Ashish Risal,⁷ Manjusha Namuduri,⁷ Jingxian Cai,⁷ Bertha Fearon-Chalet,⁷ Min Zhu,⁷ Sushmita Mukherjee,⁷ Jurriaan Brouwer-Visser,⁷ Aafia Chaudhry,⁷ Hesham Mohamed,⁷ Srikanth Ambati,⁷ Umberto Vitolo⁸

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CIRCULATING TUMOR DNA ANALYSIS ASSOCIATES WITH PFS WITH ODRONEXTAMAB MONOTHERAPY IN R/R FL AND DLBCL: IDENTIFICATION OF MRD STATUS AND HIGH-RISK SUBGROUPS FROM THE PHASE 2 ELM-2 STUDY

Jurriaan Brouwer-Visser¹, David Tucker², Jon E. Arnason³, Stefano Luminari⁴, Tae Min Kim⁵, Won Seog Kim⁶, Laura Magnano Mayer⁷, Dylan Sun¹, Jingxian Cai¹, Kamil Cygan¹, Srikanth Ambati¹, Hesham Mohamed¹, Aafia Chaudhry¹, Vladimir Jankovic⁸

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LONG TERM SURVIVAL TRENDS IN OLDER PATIENTS WITH CLASSICAL HODGKIN LYMPHOMA: AN ANALYSIS OF THE TEXAS CANCER REGISTRY

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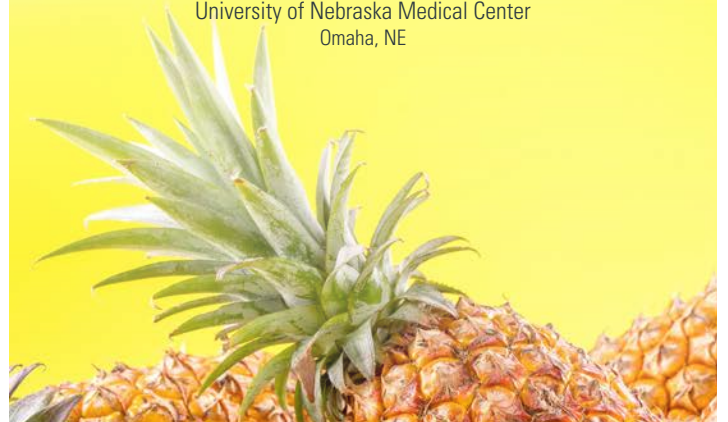
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CUTANEOUS MANTLE CELL LYMPHOMA: AN MD ANDERSON CANCER CENTER EXPERIENCE

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PRIMARY ANALYSIS OF THE PHASE 2 ELM-2 STUDY: ODRONEXTAMAB IN PATIENTS WITH RELAPSED/REFRACTORY FOLLICULAR LYMPHOMA

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EPCORITAMAB + GEMOX INDUCES DEEP, DURABLE RESPONSES IN PATIENTS WITH RELAPSED OR REFRACTORY DIFFUSE LARGE B-CELL LYMPHOMA: UPDATED RESULTS FROM THE EPCORE NHL-2 TRIAL

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EFFICACY AND SAFETY OF E7777 (IMPROVED PURITY DENILEUKIN DIFTITOX [ONTAK]) IN PATIENTS WITH RELAPSED OR REFRACTORY CUTANEOUS T-CELL LYMPHOMA: RESULTS FROM PIVOTAL STUDY 302

Francine Foss¹, MD, Youn H Kim², MD, Miles Prince³, MD, FRACP, FRCPA, Christiane Querfeld⁴, MD, PhD, Costas K Yannakou⁵ MBBS, FRACP, FRCPA, PhD, Chean Eng Ooi⁶, PhD, Dongyuan Xing⁶, PhD, Nicholas Sauter⁶, MD, Preeti Singh⁷, MD, Myron Czuczman⁷, MD, Madeleine Duvic⁸, MD

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MATCHING-ADJUSTED INDIRECT COMPARISONS OF EPCORITAMAB VS MOSUNETUZUMAB OR ODRONEXTAMAB IN RELAPSED/REFRACTORY FOLLICULAR LYMPHOMA AFTER ≥2 SYSTEMIC THERAPIES

Alexey Danilov,¹ Swetha Kambhampati,¹ Kim Linton,² Karen Cumings,³ Viktor Chirikov,³ Alex Mutebi,⁴ Savreet Bains Chawla,⁴ Fernando Rivas Navarro,⁵ Anthony Wang,⁶ Zhijie Ding,⁴ Abualbisher Alshreef,⁶ Elena Favaro,⁵ Daniela Hoehn,⁴ Anna Sureda⁷

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IMPROVING ADVANCE CARE PLANNING OUTCOMES

Jessica Davis, DNP

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TRANSCRIPTIONAL REGULATION OF CELL CYCLE PROGRESSION IN B CELL LYMPHOMA

