First-line treatment of chronic lymphocytic leukemia with ibrutinib plus obinutuzumab versus chlorambucil plus obinutuzumab: final analysis of the randomized, phase III iLLUMINATE trial

Carol Moreno,1,2 Richard Greil,1 Fatih Demirkan,4 Alessandra Tedeschi,6 Bertrand Anz,6 Loree Larratt,7 Martin Simkovic,8 Jan Novak,9 Vladimir Strugov,10 Devinder Gill,11 John G. Gribben,12 Kevin Kwei,13 Sandra Dai,13 Emily Hsu,13 James P. Dean13 and Ian W. Flinn14

1Hospital de la Santa Creu i Sant Pau, Autonomous University of Barcelona, Barcelona, Spain; 2Josep Carreras Leukemia Research Institute, Barcelona, Spain; 33rd Medical Department, Paracelsus Medical University, Salzburg Cancer Research Institute-CCCIT, Salzburg, Austria; 4Dokuz Eylul University, Division of Hematology, Izmir, Turkey; 5Niguarda Ca Granda Hospital, Milan, Italy; 6Tennessee Oncology, Chattanooga, TN, USA; 7University of Alberta Hospital, Edmonton, Alberta, Canada; 8Department of Internal Medicine, Haematology, University Hospital and Medical School Hradec, Králové, Czech Republic; 9University Hospital Kralovske Vinohrady and Third Faculty of Medicine, Charles University, Prague, Czech Republic; 10Almazov National Medical Research Centre, St. Petersburg, Russia; 11Princess Alexandra Hospital, Brisbane, Queensland, Australia; 12Barts Cancer Institute, London, UK; 13Pharmacyclics LLC, an AbbVie Company, Sunnyvale, CA, USA and 14Sarah Cannon Research Institute, Nashville, TN, USA

Abstract

iLLUMINATE is a randomized, open-label phase III study of ibrutinib plus obinutuzumab (n=113) versus chlorambucil plus obinutuzumab (n=116) as first-line therapy for patients with chronic lymphocytic leukemia or small lymphocytic lymphoma. Eligible patients were aged ≥65 years, or <65 years with coexisting conditions. Patients received oral ibrutinib 420 mg once daily until disease progression or unacceptable toxicity or six cycles of oral chlorambucil, each in combination with six cycles of intravenous obinutuzumab. After a median follow-up of 45 months (range, 0.2–52), median progression-free survival continued to be significantly longer in the ibrutinib plus obinutuzumab arm than in the chlorambucil plus obinutuzumab arm (median not reached versus 22 months; hazard ratio=0.25; 95% confidence interval: 0.16–0.39; P<0.0001). The best overall rate of undetectable minimal residual disease (<0.01% by flow cytometry) remained higher with ibrutinib plus obinutuzumab (38%) than with chlorambucil plus obinutuzumab (25%). With a median treatment duration of 42 months, 13 months longer than the primary analysis, no new safety signals were identified for ibrutinib. As is typical for ibrutinib-based regimens, common grade ≥3 adverse events were most prevalent in the first 6 months of ibrutinib plus obinutuzumab treatment and generally decreased over time, except for hypertension. In this final analysis with up to 52 months of follow-up (median 45 months), ibrutinib plus obinutuzumab showed sustained clinical benefit, in terms of progression-free survival, in first-line treatment of chronic lymphocytic leukemia, including in patients with high-risk features. ClinicalTrials.gov identifier: NCT02264574.

Introduction

Over the past decade, the introduction of novel therapies targeting B-cell receptor signaling has shifted the treatment landscape for patients with chronic lymphocytic leukemia (CLL), including for patients with high-risk genomic features who previously had limited treatment options.1 Ibrutinib is an oral, once-daily inhibitor of Bruton tyrosine kinase (BTK) approved for the treatment of patients with CLL, including in combination with the anti-CD20 monoclonal antibody obinutuzumab based on the primary results of the iLLUMINATE study.2,3 In two first-line studies ibrutinib or ibrutinib-based combination therapy was associated with superior progression-free survival (PFS) and overall survival (OS) compared with chlorambucil4 or fludarabine, cyclophosphamide, and rituximab (FCR)5 regimens in patients with CLL/small lymphocytic lymphoma (SLL). Ibrutinib (with or without rituximab) was also associated with superior PFS compared to benda-
Here we report the final analysis of iLLUMINATE with a median follow-up of 45 months.

