

**Overview of key presentations featuring Roche medicines**

| Medicine                             | Abstract title                                                                                                                                                                                                                                                                                | Abstract number/presentation details                                                                                                                                                                                                      |
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| <p>Polivy® (polatuzumab vedotin)</p> | <p>Five-Year Analysis of the POLARIX Study: Prolonged Follow-up Confirms Positive Impact of Polatuzumab Vedotin Plus Rituximab, Cyclophosphamide, Doxorubicin, and Prednisone (Pola-R-CHP) on Outcomes</p>                                                                                    | <p>#469 oral presentation</p> <p>Session: 626. Aggressive Lymphomas: Clinical and Epidemiological: CARs, Bispecifics, and ADCs: Progress and Challenges in Aggressive B Cell Lymphoma</p> <p>Sunday 8 December 2024</p> <p>9.30am PST</p> |
|                                      | <p>A Multicenter, Prospective, Observational Study of Pola-R-CHP in Patients With Previously Untreated Diffuse Large B-Cell Lymphoma (POLASTAR): A Preliminary Analysis</p>                                                                                                                   | <p>#4475 poster presentation</p> <p>Session: 626. Aggressive Lymphomas: Clinical and Epidemiological: Poster III</p> <p>Monday 9 December 2024</p> <p>6pm-8pm PST</p>                                                                     |
| <p>Lunsumio® (mosunetuzumab)</p>     | <p>A Randomized Phase II Study of Mosunetuzumab SC Plus Polatuzumab Vedotin Demonstrates Improved Outcomes Versus Rituximab Plus Polatuzumab Vedotin in Patients (Pts) with Relapsed or Refractory (R/R) Large B-Cell Lymphoma (LBCL)</p>                                                     | <p>#989 oral presentation</p> <p>Session: 627. Aggressive Lymphomas: Pharmacologic Therapies: Chemotherapy-free Combinations for Relapsed Aggressive Lymphomas</p> <p>Monday 9 December 2024</p> <p>5.30pm PST</p>                        |
|                                      | <p>Subcutaneous Mosunetuzumab Leads to High Rates of Durable Responses, Low Rates of Cytokine Release Syndrome, and Non-Inferior Exposure Compared with Intravenous Administration in Patients with Relapsed/Refractory Follicular Lymphoma: Primary Analysis of a Pivotal Phase II Study</p> | <p>#1645 poster presentation</p> <p>Session: 623. Mantle Cell, Follicular, Waldenstrom's, and Other Indolent B Cell Lymphomas: Clinical and Epidemiological: Poster I</p> <p>Saturday 7 December 2024</p> <p>5.30pm-7.30pm PST</p>        |
|                                      | <p>Mosunetuzumab Continues to Demonstrate Clinically Meaningful Outcomes in Patients with Relapsed and/or Refractory Follicular Lymphoma after ≥2 Prior Therapies Including Those with a History of POD24: 4-Year Follow-up of a Pivotal Phase II Study</p>                                   | <p>#4407 poster presentation</p> <p>Session: 623. Mantle Cell, Follicular, Waldenstrom's, and Other Indolent B Cell Lymphomas: Clinical and Epidemiological: Poster III</p> <p>Monday 9 December 2024</p>                                 |

