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Optimizing Treatment for Patients with Relapsed/Refractory Chronic Lymphocytic Leukemia

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Disclosures

| | |
|--|--|
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| Contracted Research | Ascentage Pharma, AstraZeneca Pharmaceuticals LP, MEI Pharma Inc, Novartis |
| Nonrelevant Financial Relationships | UpToDate |

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Program Steering Committee



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Key Datasets

- Thompson PA, Tam CS. **Pirtobrutinib**: A new hope for patients with **BTK inhibitor-refractory** lymphoproliferative disorders. *Blood* 2023 June 29;141(26):3137-42.
- Shah NN et al. **Pirtobrutinib monotherapy** in Bruton tyrosine kinase inhibitor-intolerant patients with B-cell malignancies: Results of the **phase I/II BRUIN trial**. *Haematologica* 2025 January 1;110(1):92-102.
- Sharman JP et al. **Phase III trial of pirtobrutinib versus idelalisib/rituximab or bendamustine/rituximab** in covalent Bruton tyrosine kinase inhibitor-pretreated chronic lymphocytic leukemia/small lymphocytic lymphoma (**BRUIN CLL-321**). *J Clin Oncol* 2025 August;43(22):2538-49.
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- Jurczak W et al. **Long-term safety with ≥12 months of pirtobrutinib** in relapsed/refractory (R/R) B-cell malignancies. EHA 2023;Abstract P618.
- Eyre TA et al. **BRUIN CLL-322**: A phase 3 open-label, randomized study of **fixed duration pirtobrutinib plus venetoclax and rituximab versus venetoclax and rituximab** in previously treated chronic lymphocytic leukemia/small lymphocytic lymphoma. ASCO 2023;Abstract TPS7583.
- Woyach J et al. **Pirtobrutinib vs ibrutinib in treatment-naïve and relapsed/refractory CLL/SLL**: Results from the first randomized **phase III study** comparing a non-covalent and covalent BTK inhibitor. ASH 2025;Abstract 683.
- Jurczak W et al. **Pirtobrutinib vs bendamustine plus rituximab (BR)** in patients with CLL/SLL: First results from a randomized **phase III study** examining a non-covalent BTK inhibitor in **untreated patients**. ASH 2025;Abstract LBA3.

Key Datasets

- Siddiqi T et al. **Lisocabtagene maraleucel (liso-cel) in R/R CLL/SLL: 24-month median follow-up of TRANSCEND CLL 004.** ASH 2023;Abstract 330.
- Wierda WG et al. **Lisocabtagene maraleucel (liso-cel) combined with ibrutinib (ibr) for patients (pts) with relapsed or refractory (R/R) chronic lymphocytic leukemia (CLL)/small lymphocytic lymphoma (SLL): Primary results from the open-label, phase 1/2 Transcend CLL 004 study.** ASH 2024;Abstract 887.

Key Datasets

- Cassanello G et al. **Trial watch: Bispecific antibodies** for the treatment of relapsed or refractory large B-cell lymphoma. *Oncoimmunology* 2024 March 3;13(1):2321648.
- Danilov A et al. **Epcoritamab monotherapy** in patients (pts) with relapsed or refractory (R/R) chronic lymphocytic leukemia (CLL): Results from **CLL expansion and optimization cohorts of Epcore CLL-1**. ASH 2024;Abstract 883.
- Ahn I et al. **Updated efficacy and safety results** of the Bruton tyrosine kinase (**BTK**) **degrader BGB-16673** in patients with relapsed/refractory chronic lymphocytic leukemia/small lymphocytic lymphoma (CLL/SLL) from the ongoing **phase 1 CaDAnCe-101 study**. ASH 2025;Abstract 85.
- Shah NN et al. **Efficacy and safety** of the Bruton's tyrosine kinase (**BTK**) **degrader NX-5948** in patients with relapsed/refractory (R/R) chronic lymphocytic leukemia (CLL): Updated results from an **ongoing phase 1a/b study**. ASH 2024;Abstract 884.
- Woyach JA et al. **First-in-human study** of the **reversible BTK inhibitor nemtabrutinib** in patients with relapsed/refractory chronic lymphocytic leukemia and B-cell non-Hodgkin lymphoma. *Cancer Discov* 2024;14(1):66-75.

Management of Double-Refractory CLL

Introduction: Sequencing of Treatment for CLL

Module 1: Clinician Survey Results

Module 2: Noncovalent BTK Inhibitor Pirtobrutinib

Module 3: Clinician Survey Results

Module 4: CAR T-Cell Therapy

Module 5: Clinician Survey Results

Module 6: Bispecific Antibodies and Promising Investigational Strategies

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Targeted Therapy Sequencing for CLL

cBTKi

BCL2i
+CD20

cBTKi + BCL2i not included here

Factors affecting timelines:

- Age
- Del(17p) / *TP53*-m
- IGHV-MS / Del(11q)
- Complex karyotype

1

2

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11

12

13

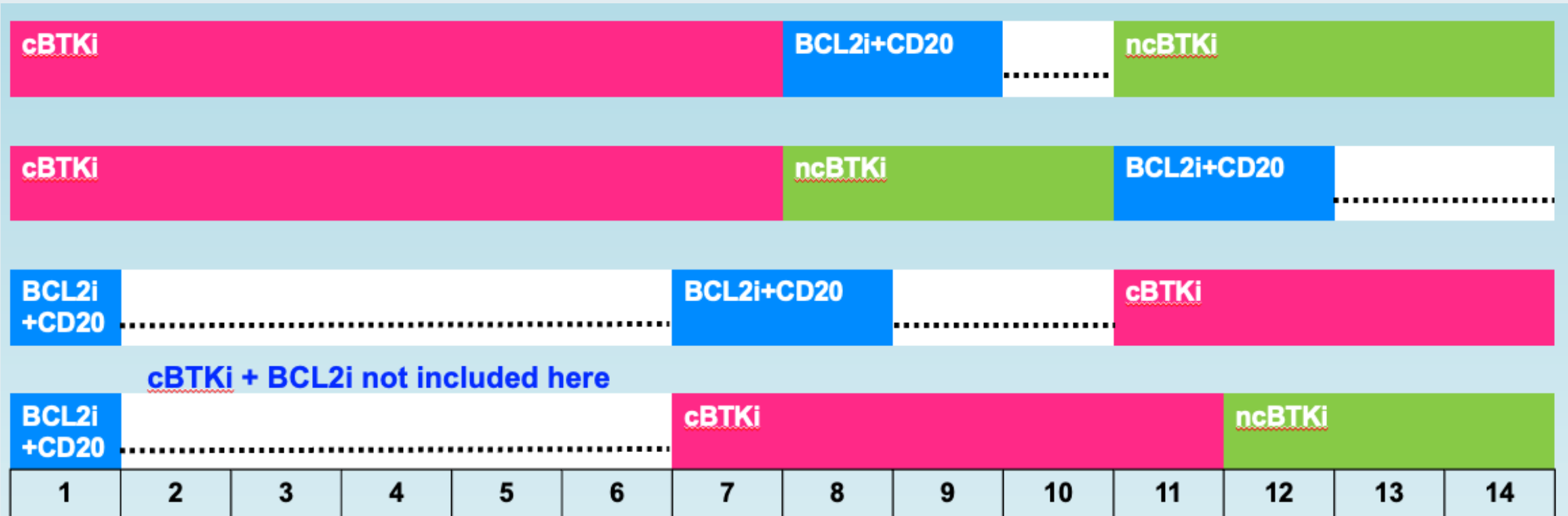
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Years

Double Exposed vs. Double Refractory:

- Exposed ≠ Refractory
- Refractory=progression on treatment

Targeted Therapy Sequencing for CLL



cBTKi + BCL2i not included here

Factors affecting timelines:

- Age
- Del(17p) / TP53-m
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- Complex karyotype

Years

Double Exposed vs. Double Refractory:

- Exposed ≠ Refractory
- Refractory=progression on treatment

FDA Approves Acalabrutinib with Venetoclax for Chronic Lymphocytic Leukemia or Small Lymphocytic Lymphoma

Press Release: February 19, 2026

“On February 19, 2026, the Food and Drug Administration approved acalabrutinib tablets and capsules in combination with venetoclax for adults with chronic lymphocytic leukemia (CLL) or small lymphocytic lymphoma (SLL).

Efficacy was evaluated in AMPLIFY (NCT03836261), a randomized, multicenter trial in adult patients previously untreated for CLL without del(17p) or TP53 mutation. Patients were randomized to receive acalabrutinib and venetoclax (AV) or Investigator’s choice of chemotherapy (fludarabine plus cyclophosphamide plus rituximab [FCR] or bendamustine plus rituximab [BR]).

The major efficacy outcome measure was progression-free survival (PFS) as assessed by independent review committee for the AV arm versus the investigator’s choice arm (FCR/BR). The median duration of PFS follow-up was 42.6 months. Median PFS was not estimable (NE) (95% CI: 51.1, NE) in the AV arm and 47.6 months (95% CI: 43.3, NE) in the FCR/BR arm (Hazard ratio 0.65 [95% CI: 0.49, 0.87]; *p*-value 0.0038). With a median follow-up of 41.0 months, there were 18 (6%) deaths in the AV arm and 42 (14%) in the FCR/BR arm.”

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Module 6: Bispecific Antibodies and Promising Investigational Strategies

75-year-old patient with relapsed CLL, no del(17p) or TP53 mutation:
Covalent BTKi 6 years → PD → venetoclax/anti-CD20 antibody 2 years →
3 years observation → PD



Dr Awan

Pirtobrutinib or venetoclax/anti-CD30 antibody



Dr Coombs

Venetoclax/anti-CD20 antibody



Dr Davids

Venetoclax/anti-CD20 antibody



Dr Fakhri

Pirtobrutinib



Dr Lamanna

Venetoclax/anti-CD20 antibody



Dr Sharman

Pirtobrutinib bridge to CAR T



Dr Woyach

Venetoclax/anti-CD20 antibody

In general, what is the minimum duration of remission after second-line venetoclax/
anti-CD20 antibody before you would consider retreatment as third-line therapy?



Dr Awan

24 months



Dr Coombs

12 months



Dr Davids

12 months



Dr Fakhri

24 months



Dr Lamanna

>24 months



Dr Sharman

24-36 months



Dr Woyach

36 months

75-year-old patient with relapsed CLL, no del(17p) or TP53 mutation:
Covalent BTKi 6 years → PD → venetoclax/anti-CD20 antibody 2 years →
1 year observation → PD



Dr Awan

Pirtobrutinib



Dr Coombs

Pirtobrutinib



Dr Davids

Pirtobrutinib



Dr Fakhri

Pirtobrutinib



Dr Lamanna

Pirtobrutinib



Dr Sharman

Pirtobrutinib bridge to CAR T



Dr Woyach

Pirtobrutinib

75-year-old patient with relapsed CLL, no del(17p) or TP53 mutation:

Covalent BTKi 2 years → responds but stops due to subdural hematoma → 3 years observation
→ PD → venetoclax/anti-CD20 antibody 2 years → 6 months observation → PD



Dr Awan

Pirtobrutinib



Dr Coombs

Pirtobrutinib



Dr Davids

Venetoclax monotherapy



Dr Fakhri

CAR T-cell therapy or epcoritamab on clinical trial



Dr Lamanna

Venetoclax monotherapy



Dr Sharman

Pirtobrutinib or CAR T



Dr Woyach

Pirtobrutinib or CAR T

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- Jurczak W et al. **Pirtobrutinib vs bendamustine plus rituximab (BR)** in patients with CLL/SLL: First results from a randomized **phase III study** examining a non-covalent BTK inhibitor in **untreated patients**. ASH 2025;Abstract LBA3.

Key Differences Between Available Covalent and Reversible Bruton Tyrosine Kinase (BTK) Inhibitors

| | Ibrutinib | Acalabrutinib | Zanubrutinib | Pirtobrutinib |
|---------------------------------|--|----------------------|--|---|
| BTK binding | Covalent C481 | Covalent C481 | Covalent C481 | Reversible ATP pocket Distant from C481 |
| Half-life | 6 hours | 1 hour | 4 hours | 20 hours >90% BTK inhibition |
| BTK Y223 autophosphorylation | Inhibited | Inhibited | Inhibited | Inhibited |
| BTK Y551 phosphorylation | No effect | No effect | No effect | Inhibited (maintenance of closed conformation) |
| BTK C481S mutation | Common | Reported | Reported | Not described Effective against C481S |
| Kinase-dead mutations | Uncommon and restricted to C481* (active against HCK) | Not reported to date | Reported: L528W > C481Y | Reported: L528W > V416L, A428D, C481R, M477I, and M437R |
| T474I/T474L gatekeeper mutation | Uncommon*; active against T474I and T474L | Reported | Not reported to date | Reported |
| Off-target hits† | BLK BMX BRK EGFR HER2 HER4 ITK JAK3 RLK TEC | HER4 | BLK BMX BRK EGFR HER4 RLK | HER4 BRK |

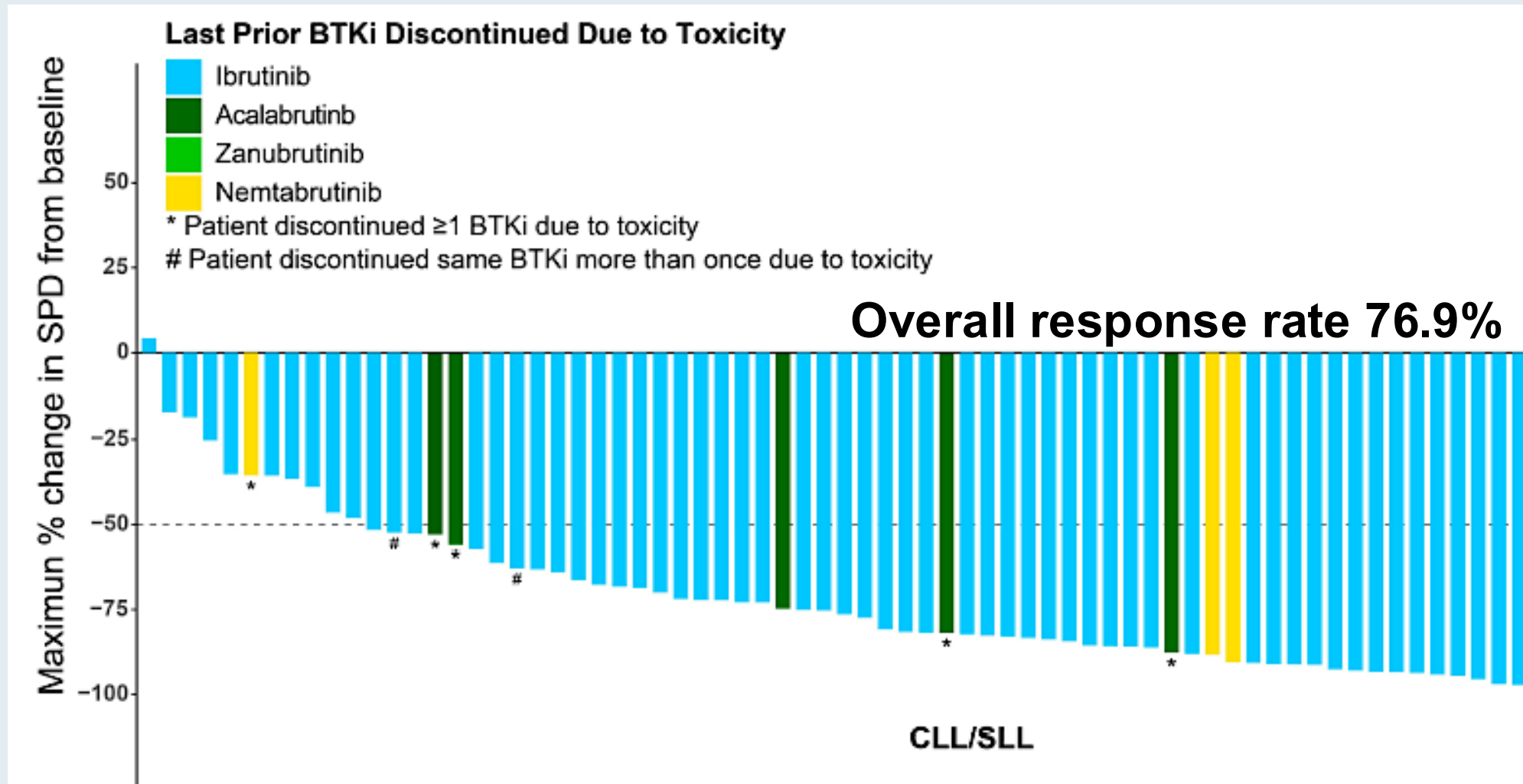


Pirtobrutinib monotherapy in Bruton tyrosine kinase inhibitor-intolerant patients with B-cell malignancies: results of the phase I/II BRUIN trial

by Nirav N. Shah, Michael Wang, Lindsey E. Roeker, Krish Patel, Jennifer A. Woyach, William G. Wierda, Chaitra S. Ujjani, Toby A. Eyre, Pier Luigi Zinzani, Alvaro J. Alencar, Paolo Ghia, Nicole Lamanna, Marc S. Hoffmann, Manish R. Patel, Ian Flinn, James N. Gerson, Shuo Ma, Catherine C. Coombs, Chan Y. Cheah, Ewa Lech-Maranda, Bitu Fakhri, Won Seog Kim, Minal A. Barve, Jonathon B. Cohen, Wojciech Jurczak, Talha Munir, Meghan C. Thompson, Donald E. Tsai, Katherine Bao, Nicholas A. Cangemi, Jennifer F. Kherani, Richard A. Walgren, Hongmei Han, Amy S. Ruppert, and Jennifer R. Brown

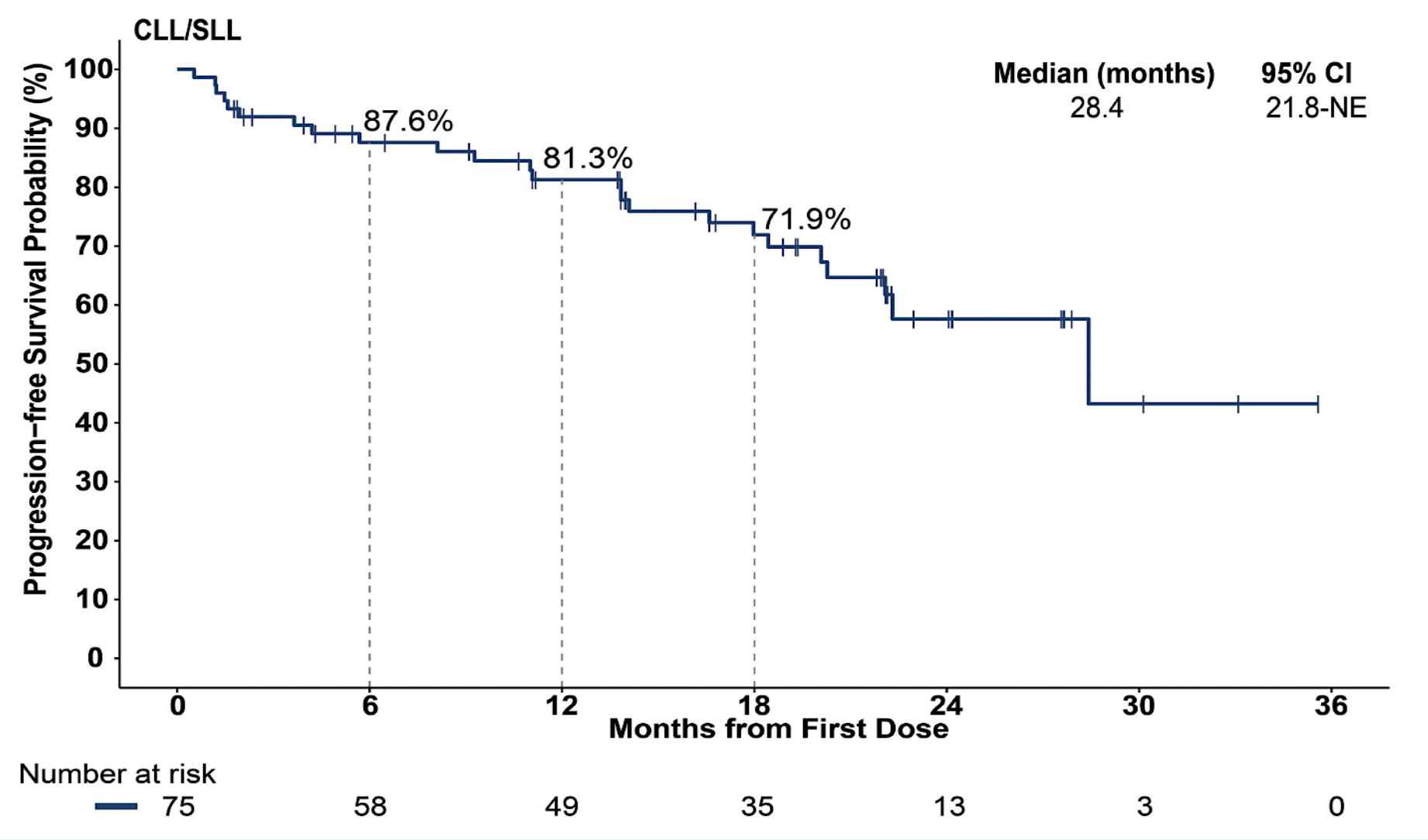
Haematologica 2025;110(1):92-102.