Methods

Study design

iLLUMINATE was a multicenter, randomized, open-label, phase III study that enrolled patients at 71 sites in Australia, New Zealand, Canada, Israel, Turkey, the European Union, and the USA (ClinicalTrials.gov identifier: NCT02264574). The study was conducted in accordance with the Declaration of Helsinki, the International Conference on Harmonisation Guidelines for Good Clinical Practice, and local regulations. The protocol was approved by the institutional review boards, research ethics boards, or independent ethics committees of participating institutions. All patients provided written informed consent. Full details of the study methodology have been published previously and are briefly described below.

Participants

Eligible patients had previously untreated CLL/SLL requiring treatment according to International Workshop on Chronic Lymphocytic Leukemia (iwCLL) criteria and were aged ≥65 years or <65 years with at least one of the following co-existing conditions: a cumulative illness rating score >6, creatinine clearance <70 mL/min, del(17p) confirmed by fluorescence in situ hybridization analysis, or TP53 mutation confirmed by next-generation sequencing. Additional eligibility criteria included an Eastern Cooperative Oncology Group (ECOG) performance status 0-2, measurable lymph node disease (longest diameter >1.5 cm), adequate hematologic function (absolute neutrophil count ≥1x10⁹ cells/L; platelet count >50x10⁹ cells/L), and adequate hepatic and renal function.

Treatments

Patients were randomized 1:1 to receive ibrutinib plus obinutuzumab or chlorambucil plus obinutuzumab, as stratified by geographic region (North America vs. rest of world), ECOG performance status (0-1 vs. 2), and cytogenetic status (del[17p] vs. del[11q] without del[17p] vs. others [neither del[11q] nor del[17p]). Patients received oral ibrutinib 420 mg once daily until disease progression or unacceptable toxicity or oral chlorambucil 0.5 mg/kg body weight on days 1 and 15 of each 28-day cycle for six cycles. Intravenous obinutuzumab (100 mg on day 1 and 900 mg on day 2, and 1,000 mg on days 8 and 15 of cycle 1, followed by 1,000 mg on day 1 of each subsequent 28-day cycle) was administered for up to six cycles.

Outcomes

The primary endpoint for the final analysis was PFS by investigator assessment based on iwCLL 2008 criteria with subsequent clarifications to account for treatment-related lymphocytosis. Secondary endpoints included overall response rate (ORR; defined as complete response [CR], CR with incomplete bone marrow (BM) recovery [CRI], nodular partial response [nPR], or partial response [PR]), rate of undetectable minimal residual disease (MRD) (<1 CLL cell per 10,000 leukocytes in peripheral blood (PB) and/or BM aspirate, as measured by central laboratory flow cytometry), OS, sustained hematologic improvement (continuous ≥2 g/dL increase in hemoglobin levels from baseline, or ≥50% increase in platelet counts from baseline for ≥56 days without blood transfusion or growth factors), and safety. Responses were assessed by the investigator for the final analysis. Non-hematologic adverse events were graded using National Cancer Institute Common Terminology Criteria for Adverse Events, version 4.03; hematologic adverse events were assessed using iwCLL criteria. Because the additional follow-up since the primary analysis extends beyond the adverse event reporting period for the chlorambucil plus obinutuzumab arm, safety findings are presented only for the ibrutinib plus obinutuzumab arm.
Results

Patient demographics and characteristics
Two hundred twenty-nine patients were randomized to receive ibrutinib plus obinutuzumab (n=113) or chlorambucil plus obinutuzumab (n=116). Patient baseline demographics and disease characteristics were generally comparable between treatment arms (Online Supplementary Table S1). The median age was 71 years (range, 40-87), with most patients (n=183; 80%) aged ≥65 years. Most patients (148 [65%]) had high-risk disease features of del(17p), TP53 mutation, del(11q), or unmutated IGHV.

Patient disposition and treatment
The final analysis was performed upon study completion. The median time on study was 45 months (range, 0.2-52 months) for the total study population, 45.5 months (range, 0.2-52 months) for the ibrutinib plus obinutuzumab arm, and 43 months (range, 1-52 months) for the chlorambucil plus obinutuzumab arm. With study completion, discontinuations across both arms were due to sponsor termination of the study (74%; n=170), death (18%; n=42), withdrawal of consent (6%; n=14), and other (1%; n=1 each: high dose of steroids, obinutuzumab allergy, and infusion reaction). Of patients treated with ibrutinib and obinutuzumab, 26 (23%) and 10 (9%) patients, respectively, discontinued these agents due to adverse events; notably, treatment duration in the ibrutinib plus obinutuzumab arm was longer than that in the chlorambucil plus obinutuzumab arm (median: 42 vs. 5 months). In the ibrutinib plus obinutuzumab arm, discontinuation of ibrutinib therapy due to disease progression was infrequent (n=5; 4%); 65 (58%) patients remained on ibrutinib until study completion (Table 1) and 100 (88%) patients completed six cycles of obinutuzumab (Online Supplementary Figure S1). In the chlorambucil plus obinutuzumab arm, 103 (89%) patients and 100 (86%) patients completed the planned six cycles of chlorambucil and obinutuzumab, respectively.