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|                       | Mosunetuzumab Monotherapy Demonstrates Encouraging Activity and a Manageable Safety Profile in Patients with Heavily Pre-Treated Relapsed or Refractory Mantle Cell Lymphoma                                                                                                                       | #1646 poster presentation<br>Session: 623. Mantle Cell, Follicular, Waldenstrom's, and Other Indolent B Cell Lymphomas: Clinical and Epidemiological: Poster I<br><br>Saturday 7 December 2024<br><br>5.30pm-7.30pm PST                 |
|                       | Fixed-Duration Subcutaneous Mosunetuzumab is Active and has a Manageable Safety Profile in Patients with Previously Untreated, Low-Tumor Burden Follicular Lymphoma: Updated Results from the Phase II MorningSun Study                                                                            | #3008 poster presentation<br>Session: 623. Mantle Cell, Follicular, Waldenstrom's, and Other Indolent B Cell Lymphomas: Clinical and Epidemiological: Poster II<br><br>Sunday 8 December 2024<br><br>6pm-8pm PST                        |
|                       | Travel Burden and Travel Costs of Bispecific Antibodies in Patients with Relapsed/Refractory Diffuse Large B-Cell Lymphoma and Relapsed/Refractory Follicular Lymphoma                                                                                                                             | #782 oral presentation<br>Session: 902. Health Services and Quality Improvement: Lymphoid Malignancies: For a Better Tomorrow - Improving Access to Blood Cancer Treatments and Trials<br><br>Monday 9 December 2024<br><br>10.45am PST |
| Columvi® (glofitamab) | Fixed-duration Glofitamab Monotherapy Continues to Demonstrate Durable Responses in Patients with Relapsed or Refractory Large B-Cell Lymphoma: 3-year Follow-Up From a Pivotal Phase II Study                                                                                                     | #865 oral presentation<br>Session: 627. Aggressive Lymphomas: Pharmacologic Therapies: New R-CHOP Combinations for Treatment Naïve DLBCL<br><br>Sunday 8 December 2024<br><br>Midday PST                                                |
|                       | Glofitamab in Combination with Polatuzumab Vedotin Maintains Durable Responses and a Manageable Safety Profile in Patients with Heavily Pre-treated Relapsed/Refractory (R/R) Large B-Cell Lymphoma (LBCL) Including High-Grade B-Cell Lymphoma (HGBCL): Extended Follow-Up of a Phase Ib/II Study | #988 oral presentation<br>Session: 627. Aggressive Lymphomas: Pharmacologic Therapies: Chemotherapy-free Combinations for Relapsed Aggressive Lymphoma<br><br>Monday 9 December 2024                                                    |

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|                  |                                                                                                                                                                                                                                                                                                                         | 5.15pm PST                                                                                                                                                                                                      |
|                  | Glofitamab in Combination with Rituximab plus Ifosfamide, Carboplatin, and Etoposide shows Favorable Efficacy and Manageable Safety in Patients with Relapsed or Refractory Diffuse Large B-cell Lymphoma, Eligible for Stem Cell Transplant or Chimeric Antigen Receptor T-cell Therapy: Results from a Phase Ib Study | #987 oral presentation<br>Session: 627. Aggressive Lymphomas: Pharmacologic Therapies: Chemotherapy-free Combinations for Relapsed Aggressive Lymphomas<br>Monday 9 December 2024<br>5pm PST                    |
|                  | Primary Results of Patient-Reported Outcomes in Patients with Relapsed/Refractory Diffuse Large B-cell Lymphoma Treated with Glofitamab plus Gemcitabine and Oxaliplatin (Glofit-GemOx) Versus Rituximab plus GemOx (R-GemOx) from the Phase III STARGLO Study                                                          | #5132 poster presentation<br>Session: 906. Outcomes Research: Lymphoid Malignancies Excluding Plasma Cell Disorders: Poster III<br>Monday 9 December 2024<br>6pm-8pm PST                                        |
|                  | 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                                                                           | #1631 poster presentation<br>Session: 623. Mantle Cell, Follicular, Waldenstrom's, and Other Indolent B Cell Lymphomas: Clinical and Epidemiological: Poster I<br>Saturday 7 December 2024<br>5.30pm-7.30pm PST |
|                  | A Healthcare Utilization Model Comparing Time Toxicity Between Glofitamab and Epcoritamab Treatment Regimens                                                                                                                                                                                                            | #3647 poster presentation<br>Session: 902. Health Services and Quality Improvement: Lymphoid Malignancies: Poster II<br>Sunday 8 December 2024<br>6pm-8pm PST                                                   |
| Englumafusp alfa | Englumafusp alfa (CD19-4-1BBL) combined with glofitamab is safe and efficacious in patients with r/r B-NHL: extended follow up analysis of the dose-escalation part of Phase 1 trial BP41072                                                                                                                            | #990 oral presentation<br>Session: 627. Aggressive Lymphomas: Pharmacologic Therapies: Chemotherapy-free Combinations for Relapsed Aggressive Lymphomas<br>Monday 9 December 2024<br>5.45pm PST                 |
| Cevostamab       | Cevostamab in Patients with Heavily Pretreated Relapsed/Refractory Multiple Myeloma (RRMM):                                                                                                                                                                                                                             | #1021 oral presentation                                                                                                                                                                                         |