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TRANSCEND FL: PHASE 2 STUDY PRIMARY ANALYSIS OF LISOCABTAGENE MARALEUCEL (LISO-CEL) AS SECOND-LINE (2L) THERAPY IN PATIENTS (PT) WITH HIGH-RISK R/R FL

F Morschhauser,¹ S Dahiya,² ML Palomba,³ AM Garcia-Sancho,⁴ JLR Ortega,⁵ J Kuruvilla,⁶ U Jäger,⁷ G Cartron,⁸ K Izutsu,⁹ M Dreyling,¹⁰ B Kahl,¹¹ H Ghesquieres,¹² K Ardeshtna,¹³ H Goto,¹⁴ AM Barbui,¹⁵ JS Abramson,¹⁶ P Borchmann,¹⁷ I Fleury,¹⁸ S Mielke,¹⁹ A Skarbnik,²⁰ S de Vos,²¹ Manali Kamdar,²² R Karmali,²³ A Viardot,²⁴ T Farazi,²⁵ M Vedal,²⁵ R Nishii,²⁵ A Avilion,²⁵ J Papuga,²⁵ LJ Nastoupil²⁶

¹Centre Hospitalier Universitaire de Lille, Lille, France; ²Stanford University School of Medicine, Stanford, CA, USA; ³Memorial Sloan Kettering Cancer Center, New York, NY, USA; ⁴Hospital Universitario de Salamanca, IBSAL, CIBERONC, Centro de Investigación del Cáncer-IBMCC (USAL-CSIC), Salamanca, Spain; ⁵Hospital Universitario Virgen del Rocío, IBIS/CSIC, Universidad de Sevilla, Seville, Spain; ⁶Princess Margaret Cancer Centre, Toronto, ON, Canada; ⁷Medical University of Vienna, Vienna, Austria; ⁸Montpellier University Hospital Center, UMR CNRS 5535, Montpellier, France; ⁹National Cancer Center Hospital, Tokyo, Japan; ¹⁰Medizinische Klinik III, Klinikum der Universität, LMU München, München, Germany; ¹¹Washington University School of Medicine in St. Louis, St. Louis, MO, USA; ¹²Hôpital Lyon Sud, Lyon, France; ¹³University College London Hospitals NHS Foundation Trust-University College Hospital, London, UK; ¹⁴Hokkaido University Hospital, Sapporo, Japan; ¹⁵Azienda Socio Sanitaria Territoriale Papa Giovanni XXIII, Bergamo, Italy; ¹⁶Massachusetts General Hospital Cancer Center, Harvard Medical School, Boston, MA, USA; ¹⁷Universität zu Köln, Köln, Germany; ¹⁸Hôpital Maisonneuve – Rosemont, Montreal, QC, Canada; ¹⁹Karolinska Institutet and University Hospital, Karolinska Comprehensive Cancer Center, Stockholm, Sweden; ²⁰Novant Health Cancer Institute, Charlotte, NC, USA; ²¹UCLA Santa Monica Medical Centre, Santa Monica, CA, USA; ²²University of Colorado Cancer Center, Aurora, CO, USA; ²³Northwestern University Feinberg School of Medicine, Robert H. Lurie Comprehensive Cancer Center, Chicago, IL, USA; ²⁴University Hospital, Ulm, Germany; ²⁵Bristol Myers Squibb, Princeton, NJ, USA; ²⁶The University of Texas MD Anderson Cancer Center, Houston, TX, USA

DREAMM-7 UPDATE: SUBGROUP ANALYSES FROM A PHASE 3 TRIAL OF BELANTAMAB MAFODOTIN (BELAMAF) + BORTEZOMIB AND DEXAMETHASONE (BVD) VS DARATUMUMAB, BORTEZOMIB, AND DEXAMETHASONE (DVD) IN RELAPSED/REFRACTORY MULTIPLE MYELOMA (RRMM)

María Victoria Mateos Manteca¹; Pawel Robak²; Marek Hus³; ChengCheng Fu⁴; Vera Zherebtsova⁵; Christopher Ward⁶; P. Joy Ho⁷; Ana Carolina de Almeida Ribas⁸; Roman Hajek⁹; Kihyun Kim¹⁰; Meletios Dimopoulos¹¹; Claudio Cerchione¹²; Nick Pirooz¹³; Astrid McKeown¹⁴; Benga Kazeem¹⁴; Hena Baig¹⁵; [Lydia Eccersley](#)¹⁶; Sumita Roy-Ghanta¹³; Joanna Opalinska¹³; Vania Hungria¹⁷

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STAGING FDG-AVIDITY IN EXTRANODAL MARGINAL ZONE LYMPHOMA (EMZL) BY DISEASE LOCATION

[E. Edelman Saul](#)¹, J.P. Alderuccio², M. Stanchina², M. Polar², R. Hennemann Sassi², W. Zhao², C.H. Moskowitz², I. Reis², R. Kuker², I. Lossos²

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HYPERAMMONEMIC ENCEPHALOPATHY IN MULTIPLE MYELOMA: SYSTEMATIC REVIEW AND CASE SERIES