Of the 116 patients randomized to chlorambucil plus obinutuzumab, 50 (43%) patients subsequently received ibrutinib with study crossover (n=45; 39%) or with commercial ibrutinib (n=6; 5%); one patient received both commercial and crossover ibrutinib. At study completion, 35 of the 45 crossover patients (78%) were still receiving single-agent ibrutinib. All patients received concomitant medications during study treatment; in the ibrutinib plus obinutuzumab arm, 63/113 (56%) patients received any anticoagulant or antiplatelet agent; of these, 36/113 (32%) patients received anticoagulants and 48/113 (42%) received antiplatelet agents. Ninety-three of 113 patients (82%) received systemic antibacterial drugs. Of the 113 patients in the ibrutinib plus obinutuzumab arm, 78 (69%) received acid-reducing medications; 71/113 (63%) received proton pump inhibitors, 14/113 (12%) received H2-receptor antagonists, and 8/113 (7%) received other acid-reducing agents.

Progression-free survival
At the final analysis, with a median follow-up of 45 months, PFS remained significantly longer in the ibrutinib plus obinutuzumab arm (median not reached; 95% confidence interval [95% CI], 49 months to not estimable [NE]) than in the chlorambucil plus obinutuzumab arm (22 months; 95% CI: 18-27 months), resulting in a 75% Table 1. Summary of treatment disposition.

| Treatment status, N (%) | Ibrutinib plus obinutuzumab (N=113) | Chlorambucil plus obinutuzumab (N=116) |
|------------------------|-------------------------------------|--------------------------------------|
|                        | Ibrutinib | Obinutuzumab | Chlorambucil | Obinutuzumab |
| Did not receive study drug | 0 | 0 | 1 (1) | 1 (1) |
| Completed/ongoing at study closure | 65 (58) | 100 (88) | 103 (89) | 100 (86) |

| Primary reason for discontinuation of study treatment, N (%) | Ibrutinib plus obinutuzumab (N=113) | Chlorambucil plus obinutuzumab (N=116) |
|-------------------------------------------------------------|-------------------------------------|--------------------------------------|
| Disease progression | 5 (4) | 1 (1) | 0 | 0 |
| Adverse events     | 25 (22) | 10 (9) | 11 (9) | 15 (13) |
| Death              | 5 (4) | 0 | 0 | 0 |
| Withdrawal of consent | 6 (5) | 1 (1) | 0 | 0 |
| Investigator decision | 3 (3) | 0 | 1 (1) | 0 |
| Other              | 4 (4) | 1 (1) | 0 | 0 |
reduction in the risk of disease progression or death with ibrutinib plus obinutuzumab (hazard ratio [HR]=0.25; 95% CI: 0.16-0.39; \( P < 0.0001 \)) (Figure 1). The estimated PFS at 42 months was 74% in the ibrutinib plus obinutuzumab arm and 33% in the chlorambucil plus obinutuzumab arm. Similar to the overall population, there was a significant PFS benefit for patients with high-risk features (del(17p), del(17p)/TP53 mutation, del(11q) and/or unmutated IGHV) treated with ibrutinib plus obinutuzumab versus chlorambucil plus obinutuzumab (HR=0.17; 95% CI: 0.10-0.28; \( P < 0.0001 \)). Within the high-risk population, PFS estimates at 42 months were significantly higher in the ibrutinib plus obinutuzumab arm than in the chlorambucil plus obinutuzumab arm.

**Figure 1.** Progression-free survival per investigator assessment in the intention-to-treat population. CI: confidence interval; mo: months; NE, not estimable; PFS, progression-free survival.

**Figure 2.** Progression-free survival per investigator assessment in the high-risk population of patients with del(17p), del(11q), TP53 mutations, and/or unmutated IGHV. CI: confidence interval; mo: months; NE: not estimable; PFS: progression-free survival.
tuzumab arm (70% vs. 12%). At 48 months, the PFS benefit was maintained since the time of the primary analysis at 30 months (70% vs. 77%, respectively) (Figure 2). The exclusion of patients with del(17p) did not affect 42-month PFS estimates for the rest of the high-risk population, which were 71% and 14%, respectively, for the two treatment arms (Online Supplementary Figure S2).