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|                                                               | Updated Results from an Ongoing Phase I Study Demonstrate Clinically Meaningful Activity and Manageable Safety and Inform the Doses and Regimen for Combination Studies                                                                      | Session: 654. Multiple Myeloma: Pharmacologic Therapies: Into the Future: New Drugs and Combinations in Multiple Myeloma<br><br>Monday 9 December 2024<br><br>4.30pm PST             |
| Venclexta®/Venclyxto® (venetoclax)*                           | 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 | #3237 poster presentation<br><br>Session: 642. Chronic Lymphocytic Leukemia: Clinical and Epidemiological: Poster II<br><br>Sunday 8 December 2024<br><br>6pm-8pm PST                |
| P-CD19CD20-ALLO1 (in collaboration with Poseida Therapeutics) | P-CD19CD20-ALLO1: Potent Fully Allogeneic CAR-T Therapy Targeting CD19 and CD20 with Superior Efficacy Over Single-Target Products                                                                                                           | #4805 poster presentation<br><br>Session: 702. CAR-T Cell Therapies: Basic and Translational: Poster III<br><br>Monday 9 December 2024<br>6pm-8pm PST                                |
| P-BCMA-ALLO1 (in collaboration with Poseida Therapeutics)     | A Phase 1 Study of P-BCMA-ALLO1, a Non-viral, Allogeneic BCMA Directed CAR-T in Relapsed/Refractory Multiple Myeloma (RRMM): Results from Optimized Lymphodepletion Cohort                                                                   | #4828 poster presentation<br><br>Session: 704. Cellular Immunotherapies: Early Phase Clinical Trials and Toxicities: Poster III<br><br>Monday 9 December 2024<br>6pm-8pm PST         |
| Hemlibra® (emicizumab)                                        | Bleed Patterns in Infants, From Birth to 12 Months of Age, with Hemophilia A Treated with Emicizumab: Exploratory Analysis of the HAVEN 7 Study                                                                                              | #1214 poster presentation<br><br>Session: 322. Hemophilia A and B: Clinical and Epidemiological: Poster I<br><br>Saturday 7 December 2024<br><br>5.30pm-7.30pm PST                   |
|                                                               | Real-World Experience With Emicizumab for Hemophilia A From the Physician Perspective Based on Survey Data                                                                                                                                   | #5078 poster presentation<br><br>Session: 905. Outcomes Research: Non-Malignant Conditions Excluding Hemoglobinopathies: Poster III<br><br>Monday 9 December 2024<br><br>6pm-8pm PST |
|                                                               | Phase III Randomized COMMODORE 2 Trial: 2-Year Efficacy and Safety of Crovalimab in Patients With                                                                                                                                            | #2687 poster presentation<br><br>Session: 508. Bone Marrow Failure:                                                                                                                  |

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| PiaSky® (crovalimab) | Paroxysmal Nocturnal Hemoglobinuria (PNH) Naive to Complement Inhibition                                                                                       | Acquired: Poster II<br>Sunday 8 December 2024<br>6pm-8pm PST                                                                     |
|                      | Phase III COMMODORE 1 Trial: 2-Year Efficacy and Safety of Crovalimab in Patients with Paroxysmal Nocturnal Hemoglobinuria (PNH) Who Switched from Ravulizumab | #4078 poster presentation<br>Session: 508. Bone Marrow Failure:<br>Acquired: Poster III<br>Monday 9 December 2024<br>6pm-8pm PST |

*\*Venclexta/Venclyxto is being developed by AbbVie and Roche. It is jointly commercialised by AbbVie and Genentech, a member of the Roche Group, in the US, and commercialised by AbbVie outside of the US.*