Final analysis from RESONATE: Up to six years of follow-up on ibrutinib in patients with previously treated chronic lymphocytic leukemia or small lymphocytic lymphoma


1Department of Haematology, St. James’s University Hospital, Leeds, UK
2CLL Center, Dana-Farber Cancer Institute, Boston, Massachusetts
3UC Irvine, Chao Family Comprehensive Cancer Center, Irvine, California
4Division of Medical Oncology and Hematology Northwell Health Cancer Institute, Lake Success, New York
5Wilmot Cancer Institute, University of Rochester Medical Center, Rochester, New York
6Vanderbilt-Ingram Cancer Center, Nashville, Tennessee
7Stanford Cancer Center, Stanford University School of Medicine, Stanford, California
8Peter MacCallum Cancer Centre, St. Vincent’s Hospital and University of Melbourne, Melbourne, Australia
9Royal North Shore Hospital, Sydney, Australia
10Division of Hematology and Hemostaseology, Medical University of Vienna, Wien, Austria
11UCSD Moores Cancer Center, San Diego, California
12Hospital de la Santa Creu i Sant Pau, Autonomous University of Barcelona, Barcelona, Spain
13Niguarda Ca’ Granda Hospital, Milan, Italy
14Department of Leukemia, University of Texas MD Anderson Cancer Center, Houston, Texas
15The Ohio State University Comprehensive Cancer Center, Columbus, Ohio
16The Leeds Teaching Hospitals, St. James Institute of Oncology, Leeds, UK
17Pharmacyclics LLC, an AbbVie Company, Sunnyvale, California

Correspondence
Talha Munir, Department of Haematology, St. James’s Hospital, Beckett St, Leeds, UK LS9 7TF.
Email: tmunir@nhs.net

Funding information
Janssen Pharmaceuticals; Pharmacyclics LLC, an AbbVie Company

Abstract
Ibrutinib, a once-daily oral inhibitor of Bruton’s tyrosine kinase, is approved in the United States and Europe for treatment of patients with chronic lymphocytic leukemia (CLL) or small lymphocytic lymphoma (SLL). The phase 3 RESONATE study showed improved efficacy of single-agent ibrutinib over ofatumumab in patients with relapsed/refractory CLL/SLL, including those with high-risk features. Here we report the final analysis from RESONATE with median follow-up on study of 65.3 months (range, 0.3-71.6) in the ibrutinib arm. Median progression-free survival (PFS) remained significantly longer for
patients randomized to ibrutinib vs ofatumumab (44.1 vs 8.1 months; hazard ratio [HR]: 0.148; 95% confidence interval [CI]: 0.113-0.196; P<.001). The PFS benefit with ibrutinib vs ofatumumab was preserved in the genomic high-risk population with del(17p), TP53 mutation, del(11q), and/or unmutated IGHV status (median PFS 44.1 vs 8.0 months; HR: 0.110; 95% CI: 0.080-0.152), which represented 82% of patients. Overall response rate with ibrutinib was 91% (complete response/complete response with incomplete bone marrow recovery, 11%). Overall survival, censored for crossover, was better with ibrutinib than ofatumumab (HR: 0.639; 95% CI: 0.418-0.975). With up to 71 months (median 41 months) of ibrutinib therapy, the safety profile remained consistent with prior reports; cumulatively, all-grade (grade ≥3) hypertension and atrial fibrillation occurred in 21% (9%) and 12% (6%) of patients, respectively. Only 16% discontinued ibrutinib because of adverse events (AEs). These long-term results confirm the robust efficacy of ibrutinib in relapsed/refractory CLL/SLL irrespective of high-risk clinical or genomic features, with no unexpected AEs. This trial is registered at www.clinicaltrials.gov (NCT01578707).

1 | INTRODUCTION

Chronic lymphocytic leukemia (CLL) and small lymphocytic lymphoma (SLL) are heterogeneous diseases with outcomes that are affected by clinical status and genetic aberrations. Patients with relapsed CLL/SLL and known high-risk factors, such as 17p deletion (del[17p]), TP53 aberrations (deletion/mutation), 11q deletion (del[11q]), or unmutated immunoglobulin heavy chain variable region (IGHV) gene, have a poor prognosis, and recent International Workshop on Chronic Lymphocytic Leukemia (IWCLL) guideline updates recommend testing for these high-risk factors to aid in treatment decisions. Chemoimmunotherapy has been at the forefront of treatment of CLL/SLL in the past decade, resulting in improved progression-free survival (PFS) and overall survival (OS) outcomes for untreated patients requiring therapy. Additionally, treatment of relapsed CLL/SLL with combination chemoimmunotherapy resulted in modest improvements in PFS and, in some cases, OS. In the past, however, once patients became refractory to chemoimmunotherapy, especially those with early relapse or high-risk genomic features, treatment options were limited and survival time was short.