BRUIN: Pirtobrutinib Efficacy in Patients with CLL or SLL Who Received Prior BTK Inhibitor (BTKi) Treatment



SLL = small lymphocytic lymphoma; SPD = sum of product diameters

BRUIN: Median Progression-Free Survival for Patients with CLL/SLL



Shah NN et al. *Haematologica* 2025;110(1):92-102.



BRUIN: Pirtobrutinib Safety Profile in the Overall Population

| AE | BTKi-intolerant (n=127) | | | |
|--|-------------------------|-----------------|--------------------------|-----------------|
| | All cause AEs, % | | Treatment-related AEs, % | |
| | Any Grade | Grade ≥3 | Any Grade | Grade ≥3 |
| Fatigue | 39.4 | 3.9 | 9.4 | 1.6 |
| Neutropenia ^a | 37.0 | 31.5 | 21.3 | 17.3 |
| Diarrhea | 29.9 | 1.6 | 12.6 | 0.8 |
| Contusion | 29.1 | 0.0 | 22.0 | 0.0 |
| Cough | 26.8 | 0.0 | 4.7 | 0.0 |
| Headache | 25.2 | 0.8 | 7.1 | 0.8 |
| COVID-19 | 22.8 | 4.7 | 0.0 | 0.0 |
| Abdominal pain | 22.0 | 2.4 | 4.7 | 0.8 |
| Dyspnea | 22.0 | 2.4 | 5.5 | 0.0 |
| Nausea | 20.5 | 0.0 | 4.7 | 0.0 |
| AEs of Interest^b | Any Grade | Grade ≥3 | Any Grade | Grade ≥3 |
| Infections ^c | 68.5 | 24.4 | 14.2 | 5.5 |
| Infections (excluding COVID-19) | 59.8 | 17.3 | 14.2 | 5.5 |
| Bruising ^d | 36.2 | 0.0 | 26.8 | 0.0 |
| Rash ^e | 22.8 | 0.8 | 8.7 | 0.8 |
| Arthralgia | 21.3 | 0.8 | 4.7 | 0.0 |
| Hemorrhage/hematoma ^f | 14.2 | 3.1 | 4.7 | 0.8 |
| Hypertension | 7.9 | 0.8 | 3.1 | 0.0 |
| Atrial fibrillation/flutter ^g | 4.7 | 1.6 | 0.8 | 0.0 |

FDA Grants Traditional Approval to Pirtobrutinib for Chronic Lymphocytic Leukemia and Small Lymphocytic Lymphoma





Press Release: December 3, 2025

“... the Food and Drug Administration granted traditional approval to pirtobrutinib for adults with relapsed or refractory chronic lymphocytic leukemia or small lymphocytic lymphoma (CLL/SLL) who have previously been treated with a covalent BTK inhibitor.

Efficacy was evaluated in BRUIN-CLL-321 (NCT 04666038), a randomized, open-label, active-controlled trial. The trial randomized 238 patients who were previously treated for CLL/SLL, including a covalent BTK inhibitor. Patients previously treated with a non-covalent BTK inhibitor were not permitted. Patients were randomized (1:1) to receive either pirtobrutinib or investigator’s choice of idelalisib plus a rituximab product (IR) or bendamustine plus a rituximab product (BR).

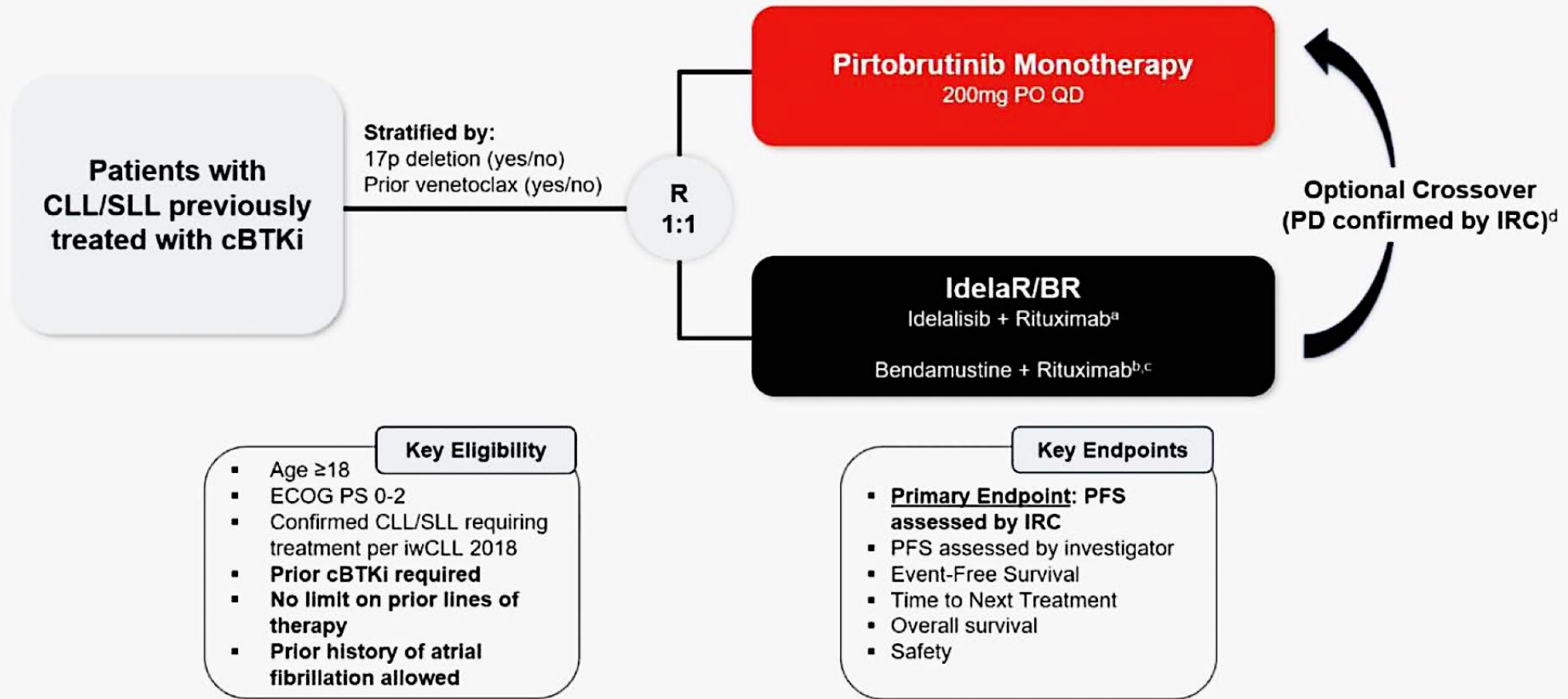
Median PFS was 11.2 months (95% CI: 9.5, 11.4) in the pirtobrutinib arm and 8.7 months (95% CI: 7.2, 10.2) in the investigator’s choice of IR/BR arm (Hazard ratio 0.58 [95% CI: 0.38, 0.89]; p-value 0.0105). Of the 119 patients in the investigator’s choice arm, 50 crossed over to receive pirtobrutinib therapy. At an updated analysis with a median follow-up time of 19.8 months, the HR for overall survival (OS) was 1.09 (95% CI: 0.68, 1.75).”

Phase III Trial of Pirtobrutinib Versus Idelalisib/Rituximab or Bendamustine/Rituximab in Covalent Bruton Tyrosine Kinase Inhibitor–Pretreated Chronic Lymphocytic Leukemia/Small Lymphocytic Lymphoma (BRUIN CLL-321)

Jeff P. Sharman, MD¹ ; Talha Munir, PhD, MBBS² ; Sebastian Grosicki, MD, PhD³; Lindsey E. Roeker, MD⁴; John M. Burke, MD⁵ ; Christine I. Chen, MHPE, MD⁶; Norbert Grzasko, MD, PhD⁷ ; George Follows, PhD, MA, BM, BCh, FRCP⁸; Zoltán Mátrai, MD, PhD⁹; Alessandro Sanna, MD¹⁰ ; Lugui Qiu, MD¹¹; Ru Feng, MD¹² ; Vu Minh Hua, PhD, MBBS, FRACP, FRCPA¹³; Wojciech Jurczak, MD, PhD¹⁴; Matthias Ritgen, MD¹⁵ ; Shuhua Yi, MD¹⁶ ; Francesc Bosch, MD, PhD¹⁷ ; Catherine C. Coombs, MD¹⁸; Katherine Bao, PhD¹⁹ ; Vishalkumar Patel, MD¹⁹; Bin Liu, MSc, MPH¹⁹; Livia Compte, MD, PhD¹⁹ ; Ananya Guntur, PhD¹⁹; Denise Y. Wang, PhD¹⁹; Marisa Hill, MS, MD¹⁹; Ching Ching Leow, PhD¹⁹; Paolo Ghia, MD, PhD²⁰ ; and Paul M. Barr, MD²¹ 

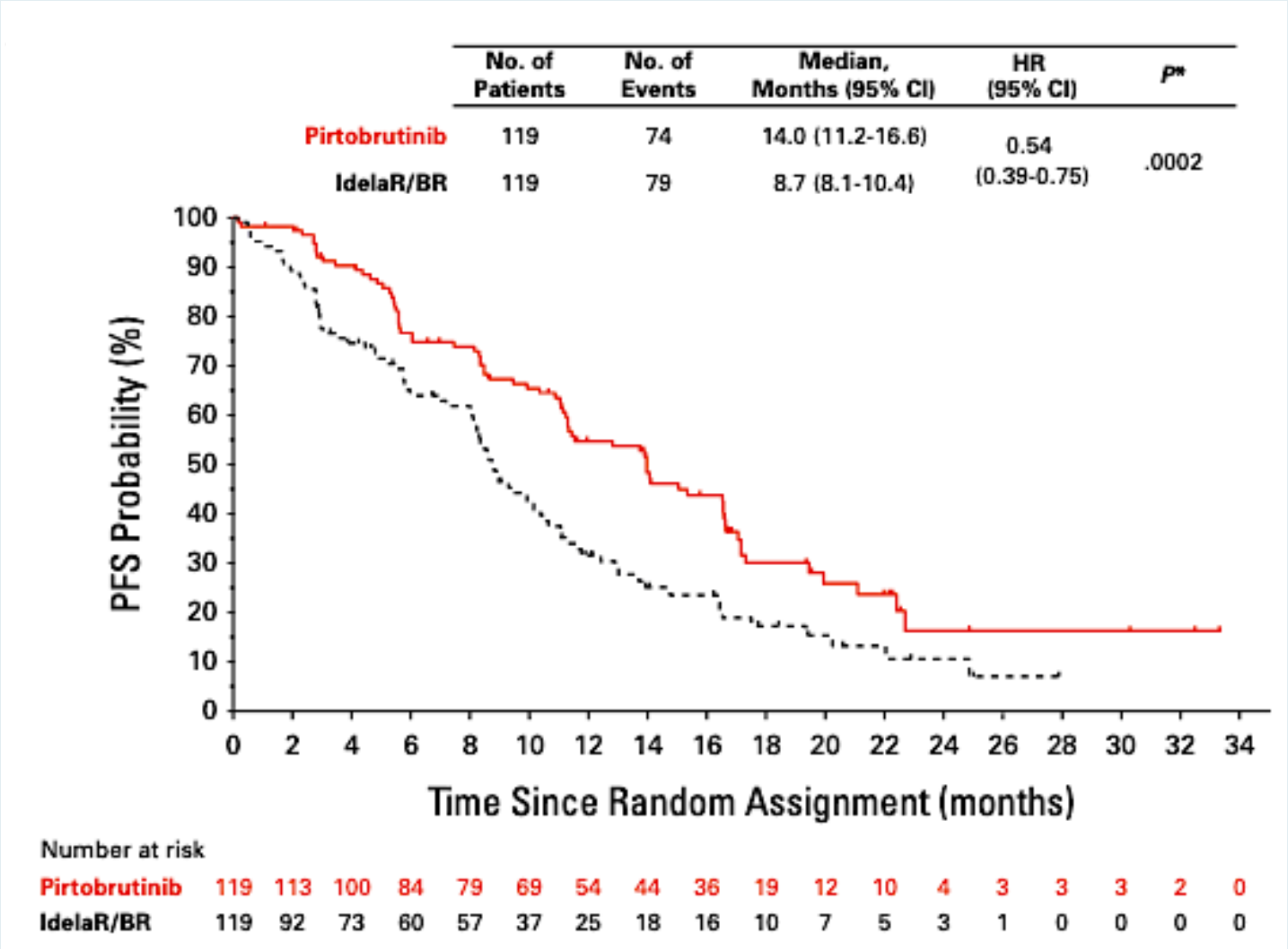
J Clin Oncol 2025;43:2538-49.

BRUIN CLL-321: A Phase III Trial of Pirtobrutinib Monotherapy for Relapsed/Refractory CLL



Treatment was given in 28-day cycles. PFS assessed based on iwCLL2018. ^aIdelalisib dosed at 150mg PO BID. Day 1 of cycle 1, first dose of rituximab at 375 mg/m², next 4 infusions at 500 mg/m² every 2 weeks, next 3 infusions at 500 mg/m² every 4 weeks. ^bBendamustine (70 mg/m²) administered IV D1, D2 of cycles 1-6. ^cDay 1 of cycle 1, first dose of rituximab at 375 mg/m², next 5 infusions day 1 of cycle 2 through cycle 6 at 500 mg/m². ^dEligible patients receiving investigator's choice of IdelaR/BR could crossover to receive pirtobrutinib monotherapy upon confirmation of PD by IRC per protocol. Abbreviations: BID, twice daily; BR, bendamustine + rituximab; cBTKi, covalent Bruton tyrosine kinase inhibitor; CLL, chronic lymphocytic leukemia; ECOG PS, Eastern Cooperative Oncology Group Performance Status; IdelaR, idelalisib + rituximab; IRC, Independent Review Committee; iwCLL, international workshop on chronic lymphocytic leukemia; mg, milligram; PD, progressive disease; PFS, progression free survival; PO, by mouth; QD, once daily; R, randomized; SLL, small lymphocytic lymphoma.

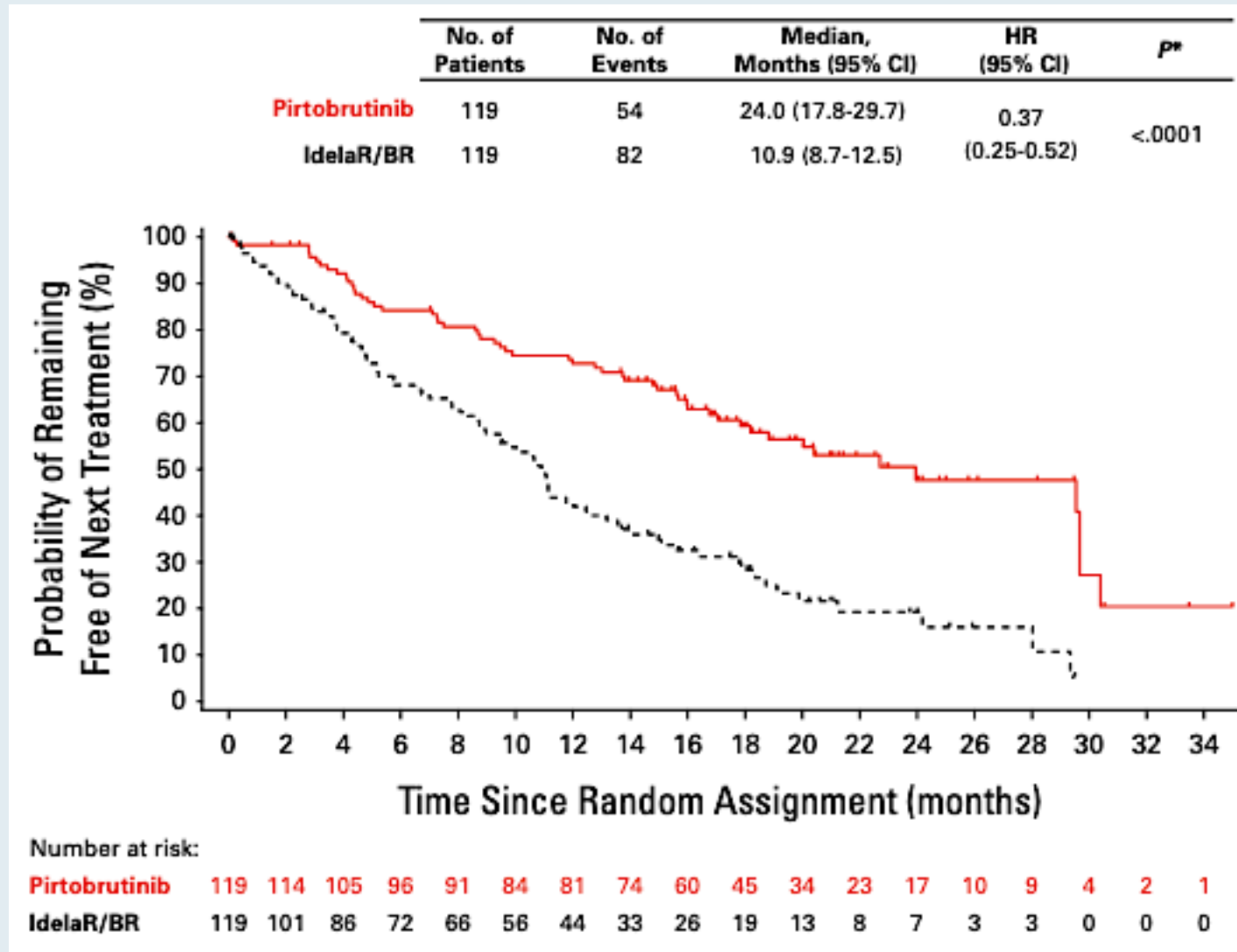
BRUIN CLL-321: IRC-Assessed Progression-Free Survival (PFS)