[Joseph Anthony El Ghoubaire](#) MD¹, [Aseel Alsouqi](#) MD¹, [Elia Abou Chawareb](#) MD², [Rita Farah MPH](#), PharmD, PhD³, [Rafic J. Farah MD](#)¹

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NASOPHARYNGEAL MALT LYMPHOMA: A CASE REPORT

[Eiman Elmileik](#) and [Carlos Galvez](#)

University of Illinois at Chicago, Chicago, IL

EPCORITAMAB WITH RITUXIMAB + LENALIDOMIDE (R2) IN PREVIOUSLY UNTREATED (1L) FOLLICULAR LYMPHOMA (FL) AND EPCORITAMAB MAINTENANCE THERAPY IN FL: NOVEL RESULTS FROM EPCORE NHL-2 ARMS 6 AND 7

[L. Falchi](#)¹, [J.S.P. Vermaat](#)², [G. Musuraca](#)³, [M. Nijland](#)⁴, [J.H. Christensen](#)⁵, [F. Offner](#)⁶, [L.A. Leslie](#)⁷, [J.D. Brody](#)⁸, [D. Hoehn](#)⁹, [J. Marek](#)⁹, [L. Wang](#)⁹, [J. Mei](#)¹⁰, [D. Belada](#)¹¹, [P. Abrisqueta](#)¹²

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USE OF BISPECIFIC T-CELL ENGAGING ANTIBODIES (BSABS) IN COMMUNITY PRACTICES: MULTIDISCIPLINARY PERSPECTIVES ON DEVELOPING LOGISTICS AND WORKFLOW FOR CYTOKINE RELEASE SYNDROME (CRS) MANAGEMENT

William Donnellan¹, Shih-Wen Lin², Jonathan Abbas¹, Jesus G. Berdeja¹, Lourenia Cassoli², Jason C. Chandler³, [Brannon Flores](#)², Sara Hall⁴, Arliene Mangalindan², Anthony Masaquel², Sharifa Patterson¹, Eileen Peng⁵, Ashley Todd¹, Chelsea Traugher¹, Lisa Raff⁶

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HEPATOSPLENIC T-CELL LYMPHOMA, $\alpha\beta$ TYPE, PRESENTING IN AN ELDERLY PATIENT STATUS POST INDOLENT B-CELL LYMPHOMA TREATMENT

[Rodolfo Garza-Morales](#)¹, Allison Rosenthal¹, Jiehao Zhou²

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THE ADVANCED PRACTICE PROVIDERS IDENTIFICATION OF GRADE 3 TOXICITIES ASSOCIATED WITH BRUTON KINASE INHIBITORS IN PATIENTS WITH RELAPSED/REFRACTORY MARGINAL ZONE LYMPHOMA

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TRIAL IN PROGRESS: PHASE 3 STUDY EVALUATING THE EFFICACY AND SAFETY OF ODRONEXTAMAB PLUS CHEMOTHERAPY VERSUS RITUXIMAB PLUS CHEMOTHERAPY IN PREVIOUSLY UNTREATED PATIENTS WITH FOLLICULAR LYMPHOMA (OLYMPIA-2)

[Colin Hardin](#)¹, [Carl Gray](#)², [Silvana Novelli](#)³, [Tuba Hacibekiroğlu](#)⁴, [Münci Yağcı](#)⁵, [Ruemu Birhiray](#)⁶, [Ashish Risal](#)⁷, [Manjusha Namuduri](#)⁷, [Jingxian Cai](#)⁷, [Melanie Ufkin](#)⁷, [Min Zhu](#)⁷, [Sushmita Mukherjee](#)⁷, [Jurriaan Brouwer-Visser](#)⁷, [Aafia Chaudhry](#)⁷, [Hesham Mohamed](#)⁷, [Srikanth Ambati](#)⁷, [Elżbieta Iskierka-Jażdżewska](#)⁸

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CNS TUBERCULOSIS IN A PATIENT WITH ANGIOIMMUNOBLASTIC T-CELL LYMPHOMA: A CASE STUDY

[Zoey Harris, MD](#)¹, [Connor Yost, MD](#)¹, [Yael Kusne, MD](#)¹, [Allison Rosenthal, DO](#)¹

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LONG-TERM SECONDARY CAUSE-SPECIFIC MORTALITY IN OLDER PATIENTS WITH CLASSICAL HODGKIN LYMPHOMA: AN ANALYSIS OF THE TEXAS CANCER REGISTRY

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DECIPHERING THE CLINICAL BENEFIT OF POLA-R-CHP VERSUS R-CHOP IN DIFFERENT GENETIC SUBTYPES BEYOND CELL OF ORIGIN IN THE POLARIX STUDY