The introduction of targeted therapy to the armamentarium of CLL/SLL therapy has dramatically improved treatment outcomes in patients refractory to chemoimmunotherapy. Ibrutinib, a first-in-class, once-daily, covalent inhibitor of Bruton’s tyrosine kinase (BTK; a B-cell receptor signaling kinase expressed by various hematopoietic cells, B-cell lymphomas, and leukemias), was one such therapy introduced in this relapsed and refractory CLL/SLL population that has since revolutionized therapy. RESONATE was the first multicenter, open-label, randomized phase 3 study that compared ibrutinib to ofatumumab treatment outcomes in previously treated patients with CLL/SLL, leading to full FDA approval of ibrutinib (after prior accelerated approval), including in patients with del(17p). The analysis from the first report of this trial demonstrated that ibrutinib significantly improved PFS, OS, and overall response rates (ORR) compared with ofatumumab and was generally well tolerated. RESONATE was also the first randomized trial that identified adverse events (AEs) associated with ibrutinib, such as atrial fibrillation and bruising, that have been followed long-term with BTK inhibitors. Herein, we report the final long-term follow-up efficacy and safety results of this landmark study, including analyses of subgroups with high-risk genomic features.

2 | METHODS

2.1 | Patients

Detailed eligibility criteria have been previously published. Briefly, patients with active relapsed/refractory CLL or SLL with measurable nodal disease by computed tomography (CT) who failed ≥1 prior line of therapy and met the International Workshop on Chronic Lymphocytic Leukemia (IWCLL) 2008 guidelines for requiring therapy, but were not candidates for purine analog therapy (see Supplementary Methods), were eligible for inclusion in the study. All patients provided written informed consent. The study was approved by the institutional review board or independent ethics committee at each participating institution and was conducted in accordance with the Declaration of Helsinki and International Conference on Harmonisation Guideline for Good Clinical Practice. The study was sponsored by Pharmacyclics LLC, an AbbVie Company, and Janssen Research and Development LLC.

2.2 | Study Design

RESONATE was a randomized, multicenter, open-label, phase 3 study that evaluated the efficacy and safety of ibrutinib compared with ofatumumab in previously treated patients with relapsed or refractory CLL/SLL (clinicaltrials.gov number NCT01578707). Enrollment was from June 2012 to April 2013. Patients were randomly assigned 1:1
to receive either oral ibrutinib 420 mg once daily (until disease progression or unacceptable toxicity) or intravenous ofatumumab for up to 24 weeks at an initial dose of 300 mg at week one, followed by a dose of 2000 mg weekly for seven weeks and then every four weeks for 16 weeks. Stratification was by purine analog refractory status (defined as no response to or relapse within 12 months of last dose of purine analog) and presence or absence of del(17p). Based on durable responses in the ongoing pivotal phase 2 study PCYC-1102, a protocol amendment was instituted in August 2013 to allow patients on the ofatumumab arm to cross over to receive ibrutinib. Following study closure, eligible patients could continue to receive ibrutinib treatment in a long-term extension study (PCYC-1145-LT; clinicaltrials.gov number NCT03229200).

2.3 | Assessments

PFS per investigator assessment was based on IWCLL 2008 criteria, with the 2012 clarification that treatment-related lymphocytosis was not considered progressive disease (PD) if in the setting of improvement in other disease parameters. Other endpoints were ORR per investigator assessment (defined as the proportion of patients who achieved complete response [CR], CR with incomplete bone marrow recovery [CRi], nodular partial response, partial response, or partial response with lymphocytosis). OS, and safety. Functional Assessment of Chronic Illness Therapy-Fatigue (FACIT-F) was a secondary endpoint; other patient-reported outcomes were exploratory endpoints. Methods for monitoring patients during follow-up have been described and were continued through long-term follow-up. Details of assessments, including patient-reported outcomes and cytogenetics, are provided in the Supplementary Methods.