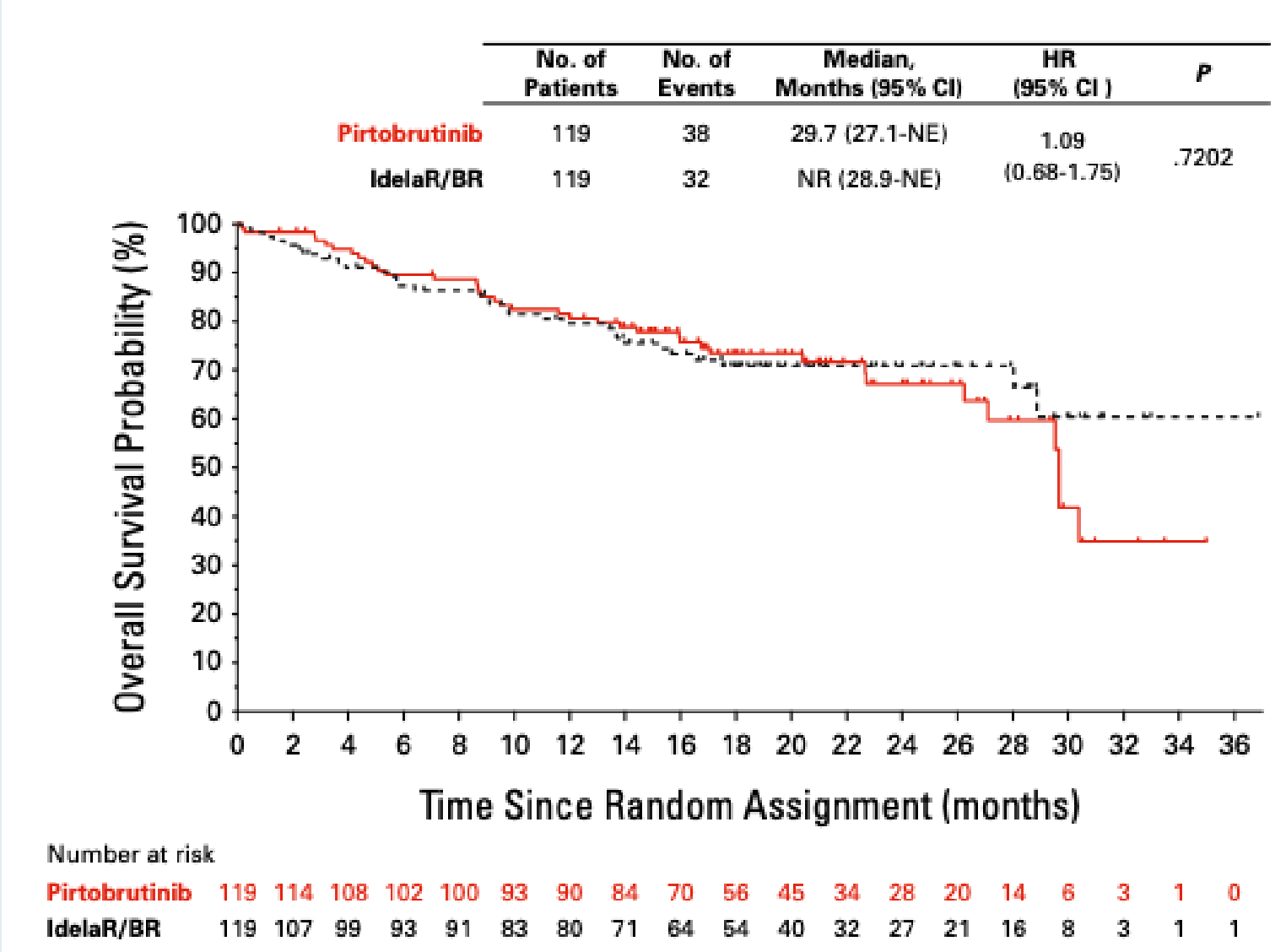
IRC = independent review committee



BRUIN CLL-321: Time to Next Treatment



BRUIN CLL-321: Overall Survival



BRUIN CLL-321: Safety Profile

| TEAE | Pirtobrutinib (n = 116), IR ^a | IdelaR or BR (n = 109), IR ^a | IRR (95% CI) ^b | P ^c |
|--------------------------|--|---|---------------------------|----------------|
| Infections ^d | 94.5 | 125.5 | 0.75 (0.53 to 1.07) | .11 |
| Pneumonia ^e | 20.4 | 19.5 | 1.04 (0.54 to 2.03) | .90 |
| COVID-19 | 11.1 | 33.4 | 0.33 (0.17 to 0.65) | .001 |
| Anemia | 18.5 | 30.3 | 0.61 (0.33 to 1.12) | .11 |
| Neutropenia ^f | 26.4 | 66.5 | 0.40 (0.25 to 0.64) | <.001 |
| Cough | 14.3 | 30.8 | 0.47 (0.25 to 0.88) | .02 |
| Diarrhea | 15.3 | 63.7 | 0.24 (0.14 to 0.42) | <.001 |
| Pyrexia | 11.1 | 52.4 | 0.21 (0.11 to 0.40) | <.001 |
| Fatigue | 9.5 | 34.2 | 0.28 (0.14 to 0.55) | <.001 |
| Nausea | 9.8 | 38.3 | 0.26 (0.13 to 0.51) | <.001 |
| Vomiting | 5.8 | 29.6 | 0.19 (0.08 to 0.44) | <.001 |
| ALT increased | 2.8 | 33.6 | 0.08 (0.03 to 0.25) | <.001 |
| Weight decreased | 2.8 | 28.5 | 0.10 (0.03 to 0.29) | <.001 |

TEAE = treatment-emergent adverse event; IR = incidence rate; IRR = IR ratio

Incidence and Management Recommendations for Select BTK Inhibitor-Associated Cardiologic Adverse Events and Bleeding

| Adverse event | BTK inhibitor | Incidence Any grade, Grade ≥3 % | Management |
|---------------------|---------------|---------------------------------------|--|
| Atrial fibrillation | Ibrutinib | 16, 2-5 | Avoid stroke; anticoagulation Better symptom control: rate vs rhythm Cardiovascular and other comorbidity management |
| | Acalabrutinib | 6-9, 1-5 | |
| | Zanubrutinib | 3-6, ≤1 | |
| | Pirtobrutinib | 2.8, 1.2 | |
| Hypertension | Ibrutinib | 16-23, 8-12 | Correct predisposing factors Antihypertensive therapy |
| | Acalabrutinib | 7-9, 3-4 | |
| | Zanubrutinib | 14-17, 6-15 | |
| | Pirtobrutinib | 9.2, 2.3 | |
| Bleeding | Ibrutinib | 36-51, 3-4 | Minor bleeding: no intervention Major bleeding: <ul style="list-style-type: none"> • Consider treatment discontinuation • Platelet transfusions regardless of platelet counts |
| | Acalabrutinib | 36-51, 3 | |
| | Zanubrutinib | 36-45, 3 | |
| | Pirtobrutinib | — | |

Incidence and Management Recommendations for Select BTK Inhibitor-Associated Noncardiovascular Adverse Events

| Adverse event | BTK inhibitor | Incidence Any grade, Grade ≥3 % | Management |
|---------------|---------------|---------------------------------------|---|
| Neutropenia | Ibrutinib | 25-39, 13-31 | Growth factor support |
| | Acalabrutinib | 21-23, 13-19 | |
| | Zanubrutinib | 37-34, 15-19 | |
| | Pirtobrutinib | 25, 20.3 | |
| Diarrhea | Ibrutinib | 22-59, <1-4 | Symptomatic treatments and dose adjustments Dietary modifications, hydration, anti-diarrheal medications Probiotics |
| | Acalabrutinib | 18-39, 1-5 | |
| | Zanubrutinib | 14-18, <1-2 | |
| | Pirtobrutinib | 24.2, 0-9 | |
| Headache | Ibrutinib | 14-18, 1-2 | Moderate dose of caffeine or acetaminophen |
| | Acalabrutinib | 22-39, <1 | |
| | Zanubrutinib | 11-12, 0-1 | |
| | Pirtobrutinib | 13.1, 0.5 | |

BRUIN CLL-322: An Ongoing Phase III Trial of Pirtobrutinib and Venetoclax/Rituximab for Relapsed/Refractory CLL

Key Inclusion Criteria

- Confirmed CLL/SLL per iwCLL 2018³
- Previously treated CLL/SLL (including a covalent BTKi or covalent BTKi naïve [limited to 20% of total enrollment])
- Known 17p status
 - If 17p status is unknown, local or central FISH test results during screening can be used
- No prior venetoclax
- ≥18 years of age and ECOG 0-2

N=600

1:1

Randomization

Arm A (PVR)
Pirtobrutinib
+ Venetoclax
+ Rituximab

Pirtobrutinib, 200 mg oral, once daily from C1D1 - C28

Rituximab, IV, 375 mg/m² on C1D1
500 mg/m² on D1 of C2-C6

Venetoclax, oral, daily from C5 - C28: 400 mg
• Dose Ramp (5 weeks) from C4D1: 20-400 mg

Arm B (VR)
Venetoclax
+ Rituximab

Rituximab, IV, 375 mg/m² on C2D1
500 mg/m² on D1 of C3-C7

Venetoclax, oral, daily from C2 - C25: 400 mg
• Dose Ramp (5 weeks) from C1D1: 20-400 mg

Stratification factors

- 17p status (deleted/wildtype)
- Prior experience of BTKi (discontinuation due to PD or other vs no prior BTKi)

Each cycle is 28 days; C1 of Arm B is 35 days

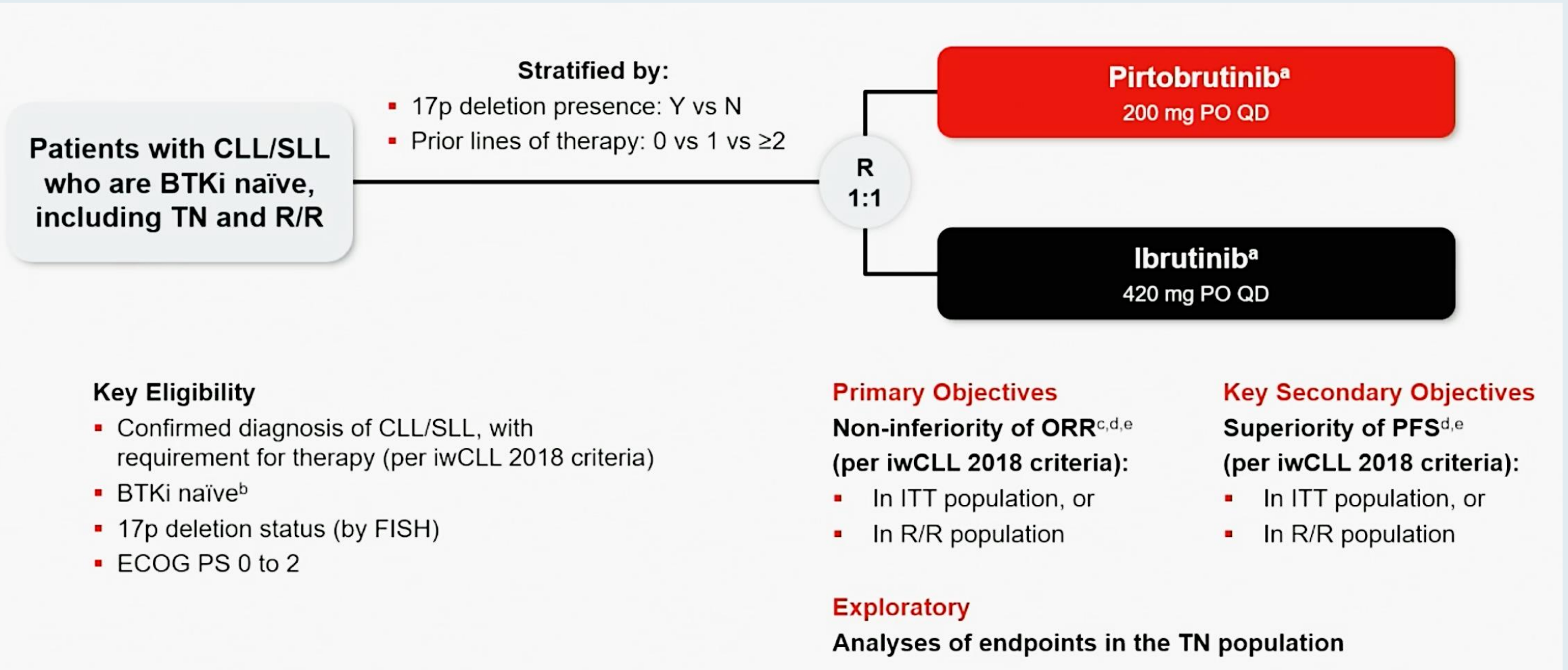
Primary endpoint: Progression-free survival per iwCLL 2018 by IRC

Pirtobrutinib vs Ibrutinib in Treatment-Naïve and Relapsed/Refractory CLL/SLL: Results From the First Randomized Phase III Study Comparing a Non-covalent and Covalent BTK Inhibitor

Jennifer A. Woyach¹, Lugui Qiu², Sebastian Grosicki³, Tomasz Wrobel⁴, Marcelo Capra⁵, Jaroslaw Czyz⁶, Shuhua Yi², Ki Seong Eom⁷, Anna Panovská⁸, Wojciech Jurczak⁹, Kamel Laribi¹⁰, Lutz Jacobasch¹¹, Ross Baker¹², Richy Agajanian¹³, Alejandro Berkovits¹⁴, Muhit Özcan¹⁵, Stéphane Lepretre¹⁶, Catherine C. Coombs¹⁷, Paula Cramer¹⁸, Katharine L. Lewis^{19,20}, Marisa Hill²¹, Katherine Bao²¹, Yuanyuan Bian²¹, Amy S. Ruppert²¹, Ching Ching Leow²¹, William G. Wierda²²

ASH 2025;Abstract 683.

BRUIN CLL-314 Study Design



TN = treatment naïve

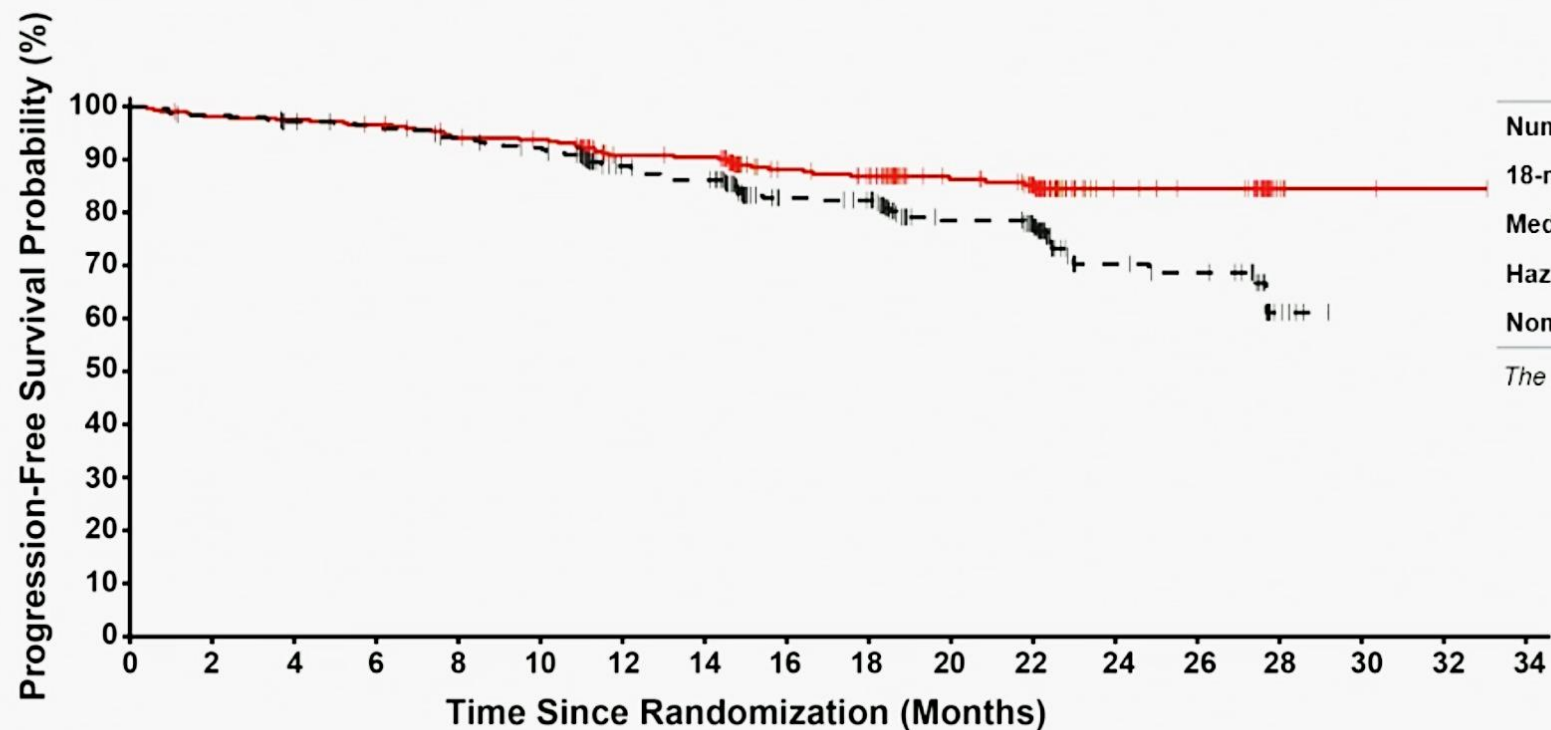
BRUIN CLL-314: Response Data

| | ITT Population | | TN Population | | R/R Population | |
|---|------------------------|--------------------|------------------------|--------------------|------------------------|--------------------|
| | Pirtobrutinib n=331 | Ibrutinib n=331 | Pirtobrutinib n=112 | Ibrutinib n=113 | Pirtobrutinib n=219 | Ibrutinib n=218 |
| ORR^a (PR or better) | | | | | | |
| % | 87.0 | 78.5 | 92.9 | 85.8 | 84.0 | 74.8 |
| 95% CI ^b | 82.90, 90.44 | 73.73, 82.85 | 86.41, 96.87 | 78.03, 91.68 | 78.48, 88.61 | 68.46, 80.39 |
| Nominal p-value ^c | 0.0035 | | 0.0886 | | 0.0175 | |
| ORR^a ratio | | | | | | |
| ORR ratio (95% CI) | 1.1080 (1.034, 1.187) | | 1.0797 (0.989, 1.179) | | 1.1233 (1.020, 1.237) | |
| p-value for NI ^d | <0.0001 | | - | | <0.0001 | |
| Best Overall Response^e, % | | | | | | |
| CR or CRi | 4.8 | 2.4 | 7.1 | 3.5 | 3.7 | 1.8 |
| PR or nPR | 82.2 | 76.1 | 85.7 | 82.3 | 80.4 | 72.9 |
| PR-L | 2.4 | 3.9 | 0.9 | 2.7 | 3.2 | 4.6 |
| SD | 5.4 | 10.9 | 2.7 | 4.4 | 6.8 | 14.2 |
| PD | 1.5 | 1.2 | 0 | 0 | 2.3 | 1.8 |
| ORR including PR-L | | | | | | |
| % | 89.4 | 82.5 | 93.8 | 88.5 | 87.2 | 79.4 |
| 95% CI ^b | 85.60, 92.52 | 77.95, 86.42 | 87.55, 97.45 | 81.13, 93.73 | 82.05, 91.33 | 73.37, 84.53 |
| Nominal p-value ^c | 0.0093 | | 0.1692 | | 0.0286 | |

ORR results presented are IRC-assessed

ORR = overall response rate; NI = noninferiority; PR-L = partial remission with lymphocytosis

BRUIN CLL-314: PFS in ITT Population



| | Pirtobrutinib (n=331) | Ibrutinib (n=331) |
|------------------------------|--------------------------|----------------------|
| Number of events, n (%) | 43 (13.0) | 69 (20.8) |
| 18-month PFS rate (95% CI) | 86.9 (82.4, 90.3) | 82.3 (77.3, 86.3) |
| Median follow-up, mo | 22.0 | 19.7 |
| Hazard ratio (95% CI) | 0.569 (0.388, 0.834) | |
| Nominal p-value ^a | 0.0034 | |

The PFS results presented are INV-assessed

Number at risk

| | | | | | | | | | | | | | | | | | | |
|---------------|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|----|----|---|---|---|---|
| Pirtobrutinib | 331 | 319 | 315 | 311 | 301 | 298 | 257 | 255 | 205 | 198 | 154 | 140 | 48 | 45 | 7 | 3 | 1 | 0 |
| Ibrutinib | 331 | 310 | 303 | 297 | 288 | 280 | 235 | 227 | 177 | 173 | 129 | 118 | 44 | 41 | 6 | 0 | 0 | 0 |

Pirtobrutinib reduced the risk of progression or death by 43%, with ibrutinib outcomes consistent with historical data

ITT = intent to treat

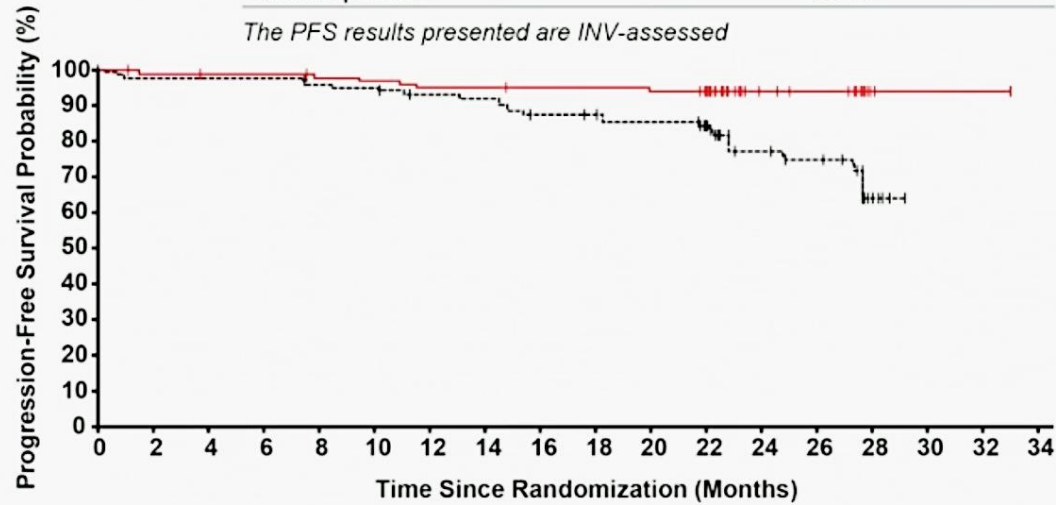


BRUIN CLL-314: PFS by Prior Treatment Status

TN population

| | Pirtobrutinib (n=112) | Ibrutinib (n=113) |
|------------------------------|--------------------------|----------------------|
| Number of events, n (%) | 6 (5.4) | 24 (21.2) |
| 18-month PFS rates (95% CI) | 95.3 (89.1, 98.0) | 87.6 (79.7, 92.6) |
| Median follow-up, mo | 22.5 | 22.4 |
| Hazard ratio (95% CI) | 0.239 (0.098, 0.586) | |
| Nominal p-value ^a | 0.0007 | |