Likewise, a PFS benefit with ibrutinib plus obinutuzumab was observed irrespective of IGHV mutation status (Figure 3). Among patients with unmutated IGHV, up to 48 months median PFS was not reached with ibrutinib plus obinutuzumab versus 15 months with chlorambucil plus obinutuzumab (HR=0.17; 95% CI: 0.10-0.29). Whereas the median PFS was not reached in either treatment arm for patients with mutated IGHV, ibrutinib plus obinutuzumab significantly reduced the risk of progression or death in this subgroup (HR=0.20; 95% CI: 0.07-0.59). Within the ibrutinib plus obinutuzumab arm, the estimated PFS at 48 months was 67% and 89% for patients with unmutated and mutated IGHV, respectively. The PFS benefit from ibrutinib plus obinutuzumab among both subgroups of unmutated and mutated IGHV patients persisted after excluding patients with del(17p) (unmutated: HR=0.17; 95% CI: 0.09-0.30; mutated: HR=0.25; 95% CI: 0.08-0.74) (Online Supplementary Figure S3). The hazard ratios of 0.20 and 0.17 observed in the mutated and unmutated IGHV subgroups of the ibrutinib plus obinutuzumab arm are consistent with the hazard ratio for the intention-to-treat population (HR=0.25) which includes other non-high-risk subgroups (Figure 1). Online Supplementary Figure S4 depicts consistently superior PFS associated with ibrutinib plus obinutuzumab across various subgroups of patients defined by age, Rai stage, ECOG performance status, and bulky disease, in addition to specified genomic risk factors mentioned above.

In patients randomized to ibrutinib plus obinutuzumab, no significant difference in PFS was shown between patients with or without del(17p)/TP53 mutation (HR=0.93; 95% CI: 0.32-2.69; P=0.895) (Figure 4). The median PFS was not reached in either subgroup, with estimated 48-month PFS...
rates of 77% and 74% in patients with and without del(17p)/TP53 mutation, respectively.

**Time to next treatment**
Over a median follow-up of 45 months, fewer patients receiving ibrutinib plus obinutuzumab initiated subsequent treatment for CLL/SLL compared with those receiving chlorambucil plus obinutuzumab (3/113 [3%] vs. 55/116 [47%] patients, respectively). The median time to next treatment was not reached in the ibrutinib plus obinutuzumab arm compared with 33 months (95% CI: 24 months-NE) in the chlorambucil plus obinutuzumab arm, corresponding to a 96% reduction in the risk of needing next-line therapy (HR=0.04; 95% CI: 0.01-0.13; \( P<0.0001 \)).

**Response rates**
Consistent with the primary analysis, the ORR by investigator assessment was higher in the ibrutinib plus obinutuzumab arm (n=103/113; 91%) than in the chlorambucil plus obinutuzumab arm (n=94/116; 81%) (Figure 5). CR rates, including CRi, in the ibrutinib plus obinutuzumab arm (47 [42%]) were slightly increased relative to rates reported at the primary analysis (46 [41%]); CR/CRi rates in the chlorambucil plus obinutuzumab arm were unchanged compared to those reported at the primary analysis (n=20 [17%]; \( P<0.0001 \) for the difference between arms). The median duration of response was not reached in the ibrutinib plus obinutuzumab arm (95% CI: NE-NE) and was 19 months (95% CI: 16-31) in the chlorambucil plus obinutuzumab arm.