2.4 | Statistical methods

Long-term efficacy, including PFS, OS, and ORR, were assessed in the intent-to-treat population. PFS and ORR were based on investigator assessment (independent review committee assessments were performed only for the primary analysis). PFS and OS (with and without censoring at crossover for ofatumumab patients) were analyzed using Kaplan-Meier methodology and hazard ratio (HR) was estimated using Cox regression model. In addition, the rank-preserving structural failure time (RPSFT) randomization-based model was employed to estimate the OS HR using counterfactual survival times that would have been observed in the absence of the extensive crossover. Ofatumumab treatment was completed at the time of the primary analysis and reported in the previous report. Long-term safety was summarized by yearly interval for ibrutinib patients.

2.5 | 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.

3 | RESULTS

3.1 | Patient features

Baseline characteristics of patients randomized to ibrutinib (n = 195) or ofatumumab (n = 196) are shown in Table S1. Similar proportions of patients in the ibrutinib and ofatumumab arms had high-risk features, including del(17p) (32% and 33%, respectively), mutated TP53 (51% and 46%, respectively), del(11q) (33% and 31%, respectively), unmutated IGHV (73% and 63%, respectively), and complex karyotype (25% and 22%, respectively) (Table S1). The majority of patients in both the ibrutinib and ofatumumab arms (86% and 79%, respectively) comprised the high-risk population, defined as having any of the following: del(17p), TP53 mutation, del(11q), and/or unmutated IGHV status.

The final analysis was performed upon study closure. Median follow-up on the study of patients initially assigned to ibrutinib was 65.3 months (range: 0.3–71.6), and median duration of ibrutinib therapy was

![Table 1](https://example.com/table1.png)

**Table 1** Patient disposition and treatment exposure during study treatment

<table>
<thead>
<tr>
<th>Parameter, n (%)</th>
<th>Ibrutinib (n = 195)</th>
<th>Ofatumumab (n = 196)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Duration of treatment, months, median (range)</td>
<td>41.0 (0.2-71.1)</td>
<td>5.3 (0.0-9.0)</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Disposition of study treatment</th>
<th>Ibrutinib</th>
<th>Ofatumumab</th>
</tr>
</thead>
<tbody>
<tr>
<td>Did not receive study drug</td>
<td>0</td>
<td>5 (2.6)</td>
</tr>
<tr>
<td>Completed treatment (ofatumumab arm only)</td>
<td>N/A</td>
<td>120 (61.2)</td>
</tr>
<tr>
<td>Discontinued</td>
<td>195 (100.0)</td>
<td>71 (36.2)</td>
</tr>
<tr>
<td>Progressive disease</td>
<td>72 (36.9)</td>
<td>36 (18.4)</td>
</tr>
<tr>
<td>Study terminated by sponsor</td>
<td>43 (22.1)</td>
<td>0</td>
</tr>
<tr>
<td>Adverse event</td>
<td>32 (16.4)</td>
<td>7 (3.6)</td>
</tr>
<tr>
<td>Patient withdrawal</td>
<td>15 (7.7)</td>
<td>6 (3.1)</td>
</tr>
<tr>
<td>Deatha</td>
<td>13 (6.7)</td>
<td>9 (4.6)</td>
</tr>
<tr>
<td>Investigator decision</td>
<td>20 (10.3)</td>
<td>13 (6.6)</td>
</tr>
</tbody>
</table>

Duration of treatment, years (randomized therapy)³

| >0 to 1 | 36 (18.5) | 191 (97.4) |
|>1 to 2 | 25 (12.8) | 0 |
|>2 to 3 | 31 (15.9) | 0 |
|>3 to 4 | 24 (12.3) | 0 |
|>4 to 5 | 22 (11.3) | 0 |
|>5 to 6 | 57 (29.2) | 0 |

³These cases of death included the following: ibrutinib arm, pneumonia (n = 3), sepsis (n = 2), unknown cause (sudden death, n = 2), neutropenic sepsis, terminal bowel cancer, lung infection, cardiac arrest, subdural hematoma, and burns and ensuing complications in one patient each; ofatumumab arm, pneumonia (n = 2), upper respiratory tract infection, squamous cell carcinoma of the neck, influenza A, aggressive squamous cell carcinoma of the scalp, nocardiosis, fever of unknown origin, and bacteremia in one patient each.