Franck Morschhauser¹, Wilfred Leung², Vibha Raghavan³, Georg Lenz⁴, Fabrice Jardin⁵, Alex F. Herrera⁶, Laurie H. Sehn⁷, Jeff P. Sharman⁸, Christopher R. Flowers⁹, Jonathan W. Friedberg¹⁰, Marek Trnėný¹¹, Hervé Tilly⁵, Charles Herbaux¹², Samuel Tracy², Christopher R. Bolen², Will Harris², Jamie H. Hirata², Calvin Lee², Yanwen Jiang², Connie Lee Batlevi², Gilles Salles¹³

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THE IMPACT OF COMMUNICATION WITH HEALTHCARE PROVIDERS ON DAILY LIVING ADJUSTMENT IN PATIENTS WITH LYMPHOMA

Hyunju Hong, Ehwa Yun, Bobae Jeong, Ahyeong Jo, Yeongme Kim, Yunseon Choe, Kwangmi Lee

Division of Hematology, National Cancer Center, 323 Ilsan-ro, Goyang-si, Gyeonggi-do, 10408, Republic of Korea

A GLOBAL PHASE 2 TRIAL OF NANATINOSTAT IN COMBINATION WITH VALGANCICLOVIR IN PATIENTS WITH EBV- POSITIVE (EBV+) RELAPSED/REFRACTORY PERIPHERAL T CELL LYMPHOMAS (NAVAL-1)

Swaminathan Iyer, MD¹, Ranjit Nair, MD², Hung Chang, MD³, Irl Brian Greenwell, MD⁴, Hui-Hua Hsiao, MD, PhD⁵, Luigi Zinzani, MD⁶, Kiran Vangipuram, PharmD⁷, Donald Strickland, MD⁷, Eva González-Barca, MD⁸

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PATIENTS WITH R/R LARGE B-CELL LYMPHOMA (LBCL) TREATED WITH LISOCABTAGENE MARALEUCEL (LISO-CEL) NONCONFORMING PRODUCT (NCP) UNDER THE EXPANDED ACCESS PROTOCOL (EAP)

BG Till, ¹C. Jacobson, ²ML Palomba, ³JS Abramson, ⁴J Arnason, ⁵FL Locke, ⁶P Caimi, ⁷N Grover, ⁸R Karmali, ⁹HC Suh, ¹⁰SJ Schuster, ¹¹M Gharibo, ¹²C Balint, ¹²T Rakhmawati, ¹³V Jude, ¹²LJ Nastoupil¹⁴

¹Fred Hutchinson Cancer Center, Seattle, WA, USA; ²Dana-Farber Cancer Institute, Boston, MA, USA; ³Memorial Sloan Kettering Cancer Center, New York, NY, USA; ⁴Massachusetts General Hospital Cancer Center, Harvard Medical School, Boston, MA, USA; ⁵Beth Israel Deaconess Medical Center, Boston, MA, USA; ⁶H Lee Moffitt Cancer Center, Tampa, FL, USA; ⁷CCF Taussig Cancer Center, Cleveland, OH, USA; ⁸Lineberger Comprehensive Cancer Center, University of North Carolina School of Medicine, Chapel Hill, NC, USA; ⁹Northwestern University Feinberg School of Medicine, Robert H. Lurie Comprehensive Cancer Center, Chicago, IL, USA; ¹⁰John Theurer Cancer Center, Hackensack University Medical Center, Hackensack, NJ, USA; ¹¹Lymphoma Program, Abramson Cancer Center, Philadelphia, PA, USA; ¹²Bristol Myers Squibb, Princeton, NJ, USA; ¹³Syneos Health, Morrisville, NC, USA; ¹⁴The University of Texas MD Anderson Cancer Center, Houston, TX, USA

CCR4 MUTATIONS DYSREGULATE T-HELPER DIFFERENTIATION AND PROMOTE LYMPHOMAGENESIS

Dylan T Jochum¹, Suchita Vishwakarma², Alyssa Bouska¹, Waseem G Lone¹, Ab Rauf Shah³, Abdul Rouf Mir¹, Sunandini Sharma¹, Mehnaz Tabassum¹, Zaina Nasser³, Amarnath Natarajan⁴, and Javeed Iqbal¹

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THE MULTICOLOR FLOW CYTOMETRY FOR PRIMARY VITREORETINAL LYMPHOMA DIAGNOSIS: A SINGLE CENTER EXPERIENCE

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PATIENT-REPORTED OUTCOMES IN PATIENTS WITH RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA TREATED WITH EPCORITAMAB

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SINGLE-AGENT EPCORITAMAB LEADS TO DEEP RESPONSES IN PATIENTS (PTS) WITH RICHTER'S TRANSFORMATION (RT): PRIMARY RESULTS FROM THE EPCORE CLL-1 TRIAL

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RETREATMENT WITH NKX019 (CD19-TARGETED CAR NK CELL THERAPY) IS FEASIBLE, SAFE, AND EFFECTIVE WITH IN SUBJECTS WITH NHL

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REAL-WORLD TREATMENT PATTERNS AND PATIENT OUTCOMES IN RELAPSED/REFRACTORY HODGKIN LYMPHOMA IN THE US