The PFS results presented are INV-assessed



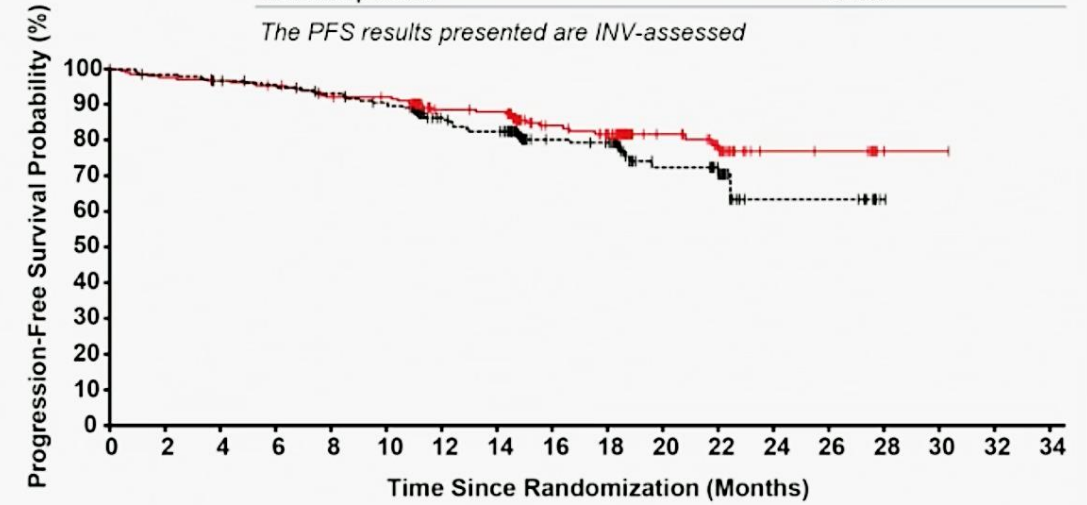
Number at risk

| | | | | | | | | | | | | | | | | | | |
|---------------|-----|-----|-----|-----|-----|-----|-----|-----|----|----|----|----|----|----|---|---|---|---|
| Pirtobrutinib | 112 | 107 | 106 | 106 | 104 | 103 | 100 | 100 | 99 | 99 | 98 | 94 | 35 | 33 | 4 | 1 | 1 | 0 |
| Ibrutinib | 113 | 105 | 105 | 105 | 102 | 101 | 97 | 96 | 90 | 89 | 86 | 81 | 32 | 29 | 5 | 0 | 0 | 0 |

R/R population

| | Pirtobrutinib (n=219) | Ibrutinib (n=218) |
|------------------------------|--------------------------|----------------------|
| Number of events, n (%) | 37 (16.9) | 45 (20.6) |
| 18-month PFS rate (95% CI) | 81.7 (75.1, 86.7) | 79.2 (72.3, 84.6) |
| Median follow-up, mo | 18.4 | 15.8 |
| Hazard ratio (95% CI) | 0.729 (0.471, 1.128) | |
| Nominal p-value ^a | 0.1563 | |

The PFS results presented are INV-assessed



Number at risk

| | | | | | | | | | | | | | | | | | | |
|---------------|-----|-----|-----|-----|-----|-----|-----|-----|-----|----|----|----|----|----|---|---|---|---|
| Pirtobrutinib | 219 | 212 | 209 | 205 | 197 | 195 | 157 | 155 | 106 | 99 | 56 | 46 | 13 | 12 | 3 | 2 | 0 | 0 |
| Ibrutinib | 218 | 205 | 198 | 192 | 186 | 179 | 138 | 131 | 87 | 84 | 43 | 37 | 12 | 12 | 1 | 0 | 0 | 0 |

Pirtobrutinib reduced the risk of progression or death by 76% in the TN population, the subgroup with the longest follow-up

BRUIN CLL-314: Safety Profile

| Preferred Term ≥10% of Participants in Either Arm | Pirtobrutinib n=330 | | Ibrutinib n=325 | |
|--|------------------------|-------------------|--------------------|-------------------|
| | Any Grade n (%) | Grade ≥3 n (%) | Any Grade n (%) | Grade ≥3 n (%) |
| Subjects with ≥1 TEAE | 320 (97.0) | 181 (54.8) | 318 (97.8) | 174 (53.5) |
| Neutropenia | 75 (22.7) | 57 (17.3) | 58 (17.8) | 43 (13.2) |
| Upper respiratory tract infection | 59 (17.9) | 2 (0.6) | 63 (19.4) | 0 (0) |
| Anemia | 50 (15.2) | 19 (5.8) | 46 (14.2) | 12 (3.7) |
| Pneumonia | 45 (13.6) | 21 (6.4) | 49 (15.1) | 28 (8.6) |
| Diarrhea | 44 (13.3) | 1 (0.3) | 62 (19.1) | 4 (1.2) |
| COVID-19 | 40 (12.1) | 4 (1.2) | 33 (10.2) | 5 (1.5) |
| Hypertension | 35 (10.6) | 11 (3.3) | 49 (15.1) | 16 (4.9) |
| Contusion | 33 (10.0) | 0 (0) | 30 (9.2) | 0 (0) |
| Arthralgia | 26 (7.9) | 0 (0) | 41 (12.6) | 0 (0) |
| Thrombocytopenia | 26 (7.9) | 9 (2.7) | 37 (11.4) | 10 (3.1) |
| Urinary tract infection | 26 (7.9) | 3 (0.9) | 40 (12.3) | 3 (0.9) |
| Atrial fibrillation | 8 (2.4) | 3 (0.9) | 41 (12.6) | 12 (3.7) |
| Dose modifications due to TEAEs | | | | |
| Reductions | | 26 (7.9) | | 59 (18.2) |
| Discontinuations | | 31 (9.4) | | 35 (10.8) |

Median time on treatment was 20.5 months with pirtobrutinib and 19.3 months with ibrutinib;
1 patient developed Richter Transformation (RT) on pirtobrutinib; 4 patients developed RT on ibrutinib

Pirtobrutinib was well-tolerated with fewer dose reductions and discontinuations due to TEAEs than ibrutinib

BRUIN CLL-314: Adverse Events of Special Interest (AESI)

| ≥10% of Participants in Either Arm | Pirtobrutinib n=330 | | Ibrutinib n=325 | |
|---|------------------------|-------------------|--------------------|-------------------|
| | Any Grade n (%) | Grade ≥3 n (%) | Any Grade n (%) | Grade ≥3 n (%) |
| Subjects with ≥1 AESI | 288 (87.3) | 127 (38.5) | 288 (88.6) | 117 (36.0) |
| Infections^a | 226 (68.5) | 56 (17.0) | 241 (74.2) | 54 (16.6) |
| Infection without COVID-19 | 214 (64.8) | 53 (16.1) | 234 (72.0) | 49 (15.1) |
| Bleeding | 115 (34.8) | 11 (3.3) | 118 (36.3) | 9 (2.8) |
| Hemorrhage ^b | 78 (23.6) | 11 (3.3) | 81 (24.9) | 9 (2.8) |
| Bruising ^c | 45 (13.6) | 0 (0) | 39 (12.0) | 0 (0) |
| Petechiae and purpura | 17 (5.2) | 0 (0) | 25 (7.7) | 0 (0) |
| Neutropenia^d | 103 (31.2) | 83 (25.2) | 76 (23.4) | 57 (17.5) |
| Anemia^e | 51 (15.5) | 20 (6.1) | 51 (15.7) | 12 (3.7) |
| Thrombocytopenia^f | 39 (11.8) | 12 (3.6) | 57 (17.5) | 13 (4.0) |
| Atrial fibrillation and atrial flutter | 8 (2.4) | 3 (0.9) | 44 (13.5) | 13 (4.0) |
| ≥75 years old ^g | 3 (4.5) | 1 (1.5) | 15 (21.4) | 5 (7.1) |

AEs were mostly low-grade and consistent with prior pirtobrutinib studies

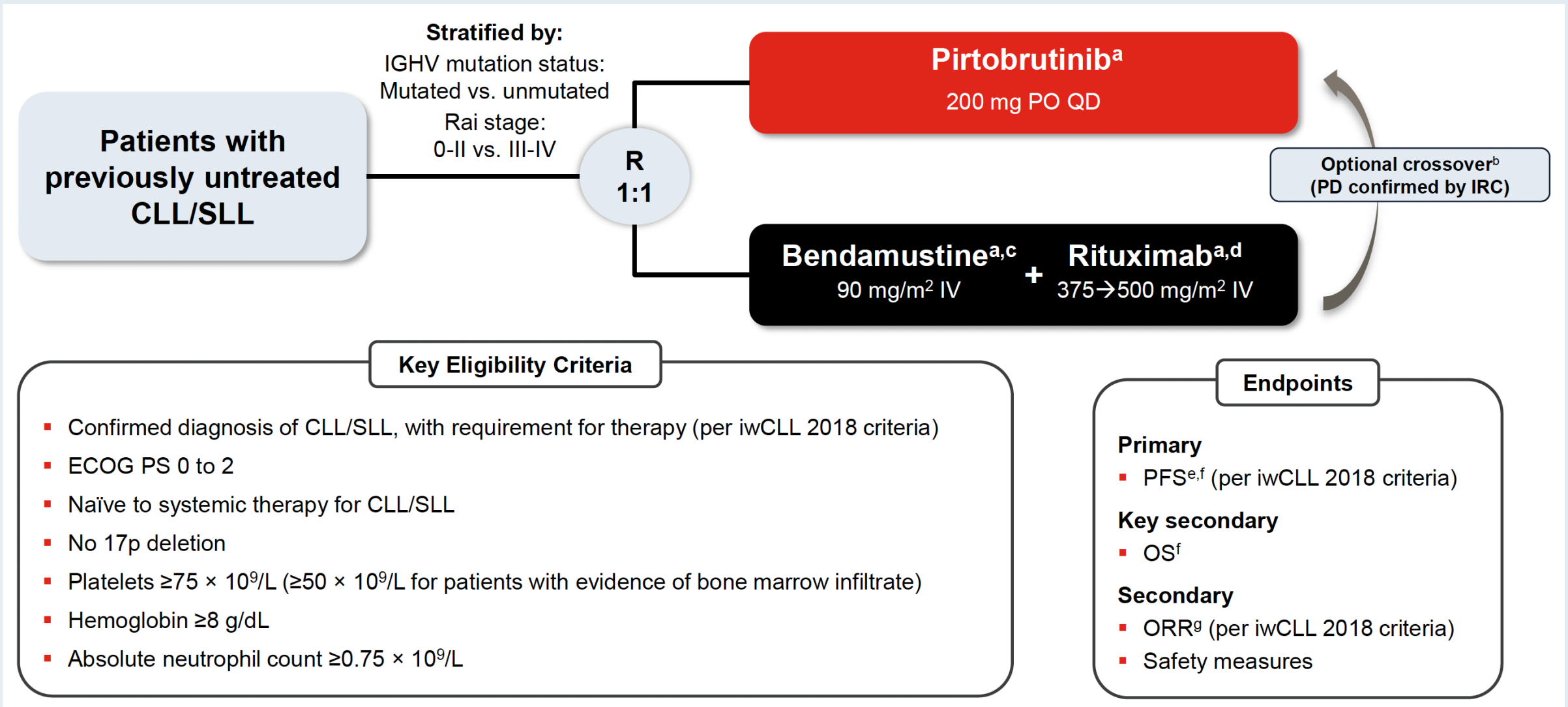
Incidence of atrial fibrillation/flutter was substantially lower with pirtobrutinib vs ibrutinib, particularly among older patients

Pirtobrutinib vs Bendamustine Plus Rituximab (BendaR) in Patients With CLL/SLL: First Results From a Randomized Phase III Study Examining a Non-Covalent BTK Inhibitor in Untreated Patients

Wojciech Jurczak¹, Michal Kwiatek², Jaroslaw Czyz³, Ederson Roberto de Mattos⁴, Ki-Seong Eom⁵, Alexander Egle⁶, Anna Panovská⁷, Zhanet Grudeva-Popova⁸, Hsuan-Jen Shih⁹, Luis Felipe Casado Montero¹⁰, Paolo Sportoletti¹¹, Vu Minh Hua¹², James T. D'Olimpio¹³, Shinsuke Iida¹⁴, Rodrigo Ito¹⁵, Katherine Bao¹⁵, Anne Fink¹⁵, Weiji Su¹⁵, Amy S. Ruppert¹⁵, Alejandro Levy¹⁵, Tomasz Wrobel¹⁶

ASH 2025;Abstract LBA3.

BRUIN CLL-313 Study Design



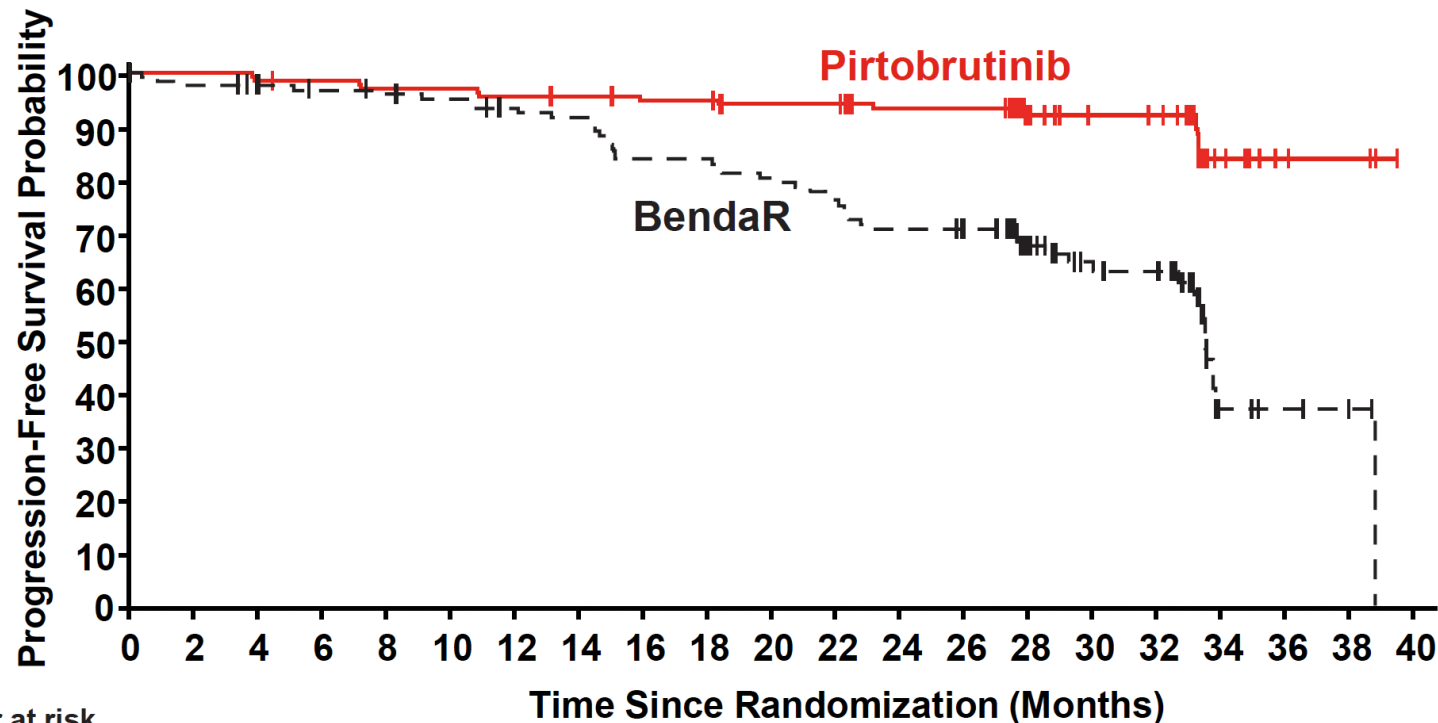
BRUIN CLL-313: Response Data

| | IRC-Assessed ORR | | INV-Assessed ORR | |
|---|--------------------------|---------------------|--------------------------|---------------------|
| | Pirtobrutinib (n=141) | BendaR (n=141) | Pirtobrutinib (n=141) | BendaR (n=141) |
| ORR^a (PR or better) | | | | |
| n (%) | 133 (94.3) | 114 (80.9) | 133 (94.3) | 116 (82.3) |
| 95% CI ^b | 89.1, 97.5 | 73.4, 87.0 | 89.1, 97.5 | 75.0, 88.2 |
| Best overall response^c, n (%) | | | | |
| CR or CRi | 19 (13.5) | 29 (20.6) | 8 (5.7) | 29 (20.6) |
| PR or nPR | 114 (80.9) | 85 (60.3) | 125 (88.7) | 87 (61.7) |
| PR-L | 0 | 0 | 2 (1.4) | 0 |
| SD | 3 (2.1) | 10 (7.1) | 1 (0.7) | 6 (4.3) |
| PD | 2 (1.4) | 1 (0.7) | 2 (1.4) | 3 (2.1) |
| Unknown/not applicable ^d | 3 (2.1) | 16 (11.3) | 3 (2.1) | 16 (11.3) |

ORR with pirtobrutinib was >90% by both IRC and INV, and higher than with BendaR

ORR = overall response rate; PR = partial response

BRUIN CLL-313: PFS Outcomes



| Number at risk | 0 | 2 | 4 | 6 | 8 | 10 | 12 | 14 | 16 | 18 | 20 | 22 | 24 | 26 | 28 | 30 | 32 | 34 | 36 | 38 | 40 |
|----------------------|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|----|----|----|----|----|----|----|
| Pirtobrutinib | 141 | 138 | 136 | 135 | 133 | 133 | 131 | 130 | 128 | 128 | 124 | 124 | 119 | 119 | 67 | 56 | 55 | 11 | 5 | 4 | 0 |
| BendaR | 141 | 122 | 120 | 116 | 114 | 111 | 107 | 105 | 96 | 96 | 92 | 87 | 81 | 77 | 50 | 38 | 36 | 6 | 4 | 3 | 0 |

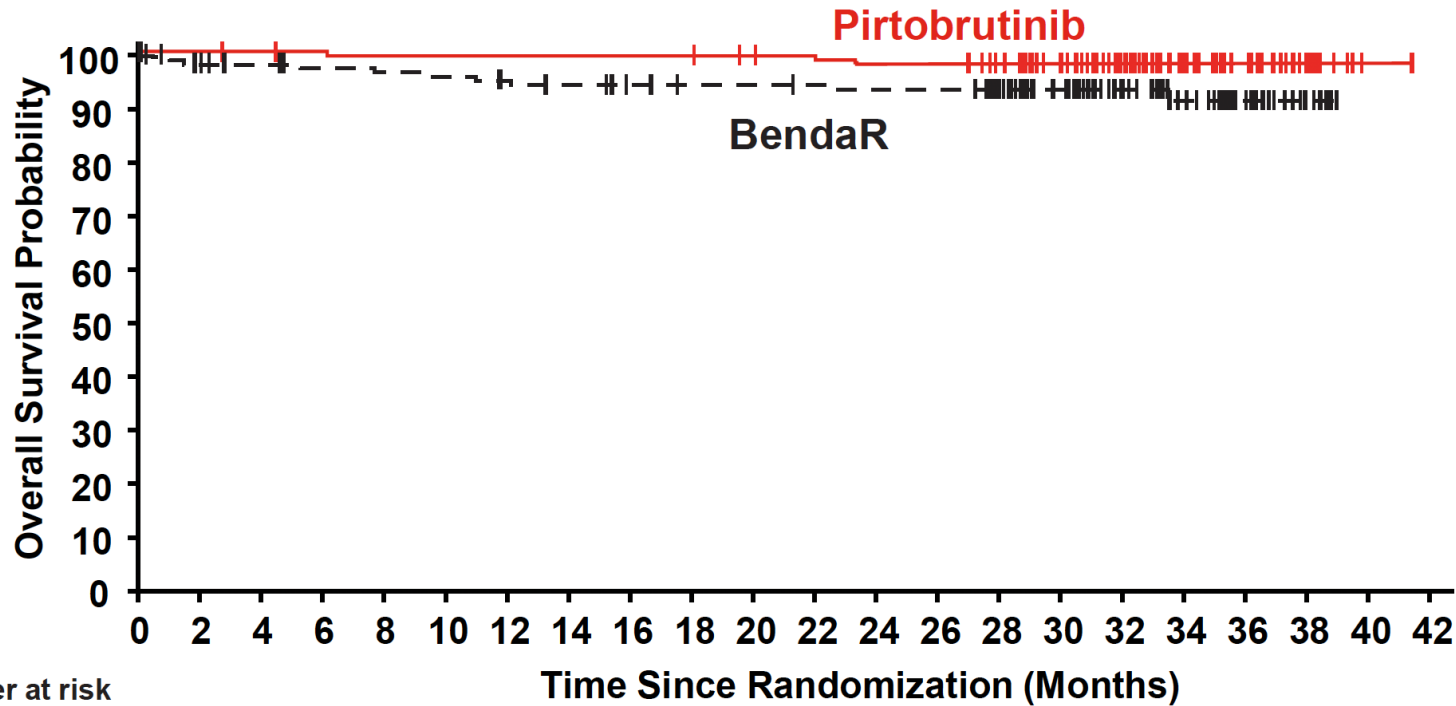
| | Pirtobrutinib (n=141) | BendaR (n=141) |
|-----------------------------|----------------------------------|-----------------------------|
| Number of events, n (%) | 13 (9.2) | 48 (34.0) |
| 24-month PFS rate, (95% CI) | 93.4 (87.6, 96.5) | 70.7 (61.5, 78.1) |
| Median follow-up, months | 28.1 | 28.3 |
| Hazard ratio (95% CI) | 0.20 (0.11, 0.37) | |
| p-value ^a | <0.0001^a | |