**Minimal residual disease**
Initially MRD information was obtained from BM from all patients at cycle 9 and upon achieving CR, but subsequent protocol amendments enabled data collection from both BM and PB, and from all responders on a scheduled basis thereafter. Overall, MRD assessments were obtained from 101/113 patients on ibrutinib plus obinutuzumab (93 BM and 90 PB) and from 92/116 patients in the chlorambucil plus obinutuzumab arm (84 BM and 60 PB). Most patients with undetectable MRD (<0.01%) were subsequently followed with at least one repeat MRD assessment (91% for ibrutinib plus obinutuzumab; 86% for chlorambucil plus obinutuzumab), with a median follow-up of 23 and 30 months, respectively, following initial attainment of undetectable MRD. Overall, 38% (43/113) of patients in the ibrutinib plus obinutuzumab arm achieved undetectable MRD in BM or PB compared with 25% (29/116) in the chlorambucil plus obinutuzumab arm (\( P=0.033 \)) (Online Supplementary Table S2). Rates of undetectable MRD for the ibrutinib plus obinutuzumab arm were 25% for BM (28/113) and 33% for PB (37/113); rates for the chlorambucil plus obinutuzumab arm were 17% (20/116) in BM and 20% (23/116) in PB. In these patients, the cumulative rate of undetectable MRD (<0.01%) increased over the first 3 years with continuous ibrutinib therapy and then remained stable through final analysis (Figure 6). As expected, given the differences in PFS for the two arms following initial attainment of undetectable MRD, 60% of patients receiving ibrutinib plus obinutuzumab maintained undetectable MRD status through last PB or BM testing compared to 31% of patients receiv-
ing chlorambucil plus obinutuzumab (Online Supplementary Figure S5A). The cumulative MRD relapse rate (defined as MRD ≥1% after previously achieving undetectable MRD) was lower in the ibrutinib plus obinutuzumab group than in the chlorambucil plus obinutuzumab group (12% vs. 24%) and the median time to MRD relapse was not estimable (range, 0.03-44.2 months) in the ibrutinib plus obinutuzumab arm versus 37.5 months (range, 0.03-44.2 months) in the chlorambucil plus obinutuzumab arm. Examined among patients who achieved a best response of CR/CRi with ibrutinib plus obinutuzumab, 28/47 patients (60%) had undetectable MRD while among those in the chlorambucil plus obinutuzumab arm achieving CR, 15/20 patients (75%) had undetectable MRD in PB or BM. By contrast, among patients who achieved a best response of PR/nPR, 14/56 patients had undetectable MRD levels in PB or BM in the ibrutinib plus obinutuzumab arm (25%) and 14/74 (19%) in the chlorambucil plus obinutuzumab arms. Although limited by the small number of evaluable patients, the median PFS was similar in patients with CR/CRi, regardless of MRD status (Online Supplementary Figure S5B, C); undetectable MRD tended to correlate positively with longer PFS in patients with PR/nPR, especially in the chlorambucil plus obinutuzumab arm.

Likewise, in the high-risk population, more than double the percentage of patients in the ibrutinib plus obinutuzumab arm maintained undetectable MRD than in the chlorambucil plus obinutuzumab group ([n=15/23]; 65% vs. [n=3/11]; 27%). In high-risk patients with >24 months of MRD follow-up, five of six patients (83%) in the ibrutinib plus obinutuzumab arm remain MRD negative compared with two of five patients (40%) in the chlorambucil plus obinutuzumab arm. As in the overall population, the cumulative MRD relapse rate in the high-risk population after achieving undetectable MRD was lower in the ibrutinib plus obinutuzumab group than in the chlorambucil plus obinutuzumab group (0%; [n=0/23] vs. 27%; [n=3/11]). The median time to MRD relapse was not estimable in either the ibrutinib plus obinutuzumab arm (range, 0.03-38.6 months) or the chlorambucil plus obinutuzumab arm (range, 0.03-33.4 months).

**Overall survival**

At the time of final analysis, the median OS was not reached in either treatment arm (HR=1.08; 95% CI: 0.60-1.97; P=0.793) (Online Supplementary Figure S6). Among the 45 patients who crossed-over to single-agent ibrutinib, the 24-month OS was 88% (95% CI: 74.0-94.9) and six deaths were reported (13%). Overall, 22 (19%) patients in the ibrutinib plus obinutuzumab arm and 21 (18%) patients in the chlorambucil plus obinutuzumab died during the study.

**Adverse events in the ibrutinib plus obinutuzumab arm**

At final analysis, the median duration of exposure to ibrutinib was 42 months (range, 0.1-52) in the ibrutinib

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**Figure 5. Best overall response per investigator assessment with ibrutinib plus obinutuzumab versus chlorambucil plus obinutuzumab.** CR: complete response; CRi, complete response with incomplete bone marrow recovery; nPR, nodular partial response; PR, partial response.
plus obinutuzumab group, which was 13 months longer than ibrutinib exposure at the primary analysis (median 29 months); 73 patients (65%) received ≥3 years of ibrutinib and 18 (16%) received ≥4 years of ibrutinib therapy. All patients had completed or discontinued obinutuzumab (on both arms) and chlorambucil before the primary analysis, thus the median durations of treatment for obinutuzumab did not change since the primary analysis (4.6 months [range, 4.6-4.9] for the chlorambucil plus obinutuzumab group; 4.6 months [range, 4.6-4.7] for the ibrutinib plus obinutuzumab group). The median duration of treatment with chlorambucil was 5.1 months (range, 5.1-5.3) in the chlorambucil plus obinutuzumab group.