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SAFETY AND EFFICACY OF BRETUXIMAB VEDOTIN WITH PROPHYLACTIC G-CSF IN CLASSICAL HODGKIN LYMPHOMA (SGN35-027, PART A)

Hun Ju Lee,¹ Ian Flinn,² Leland Metheny,³ Judah Friedman,⁴ Joseph Mace,⁵ Paul Gonzales,⁶ Ameet Patel,⁷ Mihir Raval,⁸ Asad Dean,⁹ Griffith Davis,¹⁰ Wenchuan Guo,¹⁰ Michelle Fanale,¹⁰ Chris Yasenachak¹¹

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COMPARISON OF CLONOSEQ AND MULTIPARAMETER FLOW CYTOMETRIC ANALYSIS OF MINIMAL RESIDUAL DISEASES IN MYELOMA

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CAR T ADMINISTERED IN THE OUTPATIENT SETTING FOR DLBCL: PATIENT CHARACTERISTICS, TREATMENT PATTERNS AND HEALTHCARE RESOURCE USE

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COOPERATIVE ROLE OF TET2 AND IDH2 MUTATIONS GENERATES T-CELL LYMPHOMA WITH TFH-CELL IMMUNOPHENOTYPE AND DIFFERENTIATION BLOCK IN TH1 PROGRAM

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REAL-WORLD BRUTON TYROSINE KINASE INHIBITOR TREATMENT PATTERNS AND OUTCOMES AMONG PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA OR SMALL LYMPHOCYTIC LYMPHOMA IN US COMMUNITY ONCOLOGY PRACTICES

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REAL-WORLD COST OF DISEASE PROGRESSION (PD) AFTER FRONTLINE (1L) R- CHOP IN DIFFUSE LARGE B-CELL LYMPHOMA (DLBCL): AN ANALYSIS OF A LARGE US CLAIMS DATABASE

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TRIAL IN PROGRESS: PHASE 3 STUDY EVALUATING EFFICACY AND SAFETY OF ODRONEXTAMAB PLUS CHOP VERSUS RITUXIMAB PLUS CHOP IN PREVIOUSLY UNTREATED PATIENTS WITH DIFFUSE LARGE B-CELL LYMPHOMA (DLBCL; OLYMPIA-3)

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PATIENT-REPORTED OUTCOMES (PROS) FROM THE DREAMM-7 RANDOMIZED PHASE 3 STUDY COMPARING BELANTAMAB MAFODOTIN, BORTEZOMIB, AND DEXAMETHASONE (BVD) VS DARATUMUMAB, BORTEZOMIB, AND DEXAMETHASONE (DVD) IN PATIENTS WITH RELAPSED/REFRACTORY MULTIPLE MYELOMA (RRMM)

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MOSUNETUZUMAB WITH POLATUZUMAB VEDOTIN: SUBGROUP ANALYSES IN PATIENTS (PTS) WITH PRIMARY REFRACTORY OR EARLY RELAPSED LARGE B-CELL LYMPHOMA (LBCL)

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TRANSDIFFERENTIATION WITH GENOTYPIC EVIDENCE OF A CLONAL RELATIONSHIP BETWEEN DIFFUSE LARGE B-CELL LYMPHOMA AND HISTIOCYTIC SARCOMA: A CASE REPORT

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LOGISTICAL CHALLENGES ASSOCIATED WITH CHIMERIC ANTIGEN RECEPTOR T-CELL THERAPY (CAR T) IN NON-HODGKIN LYMPHOMA (NHL): A SURVEY OF HEALTHCARE PROFESSIONALS

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EARLIER TIME POINT ASSESSMENT OF METHOTREXATE CONCENTRATIONS PREDICTIVE OF NEPHROTOXICITY IN PATIENTS RECEIVING HIGH DOSE METHOTREXATE FOR DIFFUSE LARGE B CELL LYMPHOMA

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COMPARATIVE ANALYSIS OF LUMBAR PUNCTURE UTILITY IN DIAGNOSIS OF PRIMARY VERSUS SECONDARY CNS LYMPHOMA: A RETROSPECTIVE STUDY

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LOTIS-5, AN ONGOING, PHASE 3, RANDOMIZED STUDY OF LONCASTUXIMAB TESIRINE WITH RITUXIMAB (LONCA-R) VERSUS IMMUNOCHEMOTHERAPY IN PATIENTS WITH R/R DLBCL

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A CRISPR-EDITED ALLOGENEIC ANTI-CD19 CAR-T CELL THERAPY WITH A PD-1 KNOCKOUT (CB-010) IN PATIENTS WITH RELAPSED/REFRACTORY B CELL NON-HODGKIN LYMPHOMA (R/R B-NHL): UPDATED PHASE 1 RESULTS FROM THE ANTLER TRIAL

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INCIDENCE OF CENTRAL NERVOUS SYSTEM RELAPSE IN PRIMARY MEDIASTINAL B-CELL LYMPHOMA: IMPLICATIONS FOR CENTRAL NERVOUS SYSTEM PROPHYLAXIS