The PFS results presented are IRC assessed

Pirtobrutinib demonstrated a statistically significant and clinically meaningful PFS improvement, with an 80% reduction in risk of PD or death compared with BendaR



BRUIN CLL-313: OS Outcomes



| Number at risk | 0 | 2 | 4 | 6 | 8 | 10 | 12 | 14 | 16 | 18 | 20 | 22 | 24 | 26 | 28 | 30 | 32 | 34 | 36 | 38 | 40 | 42 |
|----------------------|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|----|----|----|----|----|----|
| Pirtobrutinib | 141 | 140 | 139 | 138 | 137 | 137 | 137 | 137 | 137 | 137 | 135 | 134 | 132 | 132 | 125 | 106 | 80 | 52 | 39 | 15 | 1 | 0 |
| BendaR | 141 | 130 | 127 | 124 | 123 | 122 | 119 | 117 | 114 | 113 | 113 | 112 | 111 | 111 | 100 | 82 | 61 | 40 | 23 | 9 | 0 | 0 |

| | Pirtobrutinib n=141 | BendaR n=141 |
|----------------------------|--------------------------------|-----------------------------|
| Number of events, n (%) | 3 (2.1) | 10 (7.1) |
| 24-month OS rate, (95% CI) | 97.8 (93.3, 99.3) | 93.0 (87.0, 96.3) |
| Median follow-up, months | 32.7 | 31.7 |
| Hazard ratio (95% CI) | 0.26 (0.07, 0.93) | |
| p-value | 0.0261 ^a | |

Effective crossover rate^b:
52.9% (18/34)

OS data were immature, but trended in favor of pirtobrutinib, despite a high effective crossover rate

BRUIN CLL-313: Safety Profile

| Preferred Term ≥15% of Participants in Either Arm | Pirtobrutinib (n=140) | | BendaR (n=132) | | EAIR per 100 Person-Years | | |
|--|--------------------------|-------------------|--------------------|-------------------|---|--|-------------------------------------|
| | Any Grade n (%) | Grade ≥3 n (%) | Any Grade n (%) | Grade ≥3 n (%) | Pirtobrutinib Any Grade EAIR ^a | BendaR Any Grade EAIR ^a | EAIR Ratio (95% CI) ^b |
| Patients with ≥1 TEAE | 131 (93.6) | 56 (40.0) | 117 (88.6) | 89 (67.4) | 196.7 | 844.6 | 0.23 (0.18, 0.30) |
| Neutropenia | 17 (12.1) | 10 (7.1) | 51 (38.6) | 46 (34.8) | 5.2 | 110.0 | 0.05 (0.03, 0.08) |
| COVID-19 | 30 (21.4) | 1 (0.7) | 12 (9.1) | 2 (1.5) | 9.9 | 20.9 | 0.47 (0.24, 0.92) |
| Pyrexia | 12 (8.6) | 0 | 25 (18.9) | 0 | 3.5 | 49.1 | 0.07 (0.04, 0.14) |
| Upper respiratory tract infection | 25 (17.9) | 1 (0.7) | 9 (6.8) | 0 | 7.7 | 15.7 | 0.49 (0.23, 1.05) |
| Anemia | 13 (9.3) | 6 (4.3) | 21 (15.9) | 10 (7.6) | 3.8 | 37.7 | 0.10 (0.05, 0.20) |
| Nausea | 3 (2.1) | 0 | 31 (23.5) | 1 (0.8) | 0.8 | 65.1 | 0.01 (0.00, 0.04) |
| Infusion-related reaction | 0 | 0 | 20 (15.2) | 4 (3.0) | 0 | 39.0 | NA |

Median time on pirtobrutinib treatment was 32.3 months and on BendaR treatment was 5.6 months

Richter's transformation occurred in 1 patient treated with BendaR

Exposure adjusted TEAEs were lower with pirtobrutinib

Discontinuation due to TEAE, including deaths, occurred less frequently with pirtobrutinib (4.3%) vs. BendaR (15.4%)

Dose reduction due to TEAE also occurred less frequently with pirtobrutinib than BendaR (3.6% vs. 31.1%)

EAIR = exposure-adjusted incidence rate

BRUIN CLL-313: AEs of Special Interest

| | Pirtobrutinib (n=140) | | BendaR (n=132) | | EAIR per 100 Person-Years | | |
|---|-----------------------|----------------|-----------------|----------------|---|------------------------------------|----------------------------------|
| | Any Grade n (%) | Grade ≥3 n (%) | Any Grade n (%) | Grade ≥3 n (%) | Pirtobrutinib Any Grade EAIR ^g | BendaR Any Grade EAIR ^g | EAIR Ratio (95% CI) ^h |
| Infection^a | 80 (57.1) | 19 (13.6) | 44 (33.3) | 11 (8.3) | 38.3 | 89.7 | 0.43 (0.30, 0.62) |
| Infection without COVID-19 | 72 (51.4) | 19 (13.6) | 38 (28.8) | 9 (6.8) | 30.9 | 74.9 | 0.41 (0.28, 0.61) |
| Bleeding^b | 36 (25.7) | 1 (0.7) | 2 (1.5) | 0 (0) | 12.5 | 3.3 | 3.73 (0.90, 15.50) |
| Hemorrhage | 17 (12.1) | 1 (0.7) | 2 (1.5) | 0 (0) | 5.2 | 3.3 | 1.55 (0.36, 6.69) |
| Bruising | 16 (11.4) | 0 (0) | 0 (0) | 0 (0) | 4.8 | 0 | NE |
| Petechiae and purpura | 8 (5.7) | 0 (0) | 0 (0) | 0 (0) | 2.3 | 0 | NE |
| Neutropenia^c | 21 (15.0) | 13 (9.3) | 68 (51.5) | 60 (45.5) | 6.5 | 169.5 | 0.04 (0.02, 0.06) |
| Anemia^d | 14 (10.0) | 6 (4.3) | 21 (15.9) | 10 (7.6) | 4.1 | 37.7 | 0.11 (0.06, 0.21) |
| Thrombocytopenia^e | 12 (8.6) | 4 (2.9) | 23 (17.4) | 9 (6.8) | 3.5 | 43.1 | 0.08 (0.04, 0.16) |
| Atrial fibrillation and atrial flutter | 2 (1.4) | 1 (0.7) | 2 (1.5) | 1 (0.8) | 0.5 | 3.3 | 0.17 (0.02, 1.17) |
| ≥75 years old ^f | 1 (5.0) | 0 | 1 (4.3) | 0 | 2.2 | 10.0 | 0.22 (0.01, 3.46) |
| Hypertension | 11 (7.9) | 4 (2.9) | 6 (4.5) | 4 (3.0) | 3.2 | 10.2 | 0.31 (0.11, 0.84) |

Incidence of atrial fibrillation/flutter remains low in older patients aged ≥75 years (5.0% with pirtobrutinib and 4.3% with BendaR)

Management of Double-Refractory CLL

Introduction: Sequencing of Treatment for CLL

Module 1: Clinician Survey Results

Module 2: Noncovalent BTK Inhibitor Pirtobrutinib








Module 3: Clinician Survey Results

Module 4: CAR T-Cell Therapy

Module 5: Clinician Survey Results

Module 6: Bispecific Antibodies and Promising Investigational Strategies

Based on current clinical trial data and your personal experience, how would you compare the global efficacy and tolerability/toxicity of pirtobrutinib to that of ibrutinib, acalabrutinib and zanubrutinib for patients with relapsed/refractory CLL?

| | Efficacy | Tolerability/toxicity |
|---|--|--------------------------------------|
|  Dr Awan | About the same | Pirtobrutinib has the least toxicity |
|  Dr Coombs | There are not enough available data at this time | Pirtobrutinib has the least toxicity |
|  Dr Davids | About the same | Pirtobrutinib has the least toxicity |
|  Dr Fakhri | There are not enough available data at this time | Pirtobrutinib has the least toxicity |
|  Dr Lamanna | About the same | Pirtobrutinib has the least toxicity |
|  Dr Sharman | There are not enough available data at this time | Pirtobrutinib has the least toxicity |
|  Dr Woyach | There are not enough available data at this time | Pirtobrutinib has the least toxicity |

An older patient with newly diagnosed CLL and a significant history of atrial fibrillation has come to you for a second opinion after the recommendation of first-line treatment with pirtobrutinib. How would you respond?



Dr Awan

I do not agree with the recommendation, but it is a valid option



Dr Coombs

I do not agree with the recommendation, but it is a valid option



Dr Davids

I do not agree with the recommendation, but it is a valid option



Dr Fakhri

I do not agree with the recommendation, but it is a valid option



Dr Lamanna

I do not agree with this recommendation



Dr Sharman

I do not agree with this recommendation



Dr Woyach

I agree with this recommendation

Management of Double-Refractory CLL

Introduction: Sequencing of Treatment for CLL

Module 1: Clinician Survey Results

Module 2: Noncovalent BTK Inhibitor Pirtobrutinib

Module 3: Clinician Survey Results

Module 4: CAR T-Cell Therapy

Module 5: Clinician Survey Results

Module 6: Bispecific Antibodies and Promising Investigational Strategies

Key Datasets

- Siddiqi T et al. **Lisocabtagene maraleucel (liso-cel) in R/R CLL/SLL: 24-month median follow-up of TRANSCEND CLL 004.** ASH 2023;Abstract 330.
- Wierda WG et al. **Lisocabtagene maraleucel (liso-cel) combined with ibrutinib (ibr) for patients (pts) with relapsed or refractory (R/R) chronic lymphocytic leukemia (CLL)/small lymphocytic lymphoma (SLL): Primary results from the open-label, phase 1/2 Transcend CLL 004 study.** ASH 2024;Abstract 887.

Lisocabtagene Maraleucel in Relapsed or Refractory Chronic Lymphocytic Leukemia/Small Lymphocytic Lymphoma: 24-Month Median Follow-up of TRANSCEND CLL 004

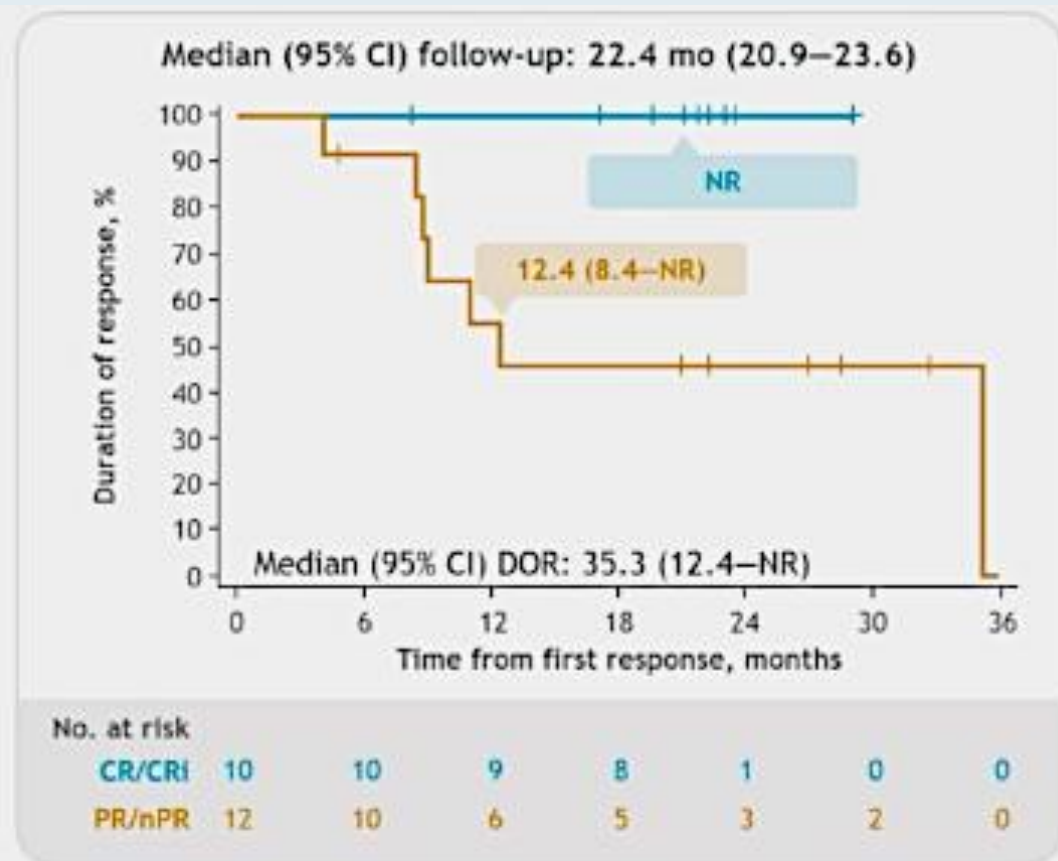
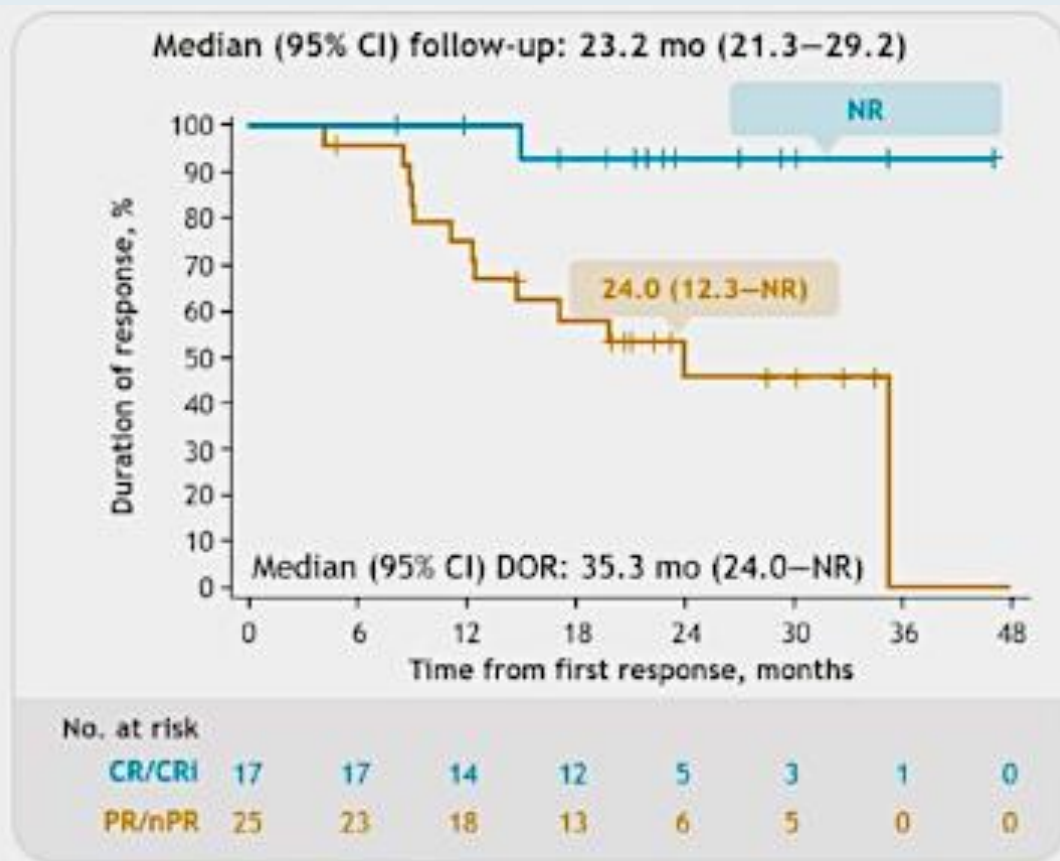
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TRANSCEND CLL 004: Duration of Response by Best Overall Response

Full Study Population at DL2 (n = 88)

PEAS (BTKi progression/venetoclax failure subset)
at DL2 (n = 50)



Data on KM curves are expressed as median (95% CI, if available). DOR, duration of response; NR, not reached.

FDA Grants Accelerated Approval to Lisocabtagene Maraleucel for Relapsed/Refractory (R/R) CLL or SLL

Press Release: March 14, 2024

“... the US Food and Drug Administration (FDA) has granted accelerated approval of lisocabtagene maraleucel (liso-cel), a CD19-directed chimeric antigen receptor (CAR) T cell therapy, for the treatment of adult patients with relapsed or refractory chronic lymphocytic leukemia (CLL) or small lymphocytic lymphoma (SLL) who have received at least two prior lines of therapy, including a Bruton tyrosine kinase (BTK) inhibitor and a B-cell lymphoma 2 (BCL-2) inhibitor. This indication is approved under accelerated approval based on response rate and duration of response. Continued approval for this indication may be contingent upon verification and description of clinical benefit in confirmatory trial(s). In R/R CLL or SLL, liso-cel is delivered through a treatment process which culminates in a one-time infusion with a single dose containing 90 to 110 x 10⁶ CAR-positive viable T cells.”

Accelerated approval was based on results from the Phase I/II open-label, single-arm TRANSCEND CLL 004 study for patients with R/R CLL or SLL.

Lisocabtagene Maraleucel Combined with Ibrutinib for Patients with Relapsed or Refractory Chronic Lymphocytic Leukemia/Small Lymphocytic Lymphoma: Primary Results from the Open-label, Phase 1/2 TRANSCEND CLL 004 Study

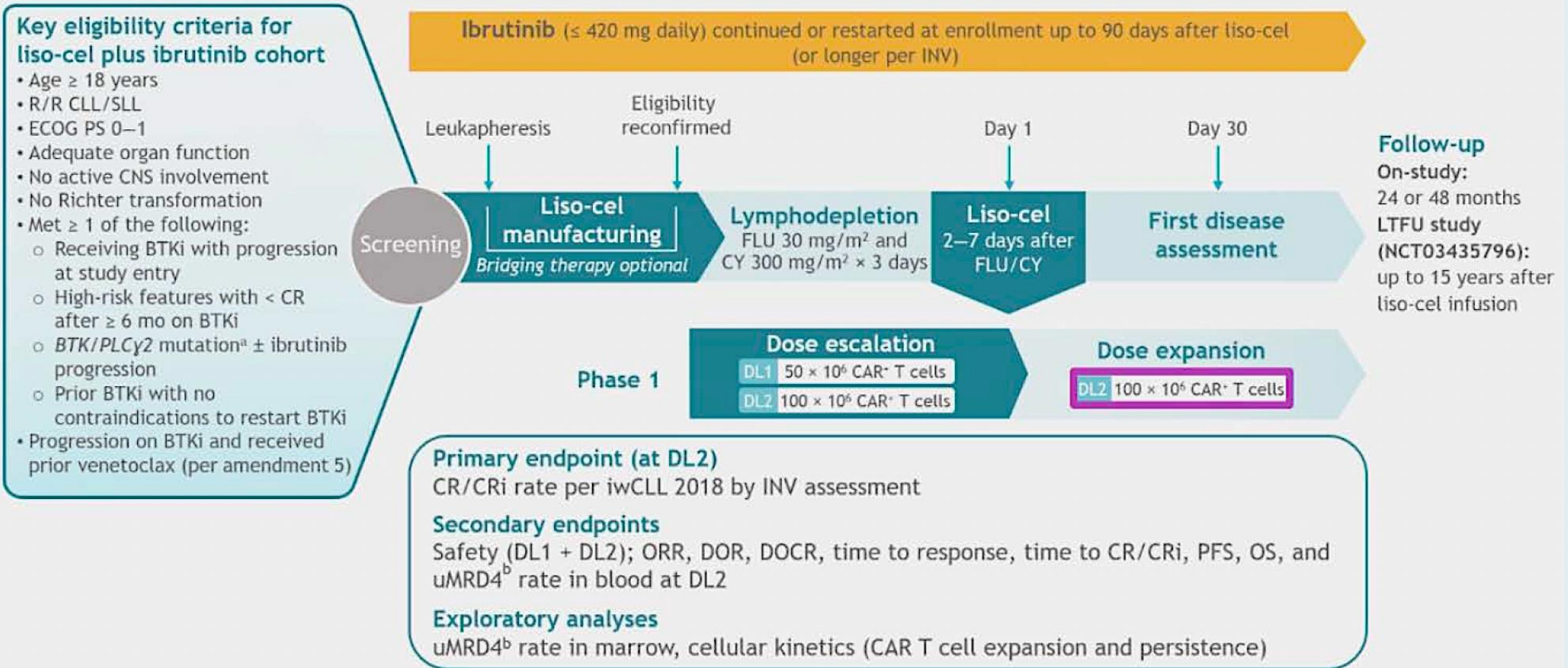
William G. Wierda, MD, PhD,¹ Kathleen Dorritie, MD,² Jordan Gauthier, MD, MSc,³ Rajneesh Nath, MD,⁴ Thomas Kipps, MD, PhD,⁵ Peter A. Riedell, MD,⁶ Herbert A. Eradat, MD,⁷ Saad S. Kenderian, MB, ChB,⁸ Mohamed A. Kharfan-Dabaja, MD, MBA,⁹ Nirav N. Shah, MD,¹⁰ Scott R. Solomon, MD,¹¹ Daniel A. Ermann, MD,¹² Jon Arnason, MD,¹³ Abhinav Deol, MD,¹⁴ Tatyana Feldman, MD,¹⁵ Charalambos Andreadis, MD, MS,¹⁶ Monalisa Ghosh, MD,¹⁷ Shuo Ma, MD, PhD,¹⁸ Stephen J. Schuster, MD,¹⁹ Usama Gergis, MD, MBA,²⁰ Julie M. Vose, MD, MBA,²¹ Jacob Soumerai, MD,²² Koen van Besien, MD, PhD,^{23*} Sherilyn A. Tuazon, MD,²⁴ Serena K. Perna, MD,²⁵ San-San Ou, MS,²⁴ Neha Rane, MD,²⁵ Eniko Papp, PhD,²⁴ Yizhe Chen, PhD,²⁵ Tanya Siddiqi, MD, MBBS²⁶