With ongoing treatment in the ibrutinib plus obinutuzumab arm, the most common adverse events (occurring in ≥10% of patients in either arm) were neutropenia (44%), thrombocytopenia (35%), diarrhea (35%), cough (29%), infusion-related reaction (25%), and arthralgia (24%) (Online Supplementary Table S3). The most common grade ≥3 adverse events in the ibrutinib plus obinutuzumab arm were neutropenia (36%), thrombocytopenia (19%), pneumonia (9%), atrial fibrillation (6%), and febrile neutropenia (5%). The prevalence of these grade ≥3 adverse events was highest during the first 6 months of treatment with ibrutinib plus obinutuzumab and generally decreased over time, with the exception of hypertension (Figure 7). The prevalence of grade ≥3 hypertension was 2% (n=2/113), 3% (n=3/98), 4% (n=3/84), and 3% (n=2/74) in years 0-1, 1-2, 2-3, and 3-4, respectively. No patient discontinued ibrutinib therapy due to hypertension of any grade. The prevalence of any grade atrial fibrillation decreased, being 10%

| Time       | Subjects with valid MRD | Cumulative Subjects with Valid MRD |
|------------|-------------------------|-----------------------------------|
|            | Ibrutinib plus obinutuzumab (n=113) | Chlorambucil plus obinutuzumab (n=116) | Ibrutinib plus obinutuzumab (n=113) | Chlorambucil plus obinutuzumab (n=116) |
| Year 1     | 85 (75)                 | 84 (72)                           | 85 (75)                           | 84 (72)                               |
| Year 2     | 72 (64)                 | 46 (40)                           | 98 (87)                           | 89 (77)                               |
| Year 3     | 85 (75)                 | 45 (39)                           | 101 (89)                          | 92 (79)                               |
| Year 4     | 67 (59)                 | 31 (27)                           | 101 (89)                          | 92 (79)                               |
| Year >4    | 3 (3)                   | 4 (3)                             | 101 (89)                          | 92 (79)                               |

Figure 6. Cumulative rates of minimal residual disease over time (in bone marrow or peripheral blood). Clb: chlorambucil; ibr: ibrutinib; MRD: minimal residual disease; Ob: obinutuzumab.
Figure 7. Prevalence of most common adverse events of any grade (≥20%) and grade ≥3 adverse events (≥3%) over time in patients treated with ibrutinib plus obinutuzumab.
(n=11/113), 10% (n=10/98), 8% (n=7/84), and 9% (n=7/74) in years 0–1, 1–2, 2–3, and 3–4, respectively; the prevalence of grade ≥3 atrial fibrillation was 4% (n=5/113), 2% (n=2/98), 0%, and 1% (n=1/74), in years 0–1, 1–2, 2–3, and 3–4, respectively. Among 17 patients with atrial fibrillation of any grade, atrial fibrillation led to discontinuation of ibrutinib in one patient (grade 3 event).

Seven major hemorrhagic adverse events (grade 1/2 for 2 events, grade 3 for 5 events, and no grade 4/5 events) occurred in five (4%) patients in the ibrutinib plus obinutuzumab arm: catheter site hematoma, ecchymosis, hematemesis, hemoptysis, post-procedural hematoma, subdural hematoma, and traumatic hematoma. Major hemorrhage led to discontinuation of ibrutinib in one patient (grade 2 hemorrhysis).

Over a median duration of 42 months of ibrutinib therapy, the only adverse event leading to discontinuation of ibrutinib in more than one patient was thrombocytopenia (n=2); all other adverse events leading to discontinuation of ibrutinib occurred in only one patient each. Of these, seven and six patients discontinued during the first 6 and 12 months of treatment, respectively. Four patients each discontinued at time intervals of >12 ≤24 months, >24 ≤36 months, and >36 ≤48 months.

In the ibrutinib plus obinutuzumab arm, 17 (15%) patients experienced a total of 32 adverse events (including 18 grade ≥3 events) that led to ibrutinib dose reduction, most commonly due to neutropenia (6 [5%] patients) (Online Supplementary Figure S7). Most (28/32 [88%]) of these adverse events resolved or recovered with dose reduction, including 94% (17/18) of grade ≥3 events.