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DOSE ADJUSTMENT OUTCOMES IN PATIENTS WITH WALDENSTRÖM MACROGLOBULINEMIA TREATED WITH IBRUTINIB

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HIGH INCIDENCE OF CYTOMEGALOVIRUS (CMV) REACTIVATION IN PATIENTS RECEIVING CAR T-CELL THERAPY

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PIRTOBRUTINIB IN RELAPSED/REFRACTORY CLL/SLL: RESULTS FROM BTKI NAÏVE COHORT IN THE PHASE 1/2 BRUIN STUDY

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ECONOMIC BURDEN OF ADVERSE EVENTS AMONG RELAPSED/REFRACTORY MANTLE CELL LYMPHOMA PATIENTS RECEIVING LISOCABTAGENE MARALEUCEL IN THE TRANSCEND MCL CLINICAL TRIAL

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STAGING PATTERNS IN BLOOD AND LYMPH NODES IN PATIENTS WITH LIMITED STAGE CUTANEOUS T-CELL LYMPHOMA (CTCL)

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NEMTABRUTINIB PLUS VENETOCLAX VERSUS VENETOCLAX PLUS RITUXIMAB IN TREATED PATIENTS WITH RELAPSED/REFRACTORY CHRONIC LYMPHOCYTIC LEUKEMIA/SMALL LYMPHOCYTIC LYMPHOMA: PHASE 3 BELLWAVE-010 STUDY

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TREATMENT OF ANTI-MYELIN-ASSOCIATED-GLYCOPROTEIN (MAG) ANTIBODY NEUROPATHY WITH ZANUBRUTINIB IN A PATIENT WITH WALDENSTROM MACROGLOBULINEMIA: A CLINICAL VIGNETTE

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AN ECONOMIC MODEL TO ESTIMATE COSTS OF SELECT ADVERSE EVENTS AMONG PATIENTS TREATED WITH CHIMERIC ANTIGEN RECEPTOR (CAR) T CELL THERAPIES FOR R/R MCL

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COFORMULATED FAVEZELIMAB AND PEMBROLIZUMAB VERSUS CHEMOTHERAPY IN PATIENTS WITH ANTI-PD-1 REFRACTORY CLASSICAL HODGKIN LYMPHOMA: A PHASE 3 KEYFORM-008 STUDY

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HEALTH-RELATED QUALITY OF LIFE (HRQOL) AND SYMPTOMS IN PATIENTS WITH RELAPSED OR REFRACTORY DIFFUSE LARGE B-CELL LYMPHOMA (R/R DLBCL) TREATED WITH ODRONEXTAMAB MONOTHERAPY IN THE PHASE 2 ELM-2 STUDY

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SAFETY AND TOLERABILITY OF E7777 (IMPROVED PURITY DENILEUKIN DIFTITOX [ONTAK]) IN PATIENTS WITH RELAPSED OR REFRACTORY CUTANEOUS T-CELL LYMPHOMA: RESULTS FROM PIVOTAL STUDY 302

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HEALTH AND ECONOMIC IMPACT OF CAR T VEIN-TO-VEIN TIME IN RELAPSED/REFRACTORY 2L LBCL PATIENTS: A US COST-EFFECTIVENESS ANALYSIS

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EFFICACY AND SAFETY OF ZILOVERTAMAB VEDOTIN IN PATIENTS WITH RELAPSED OR REFRACTORY DIFFUSE LARGE B-CELL LYMPHOMA: UPDATED RESULTS FROM THE PHASE 2 WAVELINE-004 STUDY

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MANAGEMENT OF RELAPSED CLASSICAL HODGKIN LYMPHOMA (CHL) IN PREGNANCY

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PIRTOBRUTINIB IN POST-CBTKI CLL/SLL: ~30 MONTHS FOLLOW-UP AND SUBGROUP ANALYSIS WITH/WITHOUT PRIOR BCL2I FROM THE PHASE 1/2 BRUIN STUDY

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QUANTITATIVE SYSTEMS PHARMACOLOGY MODEL PREDICTS COMBINATION ACTIVITY OF CD19-TARGETED LONCASTUXIMAB TESIRINE CO-DOSED WITH A CD20/CD3 T-CELL BISPECIFIC (EPCORITAMAB) IN DIFFUSE LARGE B-CELL LYMPHOMA

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REAL-WORLD USE OF TAFASITAMAB FOR RELAPSED OR REFRACTORY DIFFUSE LARGE B- CELL LYMPHOMA IN THE UNITED STATES BY PRIMARY REFRACTORY STATUS

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PHASE 3 TRIAL OF SUBCUTANEOUS EPCORITAMAB + R-CHOP VERSUS R-CHOP IN PATIENTS (PTS) WITH NEWLY DIAGNOSED DIFFUSE LARGE B-CELL LYMPHOMA (DLBCL): EPCORE DLBCL-2