¹The University of Texas MD Anderson Cancer Center, Houston, TX, USA; ²UPMC Hillman Cancer Center, University of Pittsburgh, Pittsburgh, PA, USA; ³Fred Hutchinson Cancer Center, Seattle, WA, USA; ⁴Banner MD Anderson Cancer Center, Gilbert, AZ, USA; ⁵Moore's UCSD Cancer Center, San Diego, CA, USA; ⁶David and Etta Jonas Center for Cellular Therapy, University of Chicago, Chicago, IL, USA; ⁷University of California, Los Angeles, Santa Monica Cancer Center, Santa Monica, CA, USA; ⁸Mayo Clinic, Rochester, MN, USA; ⁹Mayo Clinic Comprehensive Cancer Center, Jacksonville, FL, USA; ¹⁰Medical College of Wisconsin, Milwaukee, WI, USA; ¹¹Northside Hospital Cancer Institute, Atlanta, GA, USA; ¹²Huntsman Cancer Institute, University of Utah, Salt Lake City, UT, USA; ¹³Beth Israel Deaconess Medical Center, Boston, MA, USA; ¹⁴Barbara Ann Karmanos Cancer Institute, Wayne State University, Detroit, MI, USA; ¹⁵John Theurer Cancer Center at Hackensack Meridian Health, HMH School of Medicine, Hackensack, NJ, USA; ¹⁶University of California, San Francisco, San Francisco, CA, USA; ¹⁷University of Michigan Health System, Ann Arbor, MI, USA; ¹⁸Robert H. Lurie Comprehensive Cancer Center of Northwestern University, Chicago, IL, USA; ¹⁹Lymphoma Program, Abramson Cancer Center, University of Pennsylvania, Philadelphia, PA, USA; ²⁰Thomas Jefferson University, Philadelphia, PA, USA; ²¹University of Nebraska Medical Center, Omaha, NE, USA; ²²Center for Lymphoma, Massachusetts General Hospital Cancer Center, Boston, MA, USA; ²³Weill Cornell Medical College, New York, NY, USA; ²⁴Bristol Myers Squibb, Seattle, WA, USA; ²⁵Bristol Myers Squibb, Princeton, NJ, USA; ²⁶City of Hope National Medical Center, Duarte, CA, USA

*Affiliation at the time the research was conducted

ASH 2024, Presentation 887

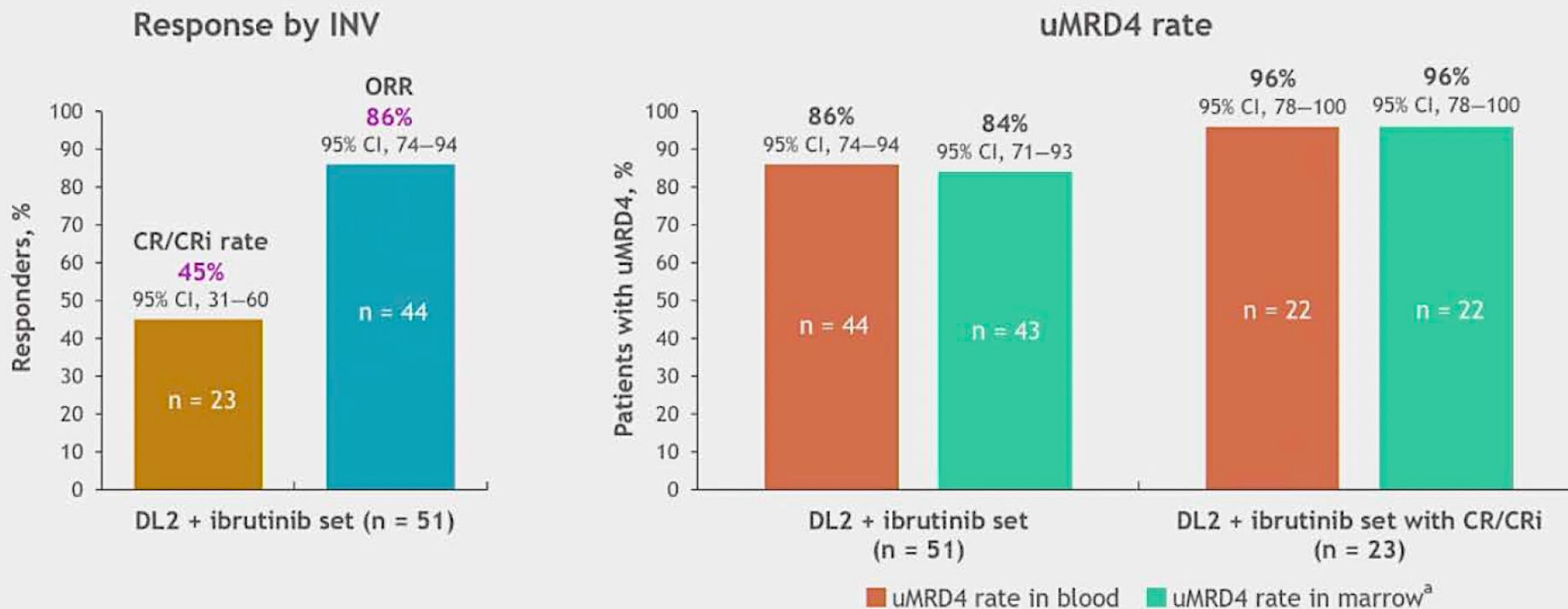
TRANSCEND CLL 004: Lisocabtagene Maraleucel (Liso-cel) and Ibrutinib Combination Cohort



^aPer local laboratory assessment; ^bMRD was assessed by next-generation sequencing using a clonoSEQ assay. Undetectable MRD was defined as < 1 CLL cell per 10,000 leukocytes at ≥ 1 time point after infusion (uMRD^b). CY, cyclophosphamide; DOR, duration of response; DOCR, duration of continued CR after initial CR; FLU, fludarabine; INV, investigator; iwCLL, International Workshop on Chronic Lymphocytic Leukemia; LTFU, long-term follow-up; uMRD^b, undetectable minimal residual disease at < 1 in 10⁻⁴ leukocytes.

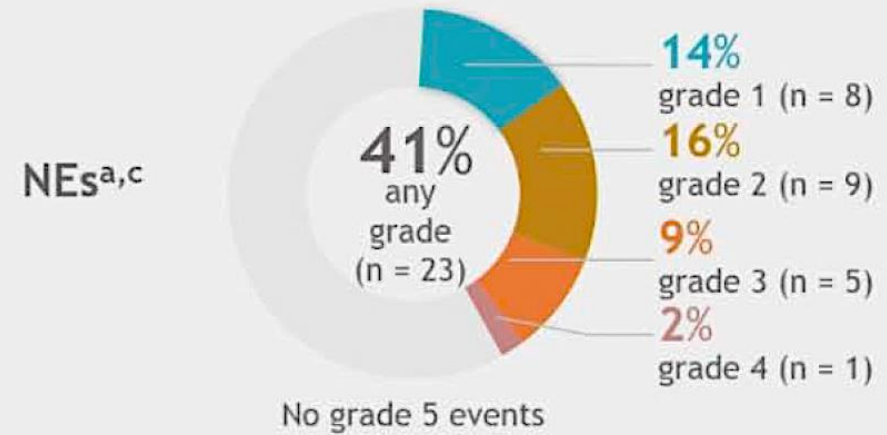
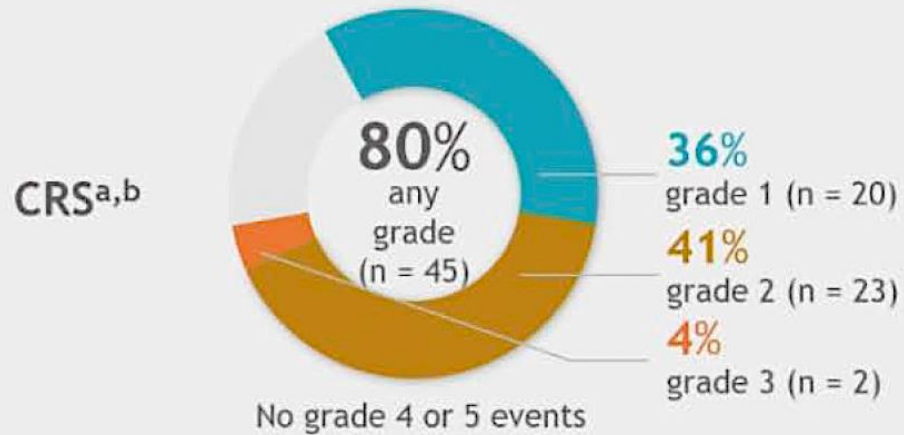
TRANSCEND CLL 004: Efficacy Outcomes with Liso-cel and Ibrutinib

- Median (IQR) on-study follow-up (including LTFU): 24.8 months (14.2–34.6)
- Median (range) time to first response: 1 month (0.9–6.0)
- Median (range) time to first CR/CRi: 3 months (0.9–12.1)



CR = complete response; CRi = CR with incomplete marrow recovery; ORR = overall response rate

TRANSCEND CLL 004: Incidence of Cytokine Release Syndrome (CRS) and Neurological Adverse Events (NEs) with Liso-cel and Ibrutinib



| | Total combination-treated set (n = 56) | | Total combination-treated set (n = 56) |
|--|--|--|--|
| Median (range) days to CRS onset | 7 (1–14) | Median (range) days to NE onset | 8 (1–15) |
| Median (range) days to CRS resolution | 5 (2–18) | Median (range) days to NE resolution | 8 (1–362) |
| Received tocilizumab and/or corticosteroids for CRS and/or NE, n (%) | 33 (59) | Received tocilizumab and/or corticosteroids for CRS and/or NE, n (%) | 33 (59) |

^aSummed percentages for grouped grades within each graph may not equal the any-grade percentage due to rounding; ^bCRS was graded based on Lee 2014 criteria; ^cNEs were defined as -INV-identified neurological AEs related to liso-cel.

Management of Double-Refractory CLL

Introduction: Sequencing of Treatment for CLL

Module 1: Clinician Survey Results

Module 2: Noncovalent BTK Inhibitor Pirtobrutinib

Module 3: Clinician Survey Results

Module 4: CAR T-Cell Therapy

Module 5: Clinician Survey Results

Module 6: Bispecific Antibodies and Promising Investigational Strategies

At what point in the treatment course are you referring patients with multiregimen-relapsed CLL for consultation regarding chimeric antigen receptor (CAR) T-cell therapy?



Dr Awan

At second relapse



Dr Coombs

After third relapse



Dr Davids

After third relapse



Dr Fakhri

At third relapse



Dr Lamanna

After third relapse



Dr Sharman








At second relapse



Dr Woyach

At second relapse

Approximately how many patients with CLL in your practice have received CAR T-cell therapy on or off protocol?

| | |
|--|------------------------|
|  Dr Awan | >10 patients |
|  Dr Coombs | 0 patients |
|  Dr Davids | 10 patients |
|  Dr Fakhri | 20 patients |
|  Dr Lamanna | 5 patients |
|  Dr Sharman | 4 patients |
|  Dr Woyach | 10 patients |

Management of Double-Refractory CLL

Introduction: Sequencing of Treatment for CLL

Module 1: Clinician Survey Results

Module 2: Noncovalent BTK Inhibitor Pirtobrutinib

Module 3: Clinician Survey Results

Module 4: CAR T-Cell Therapy

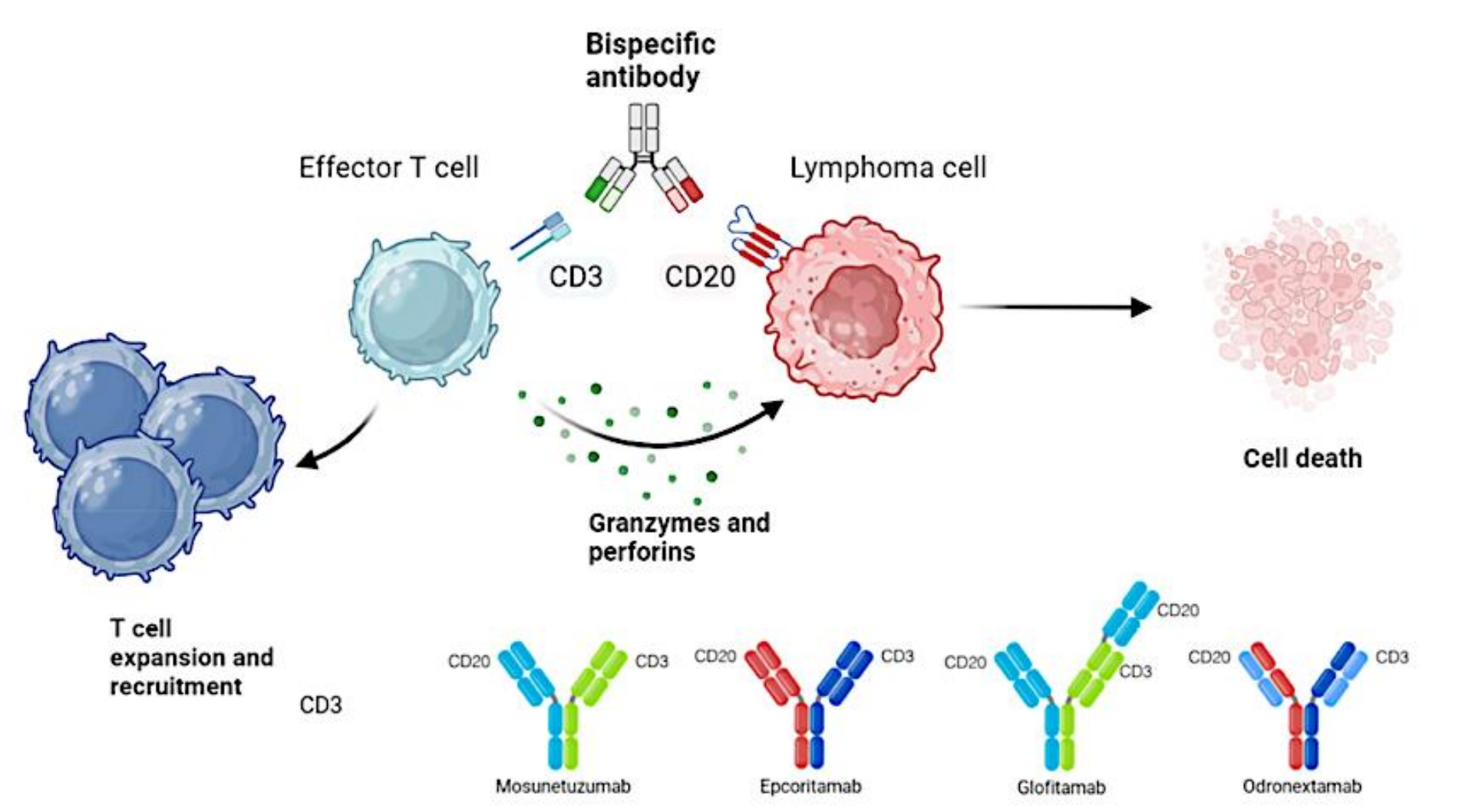
Module 5: Clinician Survey Results

Module 6: Bispecific Antibodies and Promising Investigational Strategies

Key Datasets

- Cassanello G et al. **Trial watch: Bispecific antibodies** for the treatment of relapsed or refractory large B-cell lymphoma. *Oncoimmunology* 2024 March 3;13(1):2321648.
- Danilov A et al. **Epcoritamab monotherapy** in patients (pts) with relapsed or refractory (R/R) chronic lymphocytic leukemia (CLL): Results from **CLL expansion and optimization cohorts of Epcore CLL-1**. ASH 2024;Abstract 883.
- Ahn I et al. **Updated efficacy and safety results** of the Bruton tyrosine kinase (**BTK**) **degrader BGB-16673** in patients with relapsed/refractory chronic lymphocytic leukemia/small lymphocytic lymphoma (CLL/SLL) from the ongoing **phase 1 CaDAnCe-101 study**. ASH 2025;Abstract 85.
- Shah NN et al. **Efficacy and safety** of the Bruton's tyrosine kinase (**BTK**) **degrader NX-5948** in patients with relapsed/refractory (R/R) chronic lymphocytic leukemia (CLL): Updated results from an **ongoing phase 1a/b study**. ASH 2024;Abstract 884.
- Woyach JA et al. **First-in-human study** of the **reversible BTK inhibitor nemtabrutinib** in patients with relapsed/refractory chronic lymphocytic leukemia and B-cell non-Hodgkin lymphoma. *Cancer Discov* 2024;14(1):66-75.

Mechanism of Action of CD20 x CD3 Bispecific Antibodies



Epcoritamab Monotherapy in Patients (Pts) with Relapsed or Refractory (R/R) Chronic Lymphocytic Leukemia (CLL): Results from CLL Expansion and Optimization Cohorts of EPCORE CLL-1

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Presented at the American Society of Hematology Annual Meeting; December 7–10, 2024; San Diego, CA

ASH 2024;Abstract 883

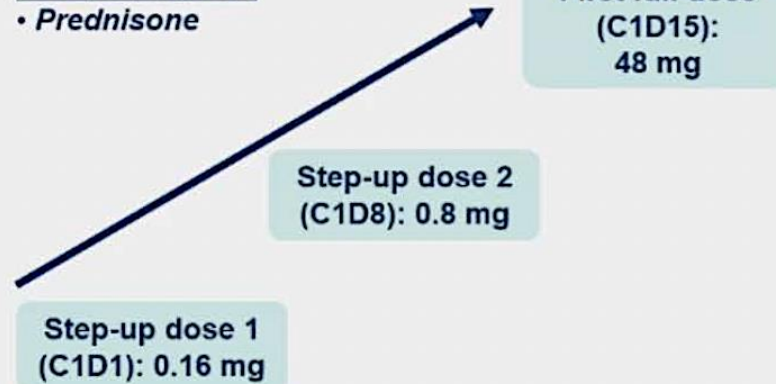
EPCORE CLL-1 Trial Expansion and Cycle 1 Optimization

Key inclusion criteria

- CD20+ R/R CLL
- ≥2 prior lines of systemic therapy
- ECOG PS 0–2
- Measurable disease with $\geq 5 \times 10^9/L$ B lymphocytes (expansion only)
- No prior allogeneic HSCT

Expansion (EXP; N=23)

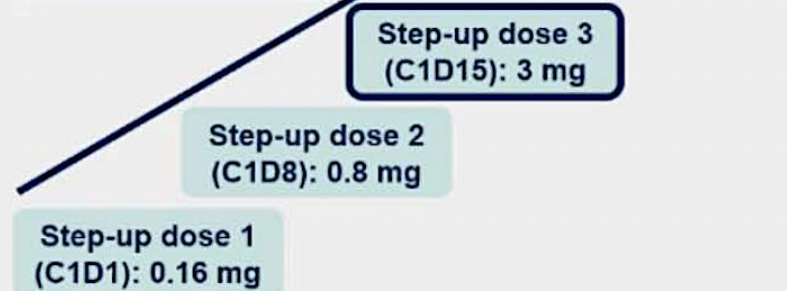
CRS prophylaxis
• Prednisone



Data cutoff: May 28, 2024
Median follow-up: 22.8 months

Cycle 1 Optimization (C1 OPT; N=17)

CRS prophylaxis
• Dexamethasone
• Adequate hydration



Data cutoff: May 28, 2024
Median follow-up: 2.9 months

- **Primary endpoint (EXP):** Overall response rate
- **Primary endpoint (C1 OPT):** Incidence and severity of CRS, ICANS, and clinical TLS
- **Key secondary endpoints (EXP):** CR rate, time to response, MRD (PBMCs using the clonoSEQ[®] assay), and safety/tolerability

- To ensure patient safety and better characterize CRS, inpatient monitoring was required for at least 24 hours after each epcoritamab dose in C1

ClinicalTrials.gov: NCT04623541; EudraCT: 2023-504828-25.