At the time of final analysis, 14 and three patients had died from treatment-emergent adverse events in the ibrutinib plus obinutuzumab and chlorambucil plus obinutuzumab arms, respectively (Online Supplementary Table S4). Of the five deaths that occurred in the ibrutinib plus obinutuzumab arm with the additional follow-up, four were due to adverse events (1 each due to respiratory tract infection, pneumonia, septic shock and one death was due to an unknown cause after the adverse event reporting period). In the chlorambucil plus obinutuzumab arm, an additional two deaths occurred since the primary analysis, after crossover to the ibrutinib plus obinutuzumab arm. One was due to sepsis, the other was due to progressive disease.

**Outcomes after discontinuation of ibrutinib**

Thirty-eight patients discontinued ibrutinib after a median of 15.5 months (range, 0.1–43.7) of treatment for reasons other than progression or death at any time on study, 25 patients discontinued due to adverse events (Table 1). Among patients who discontinued ibrutinib for reasons other than progressive disease or death, eight patients had achieved a CR or CRi, 22 patients had PR or nPR, and one patient had stable disease; seven patients had no opportunity for response assessment. With a median follow-up of 17 months (range, 0.03–42) after ibrutinib discontinuation, 18-month PFS in these 38 patients who discontinued for reasons other than progression or death was 73.8% (95% CI: 55.4–85.5).

**Discussion**

Results from the final analysis of the iLLUMINATE study with up to 52 months of follow-up (median: 45 months) confirmed the durable efficacy of ibrutinib plus obinutuzumab in previously untreated patients with CLL. Ibrutinib plus obinutuzumab resulted in sustained PFS compared to chlorambucil plus obinutuzumab, with median PFS not reached after a median follow-up of 45 months and the risk of disease progression or death reduced by 75%. Notably, durable PFS was also seen in the high-risk patient population (i.e., with del[17p], TP53 mutation, del[11q], or unmutated IGHV) treated with ibrutinib plus obinutuzumab, confirming earlier observations from the primary analysis of iLLUMINATE. At 42 months, the PFS estimates among patients with del(17p), TP53, unmutated IGHV or del(11q) mutations were significantly higher in the ibrutinib plus obinutuzumab arm than in the chlorambucil plus obinutuzumab arm. Importantly, the benefit in PFS was observed regardless of high-risk features. A similar benefit was observed in the CLL14 trial for the venetoclax plus obinutuzumab regimen in patients with unmuted and mutated IGHV. These results support current global consensus guidelines, noting ibrutinib as a preferred therapeutic regimen for older patients and/or those with comorbidities regardless of the presence of del(17p)/TP53 mutations and unmuted CLL. The follow-up for other, similarly recommended novel treatment options is more limited.

With this extended follow-up, as has been observed consistently across several large, randomized phase III studies of ibrutinib in previously untreated and relapsed/refractory patients with CLL/SLL, highly durable PFS and OS have been confirmed even in patients with high-risk disease characteristics.

In line with the results of the iLLUMINATE trial, other phase III studies have also reported the superiority of ibrutinib-based regimens over chemoimmunotherapy in the first-line treatment of CLL. Concurrently, in the phase III ALLIANCE 041202 study in patients with CLL aged ≥65 years (median follow-up: 38 months), first-line single-agent ibrutinib or ibrutinib plus rituximab significantly prolonged PFS compared with bendamustine plus rituximab; estimated 2-year PFS rates were 87% and 88% versus 74%, respectively. In patients aged ≤70 years, the phase III ECOG1912 study (median follow-up: 34 months)
demonstrated that first-line treatment with ibrutinib plus rituximab resulted in significantly longer PFS (3-year rate 89% vs. 73%) and OS (3-year rate 99% vs. 92%) compared with FCR. With additional follow-up (median: 45 months), PFS favored ibrutinib plus rituximab over FCR (HR=0.39; 95% CI: 0.26-0.57; P<0.0001). With additional follow-up since the primary analysis of iL-UMINATE, the percentage of patients treated with ibrutinib plus obinutuzumab achieving undetectable MRD increased (from 34.5% at primary analysis to 38.1% at final analysis), demonstrating deepening of remission with continued ibrutinib treatment. Furthermore, fewer patients in the ibrutinib arm experienced MRD relapse, with approximately 60% of patients maintaining undetectable MRD at a median follow-up of 2 years after first attaining undetectable MRD status. Due to the small numbers of patients in the subgroups, the association between MRD status and PFS was not further studied.