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REAL-WORLD COMPARATIVE EFFECTIVENESS OF COVALENT BRUTON TYROSINE KINASE INHIBITORS (CBTKI) AMONG PATIENTS WITH RELAPSED/REFRACTORY MANTLE CELL LYMPHOMA (R/R MCL)

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EPCORE FL-1: PHASE 3 TRIAL OF SUBCUTANEOUS EPCORITAMAB WITH RITUXIMAB AND LENALIDOMIDE (R2) VS R2 ALONE IN PATIENTS WITH RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA

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REAL-WORLD OUTCOMES FOLLOWING ADVERSE EVENTS AND DOSE REDUCTION OF FIRST- LINE IBRUTINIB IN MEDICARE BENEFICIARIES WITH CHRONIC LYMPHOCYTIC LEUKEMIA/ SMALL LYMPHOCYTIC LYMPHOMA (CLL/SLL)

Mazyar Shadman, MD¹; Bhavini P. Srivastava, MSc²; Monika Salkar, PhD²; Chadi Saifan, MD²; Barnabie, C. Agatep, MPH³; Barton Jones, MS³; Olga Ryan, DrPH, MPH, MBA²; Shafeef Bacchus, PharmD²; Deborah M. Stephens, DO⁴

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ACCESS TO CAR T-CELL THERAPY IN THE US AND ITS POTENTIAL IMPACTS ON HEALTH INEQUITIES

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LATEST RESULTS FROM AN ONGOING FIRST-IN-HUMAN PHASE 1A/B STUDY OF NX-5948, A SELECTIVE BRUTON'S TYROSINE KINASE (BTK) DEGRADER, IN PATIENTS WITH RELAPSED/ REFRACTORY CLL AND OTHER B-CELL MALIGNANCIES

¹Nirav N. Shah¹, ²Kim Linton², ³Graham P. Collins², ⁴Francesco Forconi⁵, ⁵Karan Dixit⁶, ⁶Talha Munir⁷, ⁷Zulfa Omer⁸, ⁸Dima El-Sharkawi⁹, ⁹Jeanette Doorduijn¹⁰, ¹⁰Alvaro Alencar¹¹, ¹¹Pam McKay¹², ¹²John Riches¹³, ¹³Mary Gleeson¹⁴, ¹⁴David Lewis¹⁵, ¹⁵Allison Winter¹⁶, ¹⁶Sarah Injac¹⁶, ¹⁶Ted Shih¹⁶, ¹⁶Srinand Nandakumar¹⁶, ¹⁶May Tan¹⁶, ¹⁶Ganesh Cherala¹⁶, ¹⁶Erin Meredith¹⁷, ¹⁷Alexey Danilov

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INTEGRATIVE FUNCTIONAL GENOMICS AND SPATIAL TUMOR MICROENVIRONMENT ANALYSIS OF MANTLE CELL LYMPHOMA REVEAL PROGNOSTIC BIOLOGICAL SUBTYPES

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THE EFFICACY OF SUBCUTANEOUS EPCORITAMAB VS STANDARD OF CARE (SCHOLAR-5) IN PATIENTS WITH RELAPSED/REFRACTORY FOLLICULAR LYMPHOMA AFTER ≥2 SYSTEMIC THERAPIES: AN INDIRECT TREATMENT COMPARISON

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QUANTITATIVE SYSTEMS PHARMACOLOGY MODELING OF LONCASTUXIMAB TESIRINE COMBINED WITH MOSUNETUZUMAB AND GLOFITAMAB HELPS GUIDE DOSING FOR PATIENTS WITH DLBCL

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FINAL ANALYSIS OF THE RESONATE-2 STUDY: UP TO 10 YEARS OF FOLLOW-UP OF FIRST-LINE IBRUTINIB TREATMENT IN PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA/SMALL LYMPHOCYTIC LYMPHOMA

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EXTENDED FOLLOW-UP BEYOND 2.5 YEARS SHOWS LONG-TERM EFFICACY IN COMPLETE RESPONDERS FOLLOWING EPCORITAMAB MONOTHERAPY IN RELAPSED OR REFRACTORY LARGE B-CELL LYMPHOMA

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Sutton, UK; ⁷MSC National Research Institute of Oncology, Kraków, Poland; ⁸John Theurer Cancer Center at Hackensack Meridian Health, Hackensack Meridian Health School of Medicine, Hackensack, NJ, USA; ⁹The Christie NHS Foundation Trust, Manchester Cancer Research Centre, and Division of Cancer Sciences, University of Manchester, Manchester, UK; ¹⁰University of Nebraska Medical Center, Omaha, NE, USA; ¹¹University of Iowa, Iowa City, IA, USA; ¹²Samsung Medical Center, Seoul, Republic of Korea; ¹³AbbVie, North Chicago, IL, USA; ¹⁴Genmab, Plainsboro, NJ, USA; ¹⁵Cancer Research Center of Toulouse, Toulouse, France; ¹⁶On behalf of the Lunenburg Lymphoma Phase I/II Consortium-HOVON/LLPC, Erasmus MC Cancer Institute, University Medical Center, Department of Hematology, Rotterdam, Netherlands; ¹⁷Rigshospitalet and University of Copenhagen, Copenhagen, Denmark