ICANS = immune effector cell-associated neurotoxicity syndrome; TLS = tumor lysis syndrome; PMBC = peripheral blood mononuclear cell

EPCORE CLL-1: Response Across Subgroups

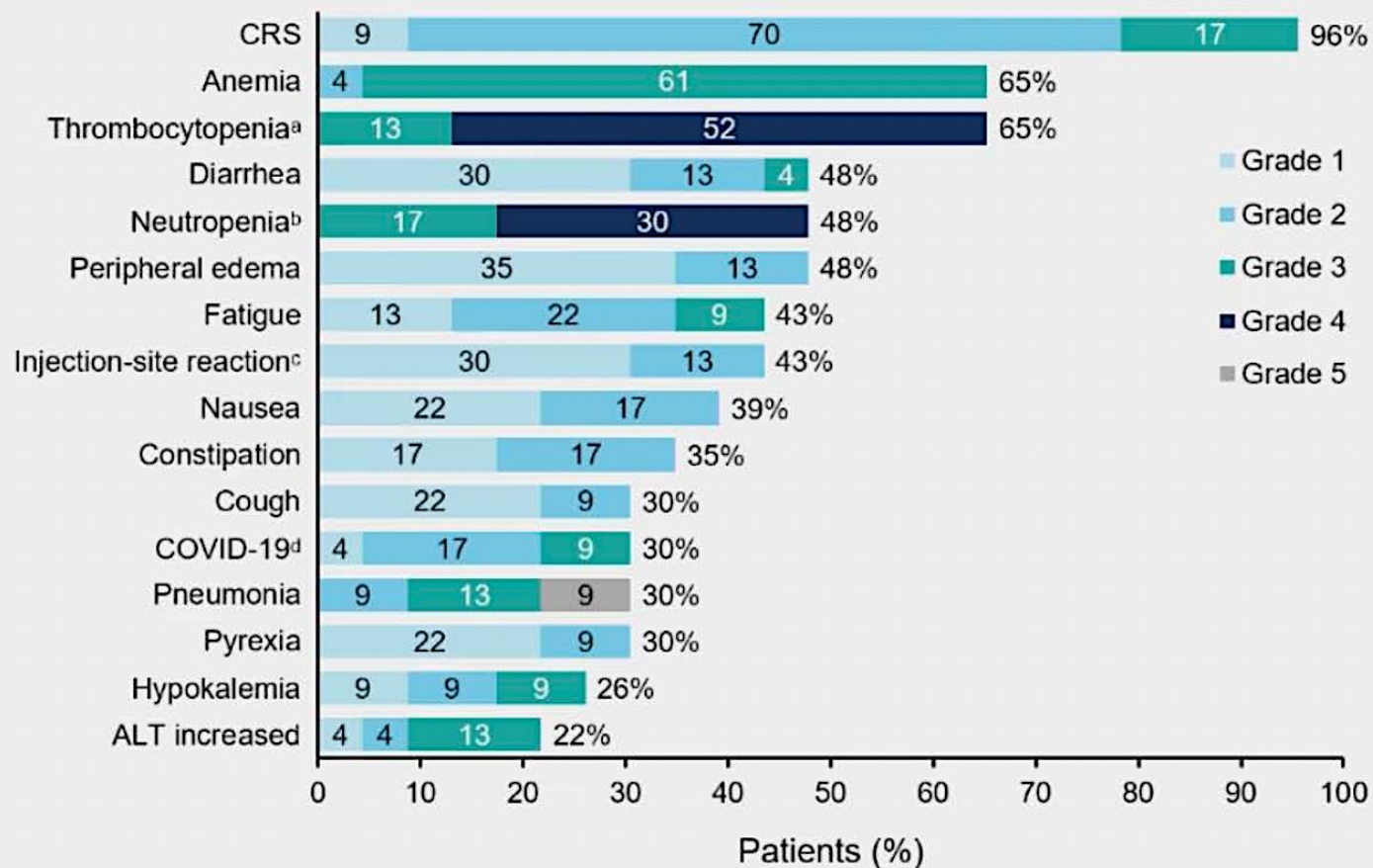
| Response, n (%) | EXP mFU: 22.8 months | | | | | C1 OPT mFU: 2.9 months |
|-------------------------------------|---------------------------|----------------------------|-----------------------------------|----------------------------------|-------------------------------------|----------------------------|
| | Full Analysis Set N=23 | Response Evaluable n=21 | <i>TP53</i> Aberration n=15 | <i>IGHV</i> Unmutated n=16 | Double Exposed ^a n=19 | Response Evaluable n=10 |
| Overall response^b | 14 (61) | 14 (67) | 10 (67) | 10 (63) | 10 (53) | 6 (60) |
| Complete response | 9 (39) | 9 (43) | 5 (33) | 7 (44) | 7 (37) | 1 (10) |
| Partial response | 5 (22) | 5 (24) | 5 (33) | 3 (19) | 3 (16) | 5 (50) |
| Stable disease | 4 (17) | 4 (19) | 2 (13) | 3 (19) | 4 (21) | 2 (20) |
| Progressive disease | 1 (4) | 1 (5) | 1 (7) | 0 | 1 (5) | 1 (10) |

- With limited follow-up, the C1 OPT regimen does not appear to affect epcoritamab efficacy
- uMRD4 in PBMCs was observed in most responders, including all patients with CR who were tested for MRD

| EXP MRD Negativity, n/n (%) ^c | uMRD4 | uMRD6 ^d |
|--|-----------|--------------------|
| Overall response ^b | 9/12 (75) | 8/12 (67) |
| Complete response | 7/7 (100) | 6/7 (86) |
| Partial response | 2/5 (40) | 2/5 (40) |
| Full analysis set | 9/23 (39) | 8/23 (35) |

Four patients (*TP53* aberration, n=2; *IGHV* unmutated, n=3; double exposed, n=4) in EXP and 1 in C1 OPT shown above were not evaluable or had no assessment, including 3 in EXP (*TP53* aberration, n=2; *IGHV* unmutated, n=2; double exposed, n=3) and 1 in C1 OPT who died without postbaseline assessment. ^aPatients previously treated with both a BTK inhibitor and a BCL-2 inhibitor. ^bResponse assessment according to iwCLL criteria. ^cPatients evaluated for MRD had at least 1 on-treatment MRD result and were not MRD negative at baseline. MRD was only evaluated in patients with CR or PR. ^dTwo of 3 evaluated patients had uMRD6 in bone marrow at or shortly after the first CR assessment. mFU, median follow-up.

EPCORE CLL-1: Treatment-Emergent Adverse Events (TEAEs)

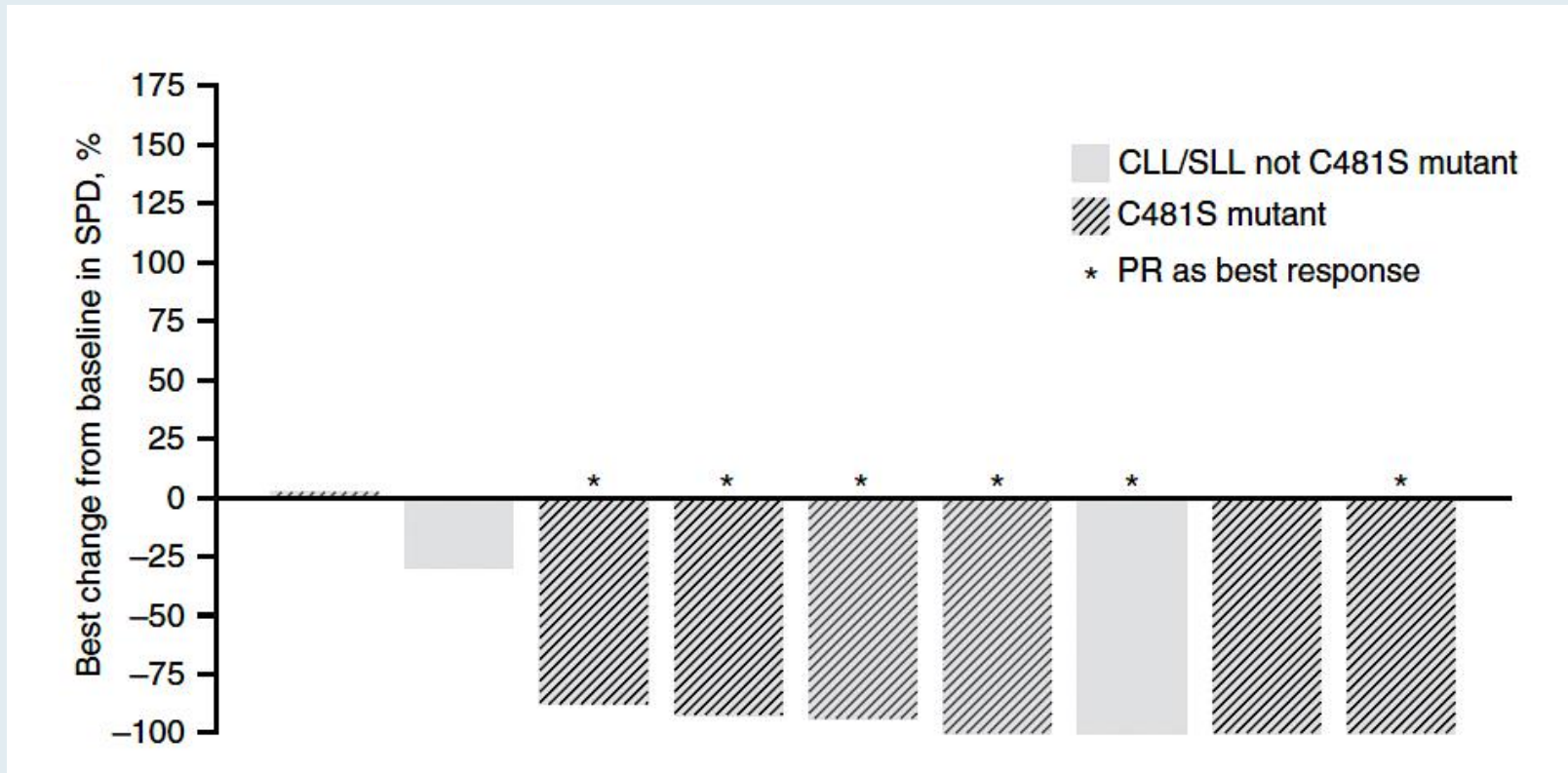


| Patients With ≥1 Event, n (%) | EXP N=23 |
|-------------------------------|-------------|
| Anemia | 15 (65) |
| At study entry | 14 (61) |
| In first 8 weeks | 15 (65) |
| Thrombocytopenia | 15 (65) |
| At study entry | 14 (61) |
| In first 8 weeks ^a | 14 (61) |
| Neutropenia | 11 (48) |
| At study entry | 1 (4) |
| In first 8 weeks ^b | 11 (48) |

- TEAEs were primarily low grade (G1–2)
- TEAEs led to treatment discontinuation in 5 patients from EXP and 1 patient from C1 OPT
- 4 fatal TEAEs^e occurred in EXP; none occurred in C1 OPT

^aCombined term includes thrombocytopenia and decreased platelet count. ^bCombined term includes neutropenia, decreased neutrophil count, and febrile neutropenia. Three patients had febrile neutropenia (EXP, n=2 [grades 1 and 3]; C1 OPT, n=1 [grade 3]). ^cCombined term includes injection-site reaction, bruising, erythema, rash, and swelling. ^dCombined term includes COVID-19 and COVID-19 pneumonia. ^eFatal TEAEs were pneumonia (n=2), sepsis (n=1), and squamous cell carcinoma of the skin (n=1); 1 case of pneumonia was considered related to epcoritamab.

Nemtabrutinib: A Novel Reversible BTK Inhibitor

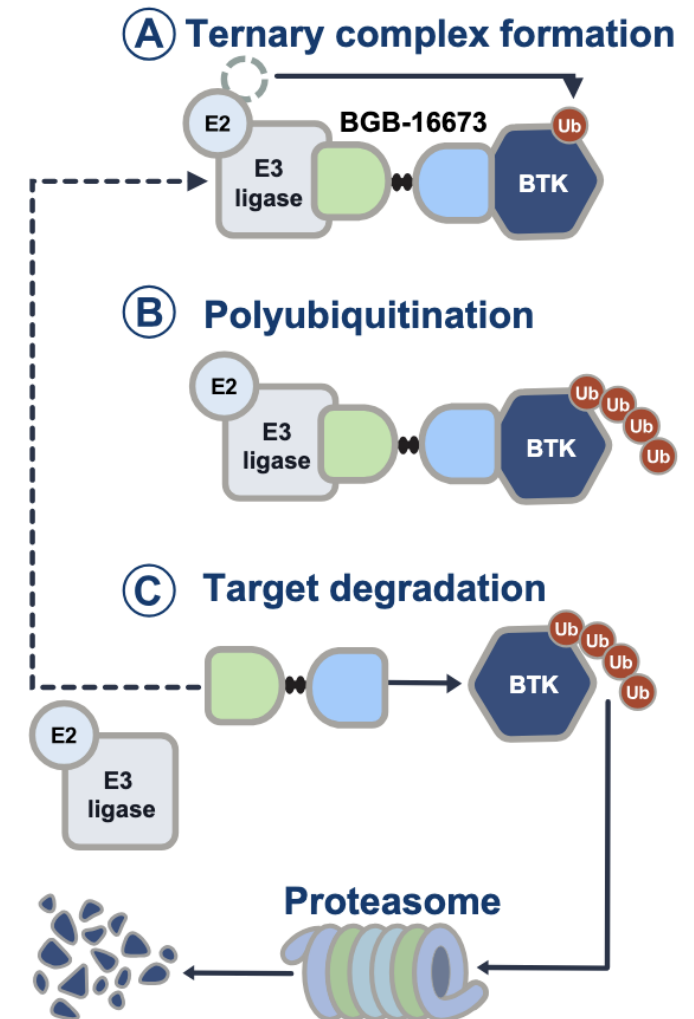


In a Phase I study, among the 22 patients with CLL, 8 (36.4%) achieved at least a partial remission with lymphocytosis as best response.

Among the 47 patients with CLL or non-Hodgkin lymphoma in the safety analysis, atrial fibrillation was reported in 1 patient (Grade 3), and no ventricular arrhythmias or unexplained deaths were reported. Hypertension was reported in 32% of patients (Grade 2 in 17%, Grade 3 in 15%).

BGB-16673: A Chimeric Degradation Activating Compound

BGB-16673 is an orally available protein degrader that blocks BTK signaling by tagging BTK for degradation through the cell's proteasome pathway, leading to tumor regression

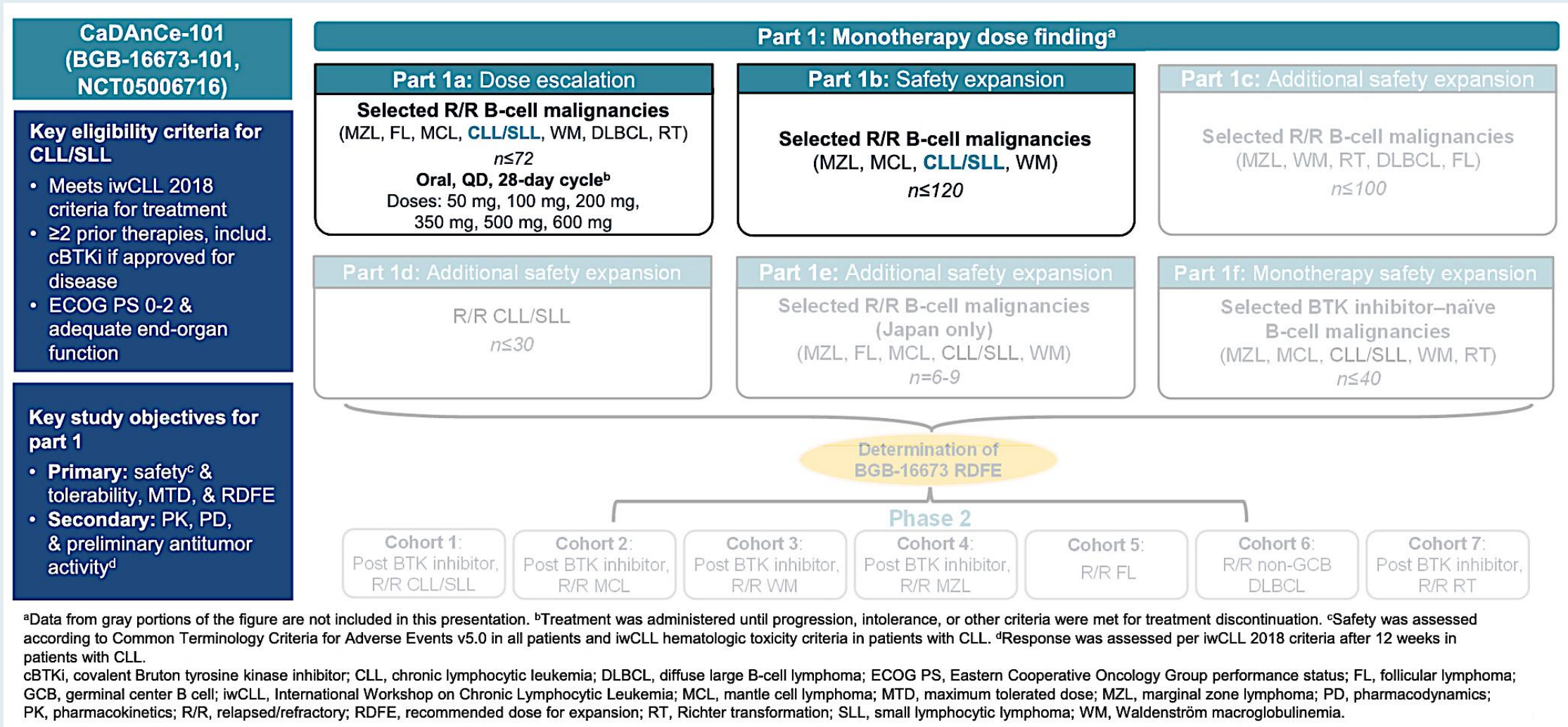


BTK, Bruton tyrosine kinase; cBTK, covalent Bruton tyrosine kinase; CLL/SLL, chronic lymphocytic leukemia/small lymphocytic lymphoma;

CNS, central nervous system; ncBTK, noncovalent Bruton tyrosine kinase; R/R, relapsed/refractory; ub, ubiquitin.

1. Moreno C. *Hematol Am Soc Hematol Educ Program*. 2020;2020:33-40; 2. Woyach JA, et al. *N Engl J Med*. 2014;370:2286-2294; 3. Wang E, et al. *N Engl J Med*. 2022;386:735-743; 4. Feng X, et al. EHA 2023. Abstract P1239; 5. Wang H, et al. EHA 2023. Abstract P1219; 6. Seymour JF, et al. ASH 2023; Abstract 4401.