Few patients discontinued treatment during the additional 13 months of follow-up after the primary analysis (n=14), and no new safety signals were reported with ongoing ibrutinib treatment. This is consistent with prior reports that rates of most adverse events are highest during the first year of ibrutinib-based treatment and decrease in frequency thereafter. Most patients had improvement or resolution of adverse events following dose reduction, suggesting that many such events are managed effectively by dose modification, allowing patients to stay on therapy and continue to maintain disease control. The discontinuation rate due to adverse events (22%) is comparable to that for first-line, single-agent ibrutinib after 4 years of follow-up in the RESONATE-2 study (19%) and is supported by real-world studies showing that treatment discontinuation is similar in patients with CLL/SLL receiving single-agent ibrutinib or ibrutinib-based combination therapy.

Overall, first-line ibrutinib plus obinutuzumab was well tolerated, which is important given that the median age at diagnosis of CLL is over 70 years; a demographic that often presents with comorbidities that preclude the use of chemotherapy-containing regimens. In patients receiving concomitant medications, ibrutinib was well tolerated. Notably, ibrutinib can be co-administered with acid-reducing medications, offering an advantage over other kinase inhibitors known to have drug drug interactions with acid-reducing agents. Overall, these results strengthen and build upon the broad experience with ibrutinib. Several studies have assessed whether adding anti-CD20 therapy to ibrutinib results in a clinically relevant improvement in efficacy in CLL. Evidence suggests that addition of anti-CD20 therapy to ibrutinib may increase depth of response and decrease the time to absolute lymphocyte count normalization relative to single-agent ibrutinib therapy, although survival outcomes have not been significantly improved over the impressive results with single-agent ibrutinib. The lack of data showing quality of life or survival outcome benefits notwithstanding, the addition of an anti-CD20 agent may nonetheless provide reassurance for patients concerned with increased absolute lymphocyte count. In a cross-trial comparison of RESONATE-2 and iLUMINATE, patients treated with ibrutinib plus obinutuzumab combination therapy had higher CR/CRI rates (44% vs. 27%; P=0.006) and shorter time to absolute lymphocyte count normalization (8 vs. 55 weeks) than patients treated with single-agent ibrutinib. Compared to single-agent ibrutinib, the combination of ibrutinib plus rituximab was demonstrated by Burger et al. to improve CR/CRI rates (particularly in the first-line setting: 50% vs. 20%) and shortened time to absolute lymphocyte count normalization (24 vs. 48 weeks) in a phase II study with a median follow-up of 36 months. In the phase III ALLIANCE 041202 study, the CR rates (12% and 7%) and rates of undetectable MRD (4% and 1%) were slightly higher with ibrutinib plus rituximab than with single-agent ibrutinib. The difference in response rates tended to be greater in patients with high-risk disease, suggesting that combination therapy may be most useful for patients with high-risk or bulky disease. In the relapsed/refractory high-risk setting, the GENUINE study demonstrated significantly higher ORR with ublituximab plus ibrutinib compared to single-agent ibrutinib after a median follow-up of 41.6 months (83% vs. 65%; P=0.02). In the setting of other BTK inhibitors, the ELEVATE-TN study reported investigator-assessed CR/CRI rates of 24% with acalabrutinib plus obinutuzumab compared to 8% for acalabrutinib monotherapy and 13% for obinutuzumab plus chlorambucil. In terms of PFS, 48-month estimates between RESONATE-2 and the current study were similar across ibrutinib-based treatment arms (76% vs. 74%), although it is worth noting that the current study included patients with high-risk genomic features whereas RESONATE-2 excluded those with del17p. Regardless, these broad clinical data clearly refute preclinical hypotheses of ibrutinib negating anti-CD20 efficacy, although anti-CD20 use may be important to optimize efficacy of other BTK inhibitors. In conclusion, ibrutinib plus obinutuzumab remains an effective chemotherapy-free regimen for patients with CLL/SLL that provides sustained efficacy and significantly reduces the risk of disease progression or death compared with chlorambucil plus obinutuzumab, including in patients with high-risk genomic features. With ongoing once-daily dosing, long-term ibrutinib therapy was well tolerated with no new safety signals observed.

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**Contributions**
As members of the Steering Committee, CM, DG, JGG, and IWF collaborated with the study sponsors to design the study and protocol; CM, RG, RD, AT, BA, LL, MS, JN, VS, DG, JGG, and IWF collected the study data; SD analyzed the data; KK collected and tested the high-risk genomics data for the study; SD, EH, and JD confirmed the accuracy of the data and compiled it for analysis. All authors had access to the data and were involved in the interpretation of data, contributed to the manuscript review and revisions, and approved the final version for submission.

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**Data-sharing statement**
Requests for access to individual participant data from clinical studies conducted by Pharmacyclics LLC, an AbbVie Company, can be submitted through Yale Open Data Access (YODA) Project site at http://yoda.yale.edu.
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