PHASE 1B OPEN-LABEL STUDY OF LONCASTUXIMAB TESIRINE IN COMBINATION WITH OTHER ANTICANCER AGENTS IN PATIENTS WITH RELAPSED/REFRACTORY B-CELL NON-HODGKIN LYMPHOMA (LOTIS-7)

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MAINTENANCE OF MODERATE TO HIGH LEVELS OF FUNCTIONING AND QUALITY OF LIFE (QOL) WITH ODRONEXTAMAB MONOTHERAPY IN PATIENTS WITH RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA (R/R FL)

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EPCORITAMAB INDUCES DEEP RESPONSES IN RELAPSED OR REFRACTORY (R/R) FOLLICULAR LYMPHOMA (FL): SAFETY AND POOLED EFFICACY DATA FROM EPCORE NHL-1 PIVOTAL AND CYCLE (C) 1 OPTIMIZATION (OPT) FL COHORTS

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TRIAL IN PROGRESS: PHASE 3 STUDY OF ODRONEXTAMAB PLUS LENALIDOMIDE VERSUS RITUXIMAB PLUS LENALIDOMIDE IN PATIENTS WITH RELAPSED/REFRACTORY (R/R) FOLLICULAR LYMPHOMA (FL) AND MARGINAL ZONE LYMPHOMA (MZL) (OLYMPIA-5)

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PIRTOBRUTINIB IN PRIOR CBTKI R/R MCL: PHASE 1/2 BRUIN STUDY UPDATES INCLUDING HIGH-RISK SUBGROUP ANALYSES

Yucui Wang¹, Jonathon B. Cohen², Nirav N. Shah³, Wojciech Jurczak⁴, Pier Luigi Zinzani⁵, Chan Y. Cheah⁶, Toby A. Eyre⁷, Chaitra S. Ujjani⁸, Youngil Koh⁹, Won Seog Kim¹⁰, Sunita D. Nasta¹¹, Ian Flinn¹², Benoit Tessoulin¹³, Shuo Ma¹⁴, Alvaro J. Alencar¹⁵, David J. Lewis¹⁶, Jennifer A. Woyach¹⁷, Kami J Maddocks¹⁷, Krish Patel¹⁸, Joanna Rhodes¹⁹, Constantine S. Tam²⁰, John F. Seymour²¹, Hirokazu Nagai²², Julie M. Vose²³, Bita Fakhri²⁴, Marc S. Hoffmann²⁵, Francisco Hernandez-Ilizaliturri²⁶, Andrew D. Zelenetz²⁷, Anita Kumar²⁷, Talha Munir²⁸, Donald Tsai²⁹, Minna Balbas²⁹, Bin Liu²⁹, Amy S. Ruppert³⁰, Bastien Nguyen²⁹, Lindsey E. Roeker²⁷, Michael L. Wang³¹

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PRIMARY THYROID DIFFUSE LARGE B CELL LYMPHOMA MASKING AS LYMPHOCYTIC THYROIDITIS IN A PATIENT WITH HASHIMOTO'S THYROIDITIS AND SYSTEMIC LUPUS ERYTHEMATOSUS

Chalothorn Wannaphut MD¹, Ryon Nakasone MD², Jodi Kagihara MD²

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ALPHA3: A PHASE 2 STUDY EVALUATING THE SAFETY AND EFFICACY OF FIRST-LINE (1L) CONSOLIDATION WITH CEMACABTAGENE ANSEGEDLEUCEL (CEMA-CEL) IN PATIENTS WITH LARGE B-CELL LYMPHOMA (LBCL) AND MINIMAL RESIDUAL DISEASE (MRD) AFTER RESPONSE TO STANDARD THERAPY

Jason Westin, MD¹; John M. Burke, MD²; Houston E. Holmes, MD³; Edward J. Licitra, MD, PhD⁴; Rushang D. Patel, MD, PhD⁵; Amy Feng, PhD⁶; Lynn Navale, MS⁶; Srinivas Ghatta, PhD⁶; John B. Le Gall, MD⁶; Alex F. Herrera, MD⁷ on behalf of the ALPHA3 investigators

¹University of Texas MD Anderson Cancer Center, Houston, TX, USA; ²US Oncology Hematology Research Program, Rocky Mountain Cancer Centers, Aurora, CO, USA; ³Texas Oncology, Dallas, TX, USA; ⁴Astera Cancer Center, East Brunswick, NJ, USA; ⁵AdventHealth Cancer Institute, Orlando, FL, USA; ⁶Allogene Therapeutics, San Francisco, CA, USA; ⁷City of Hope National Medical Center, Duarte, CA, USA



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Reference: 1. Sharman JP, et al. Abstract presented at: ASH annual meeting; December 9-12, 2023; San Diego, CA. Abs 636.



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