CaDAnCe-101: A Phase I/II Dose-Escalation/Expansion Study of BGB-16673 for R/R B-Cell Cancers



CaDAnCe-101: Response Data

| | 50 mg (n=1) | 100 mg (n=22) | 200 mg (n=18) | 350 mg (n=15) | 500 mg (n=12) | Total (N=68) |
|---|---------------------|--------------------|--------------------|--------------------|--------------------|--------------------|
| Best overall response, n (%) | | | | | | |
| CR/CRI | 0 | 1 (4.5) | 1 (5.6) | 0 | 0 | 2 (2.9) |
| PR ^a | 1 (100) | 14 (63.6) | 12 (66.7) | 11 (73.3) | 11 (91.7) | 49 (72.1) |
| PR-L | 0 | 2 (9.1) | 4 (22.2) | 0 | 1 (8.3) | 7 (10.3) |
| SD | 0 | 5 (22.7) | 0 | 0 | 0 | 5 (7.4) |
| PD | 0 | 0 | 1 (5.6) | 1 (6.7) | 0 | 2 (2.9) |
| Discontinued prior to first assessment | 0 | 0 | 0 | 3 (20.0) | 0 | 3 (4.4) |
| ORR, n (%)^b | 1 (100) | 17 (77.3) | 17 (94.4) | 11 (73.3) | 12 (100) | 58 (85.3) |
| Time to first response, median (range), months^c | 2.9 (2.9-2.9) | 2.8 (2.0-6.2) | 2.9 (2.6-8.3) | 2.9 (2.6-19.4) | 2.8 (2.7-13.8) | 2.8 (2.0-19.4) |
| Time to best response, median (range), months | 2.9 (2.9-2.9) | 2.9 (2.0-11.1) | 3.0 (2.6-13.8) | 5.6 (2.6-19.4) | 8.4 (2.7-13.8) | 4.2 (2.0-19.4) |
| Duration of exposure, median (range), months | 29.6 (29.6-29.6) | 12.3 (3.4-25.4) | 14.4 (2.9-30.3) | 19.8 (0.2-28.5) | 20.4 (6.8-27.1) | 13.6 (0.2-30.3) |

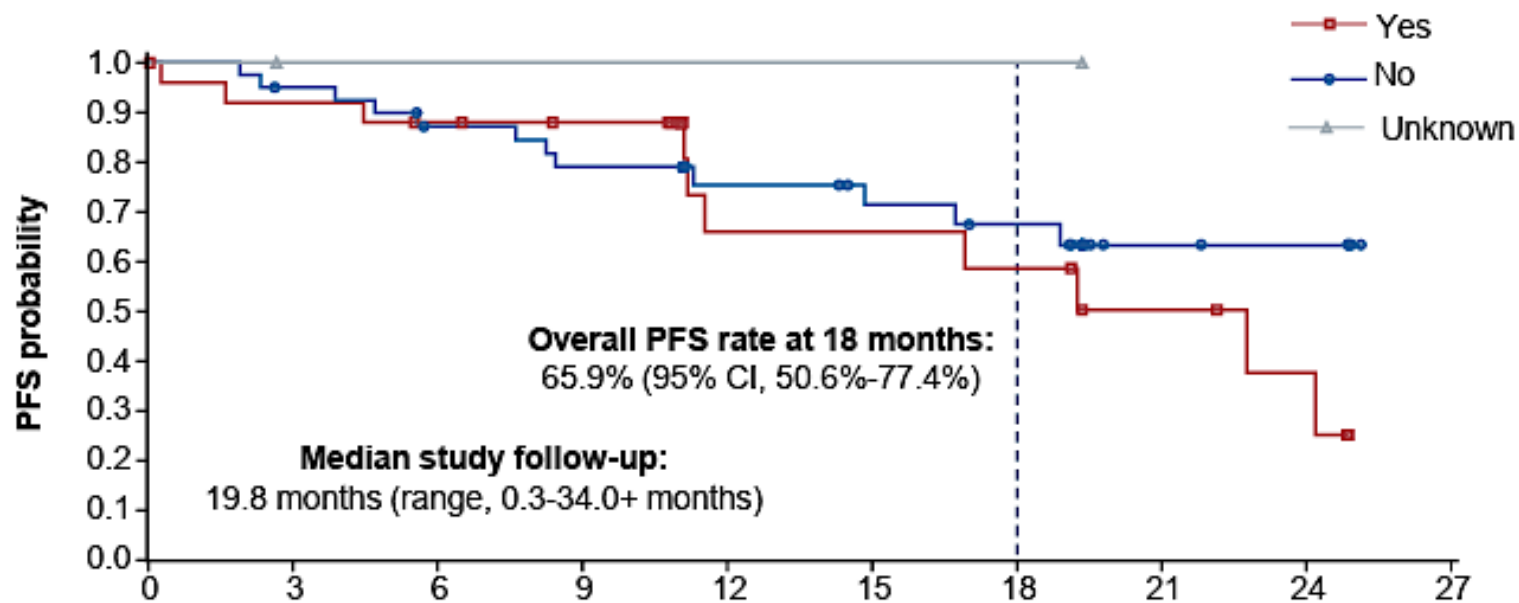
CaDAnCe-101: Overall Response Rate in High-Risk Subgroups

| Characteristic, n/N with known status (%) | ORR |
|---|--------------|
| Prior cBTKi + BCL2i | 41/44 (93.2) |
| Prior cBTKi + BCL2i + ncBTKi | 9/12 (75.0) |
| 6 or more prior lines of therapy | 13/16 (81.3) |
| del(17p) and/or <i>TP53</i> mutation | 37/46 (80.4) |
| Complex karyotype (≥ 3 abnormalities) | 16/22 (72.7) |
| <i>BTK</i> mutations | 20/26 (76.9) |
| <i>PLCG2</i> mutations | 9/10 (90.0) |

cBTKi = covalent BTK inhibitor; BCL2i = Bcl-2 inhibitor; ncBTKi = noncovalent BTK inhibitor

CaDAnCe-101: PFS Outcomes

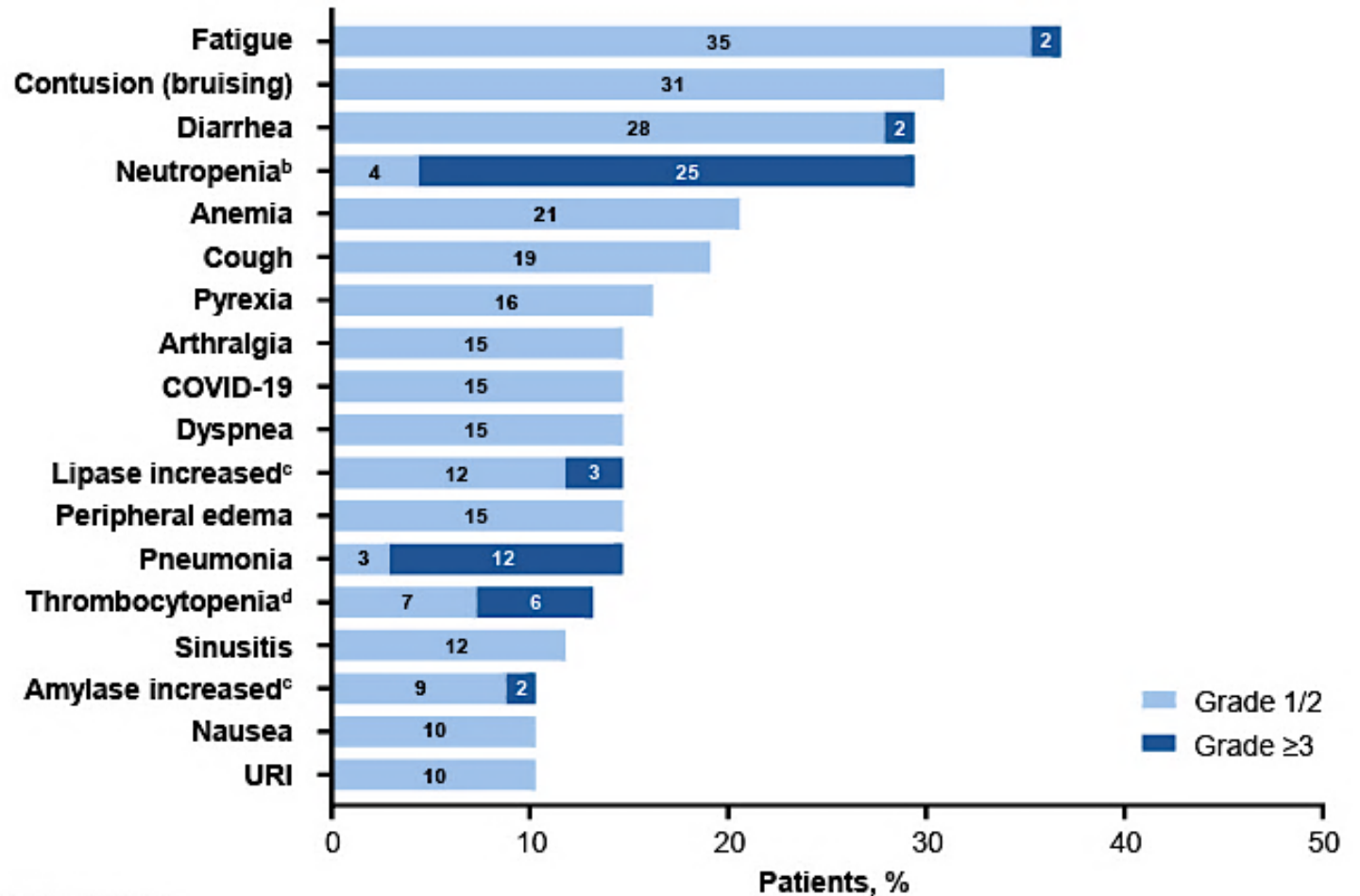
| BTK mutation status | 18-month PFS rate, % (95% CI) |
|----------------------------|--------------------------------------|
| Yes | 58.7 (30.6-78.7) |
| No | 67.5 (48.2-80.9) |
| Unknown | 100 (100-100) |



| No. at risk | Months | | | | | | | | | |
|-------------|--------|----|----|----|----|----|----|----|----|----|
| | 0 | 3 | 6 | 9 | 12 | 15 | 18 | 21 | 24 | 27 |
| Yes | 26 | 23 | 21 | 19 | 9 | 9 | 8 | 5 | 3 | 0 |
| No | 40 | 37 | 32 | 29 | 21 | 18 | 16 | 5 | 4 | 0 |
| Unknown | 2 | 1 | 1 | 1 | 1 | 1 | 1 | 0 | 0 | 0 |

CaDAnCe-101: Safety Profile

- The most common TEAEs were fatigue (36.8%) and contusion (bruising; 30.9%)
- Grade ≥ 3 neutropenia: n=17 (25.0%); 16 patients (23.5%) had grade ≥ 2 neutropenia at baseline
 - Neutropenic fever: n=1
- Atrial fibrillation: n=3 (grade 1, n=1; grade 2, n=2; all transient (2 of them lasting 1 day) in the context of infection and PD, assessed as unrelated to treatment)
- Treatment-related major hemorrhage^a: n=2 (one grade 3 subdural hemorrhage and one grade 3 post-procedural hematuria)

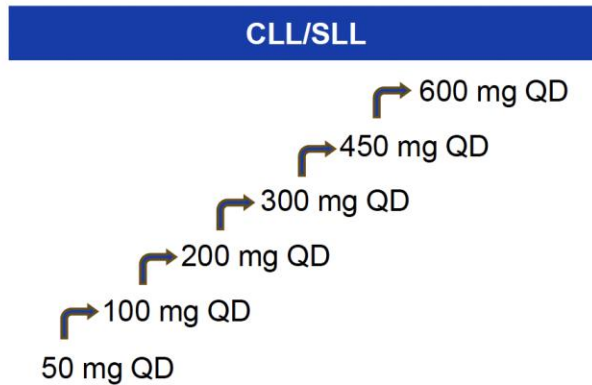


Median follow-up in safety-evaluable patients: 19.8 months (range, 0.3-34.0+ months).

^aGrade ≥ 3 , serious, or any central nervous system bleeding. ^bNeutropenia combines preferred terms *neutrophil count decreased* and *neutropenia*. ^cAll events were laboratory findings and were transient, mostly occurring during the first 1-3 cycles of treatment, with no clinical pancreatitis. ^dThrombocytopenia combines preferred terms *platelet count decreased* and *thrombocytopenia*. PD, progressive disease; TEAE, treatment-emergent adverse event; URI, upper respiratory tract infection.

Bexobrutideg (NX-5948): A Novel BTK Degradator Phase Ia/b Trial Design

Phase 1a dose escalation (fully enrolled)



CLL Phase 1b randomized cohort 1 (fully enrolled; 200 vs 600 mg)

CLL/SLL 200 mg QD
Prior BTKi and BCL2i

CLL/SLL 600 mg QD
Prior BTKi and BCL2i

CLL Phase 1b expansion, other cohorts (ongoing; all 600 mg)

Non-C481S BTK
mutations, prior
BTKi and BCL2i

Prior non-covalent
BTKi, no BCL2i

TP53 or 17p
deletion, 2L, prior
BTKi, no BCL2i

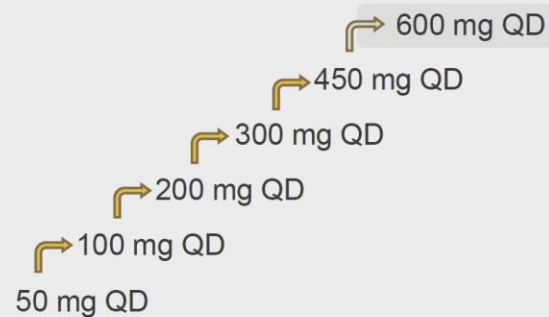
2L+, prior BTKi,
no BCL2i

BTKi-naïve

With wAIHA,
prior BTKi

With CNS
involvement,
prior BTKi

WM/NHL



NHL/WM Phase 1b expansion cohorts (600 mg)

MZL

Marginal zone
lymphoma

FL

Follicular lymphoma

WM

Waldenström
macroglobulinemia

MCL

Mantle cell lymphoma

DLBCL

Diffuse large B-cell
lymphoma

PCNSL

Primary CNS
lymphoma

2L+, second line +; BCL2i, B-cell lymphoma 2 inhibitor; BTKi, Bruton's tyrosine kinase inhibitor; CLL, chronic lymphocytic leukemia; CNS, central nervous system; NHL, non-Hodgkin's lymphoma; QD, once daily; SLL, small lymphocytic lymphoma; wAIHA, warm autoimmune hemolytic anemia; WM, Waldenström macroglobulinemia

Bexobrutideg (NX-5948): Overall Safety Summary

Tolerable safety profile, consistent between the RP2D 600 mg and overall study population

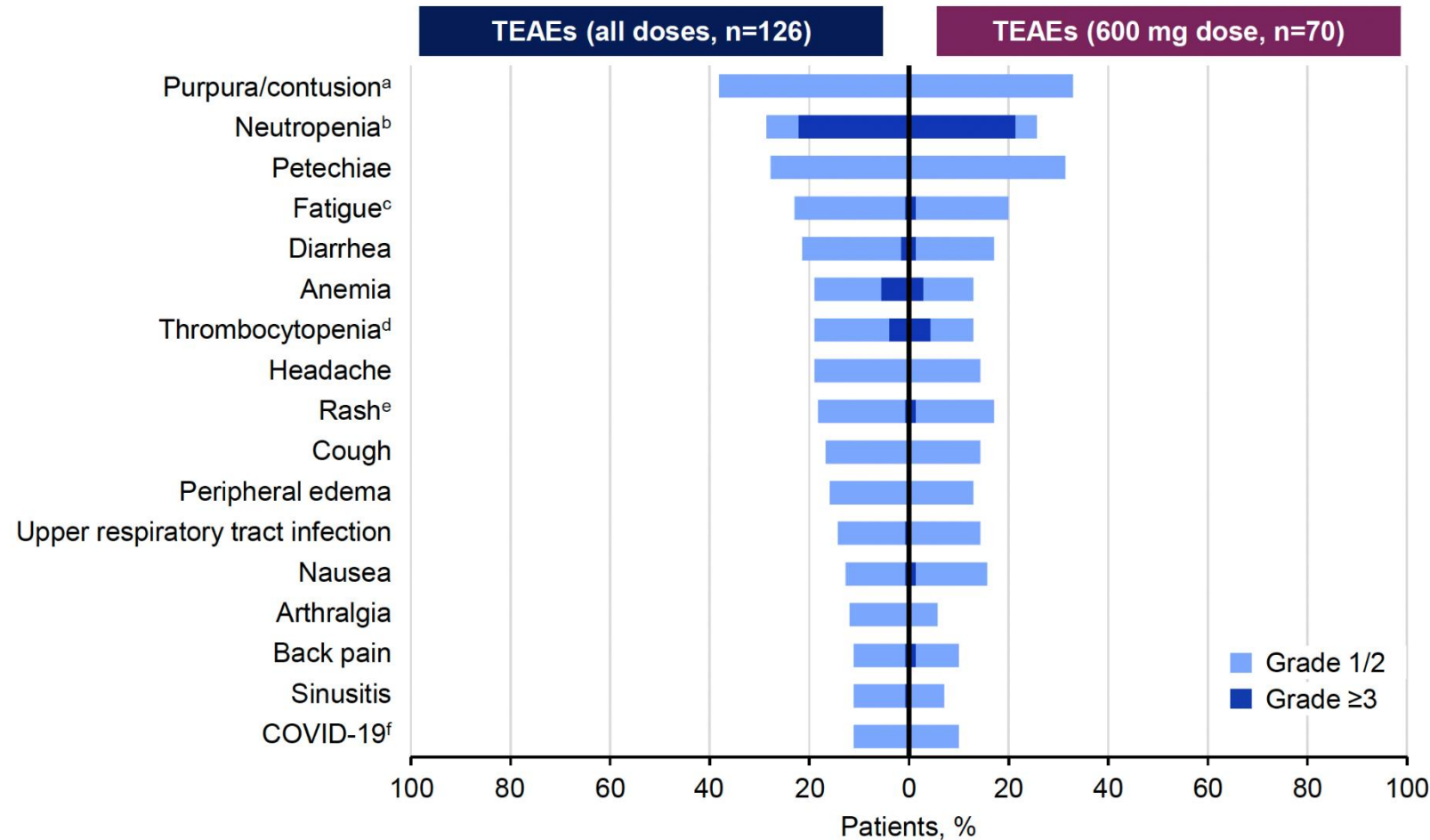
| | Phase 1a/b – all patients (n=126) | Phase 1a/b 600 mg (n=70) |
|---|--------------------------------------|-----------------------------|
| Any TEAE, n (%) | 114 (90.5) | 60 (85.7) |
| Treatment related | 95 (75.4) | 51 (72.9) |
| Grade ≥3 | 62 (49.2) | 31 (44.3) |
| Treatment-related | 31 (24.6) | 18 (25.7) |
| SAE | 27 (21.4) | 10 (14.3) |
| Treatment-related | 7 (5.6) | 3 (4.3) |
| Grade 5^a | 3 (2.4) | 1 (1.4) |
| Treatment-related | 0 | 0 |
| Leading to treatment discontinuation | 8 (6.3) | 4 (5.7) |
| Treatment-related | 5 (4.0) | 2 (2.9) |
| DLT | 0 | 0 |
| Median duration of treatment, months (range) | 7.1 (0.0–32.3) | 3.6 (0.0–18.0) |

^aGrade 5 AEs: pulmonary embolism; death not otherwise specified; pneumonia
 AE, adverse event; DLT, dose-limiting toxicity; RP2D, recommended Phase 2 dose; SAE, serious AE; TEAE, treatment-emergent AE

Data cutoff: 19 Sep 2025

Bexobrutideg (NX-5948): Treatment-Emergent Adverse Events

Comparable AE profile for patients at the RP2D 600mg dose and overall population



- Tolerable safety profile consistent with prior disclosures
- No dose-limiting toxicities
- No systemic fungal infections or Grade 4 infections of any kind reported
- Single event of new onset atrial fibrillation consistent with the rate in the age-matched general population
- 3 Grade 5 AEs (death not otherwise specified; pulmonary embolism; pneumonia; all deemed not related to bexobrutideg)

^aPurpura/contusion includes episodes of contusion or purpura; ^bAggregate of 'neutrophil count decreased' or 'neutropenia'; ^cFatigue was transient; ^dAggregate of 'thrombocytopenia' and 'platelet count decreased'; ^eAggregate of 'rash' and 'rash maculopapular' and 'rash pustular'; ^fAggregate of 'COVID-19' and 'COVID-19 pneumonia'
 AE, adverse event; NOS, not otherwise specified; RP2D, recommended Phase 2 dose; TEAE, treatment-emergent adverse event

Data cutoff: 19 Sep 2025

Bexobrutideg (NX-5948): Objective Response Rate and Median Duration of Response

Encouraging ORR and long median duration of response

| Response-evaluable patients | Phase 1a (n=47) |
|--|-------------------------|
| Objective response rate (ORR),^a % (95% CI) | 83.0 (69.2–92.4) |
| Disease control rate (DCR),^b % (95% CI) | 95.7 (85.5–99.5) |
| Best response,^c n (%) | |
| Complete response (CR) | 2 (4.3) |
| Nodal partial response (nPR) | 1 (2.1) |
| Partial response (PR/PR-L) | 36 (76.6) |
| Stable disease (SD) | 6 (12.8) |
| Progressive disease (PD) | 2 (4.3) |
| Median follow-up,^d months (range) | 19.0 (13.5–32.3) |
| Median duration of response, months (95% CI) | 20.1 (12.2–NE) (n=39) |

^aObjective response rate includes CR + nPR + PR + PR-L; ^bDisease control rate includes CR + nPR + PR/PR-L + SD; ^cPercentages are based on the number of patients dosed who had at least one post-baseline disease assessment or documented clinical PD; ^dTime from treatment start to data cutoff
CI, confidence interval; **CR**, complete response; **DCR**, disease control rate; **NE**, not evaluable; **nPR**, nodal partial response; **ORR**, objective response rate; **PD**, progressive disease; **PR**, partial response; **PR-L**, partial response with lymphocytosis; **SD**, stable disease

Data cutoff: 19 Sep 2025

Questions?