³Planned duration of treatment with ofatumumab was up to 24 weeks.
41 months (range: 0.2-71.1), with 41% of patients receiving more than four years of therapy (Table 1). Among patients initially assigned to ibrutinib, 43/195 (22%) remained on therapy until study closure (Table 1 and Figure S1). The most common reasons for ibrutinib discontinuation prior to study closure were PD (72/195, 37%) and AEs (32/195, 16%). Progressive disease was evidenced as Richter’s transformation in 20 patients (10%) in the ibrutinib arm, occurring in eight, four, six, zero, and two patients in years zero to one, one to two, two to three, three to four, and for five, respectively.

Median follow-up of the 196 patients initially assigned to the ofatumumab arm was 65.6 months (range: 0.1-73.9) and median duration of ofatumumab treatment was 5.3 months (range: 0-9.0). In total, 133 of 196 patients (68%) crossed over to receive ibrutinib; 36% of these patients (48/133) had received more than 4 years of next-line ibrutinib therapy, and 35% (47/133) were still receiving ibrutinib at the time of study closure. In addition, 15 of 196 patients (8%) had PD but exited the study on or before the protocol amendment allowing for crossover to receive ibrutinib.

3.2 | Progression-free survival

With an overall follow-up of 74 months, the median investigator assessed PFS per IWCLL criteria was 44.1 months (95% confidence interval [CI]: 38.5-56.2) in the ibrutinib arm and 8.1 months (95% CI: 7.8-8.3) in the ofatumumab arm. Similar to the prior analyses, PFS was significantly longer for patients assigned to ibrutinib than for patients assigned to ofatumumab (HR: 0.148; 95% CI: 0.113-0.196; P<0.0001; Figure 1A). At the time of this analysis, 92% (180/196) of patients initially assigned to ofatumumab had a PFS event. At the 60-month landmark, the estimated PFS rates were 40% in the ibrutinib arm and 3% in the ofatumumab arm. Median PFS in patients in the high-risk population (with del[17p], TP53 mutation, del[11q], and/or unmutated IGHV status) was 44.1 months (95% CI: 38.5-56.9) with ibrutinib and 8.0 months (95% CI: 6.4-8.2) with ofatumumab (HR: 0.110; 95% CI: 0.080-0.152) (Figure 1B). The PFS benefit of ibrutinib vs ofatumumab was consistent across subgroups defined by patient baseline clinical and genomic risk factors; HRs ranged from 0.042 to 0.306 (Figure 1C).

**FIGURE 1** Progression-free survival (A) in the ITT population and (B) in the high-risk population (patients with del[17p], TP53 mutation, del[11q], and/or unmutated IGHV status). (C) Forest plot of HRs for progression-free survival by baseline subgroups (ITT population). CI, confidence interval; ECOG, Eastern Cooperative Oncology Group; IGHV, immunoglobulin heavy-chain variable region gene; ITT, intention-to-treat; PFS, progression-free survival.

*Patients with del[17p], TP53 mutation, del[11q], and/or unmutated IGHV status
Within the ibrutinib arm, median PFS by line of therapy was not reached in patients who received 1 prior line of therapy (n = 35; 95% CI: 44.4-not estimable [NE]) and was 67.3 (n = 57; 95% CI: 36.0-NE), 44.1 (n = 32; 95% CI: 25.4-NE), 33.0 (n = 27; 95% CI: 13.6-NE), and 27.3 months (n = 44; 95% CI: 22.0-40.8) for patients who received two, three, four, or five or more prior lines of therapy, respectively (Figure 2A).

![Graph A](image1)

**Figure 2**  Progression-free survival by lines of therapy and genomic subgroups in the ibrutinib arm (ITT population). (A) Analysis by number of prior lines of therapy. (B) Analysis by del(17p) and del(11q).* (C) Analysis by CK. (D) Analysis by IGHV mutation status. (E) Analysis by TP53 mutation status. (F) Analysis by del(17p) and/or TP53 mutation status. CI, confidence interval; CK, complex karyotype; FISH, fluorescence in situ hybridization; IGHV, immunoglobulin heavy-chain variable region gene; NE, not estimable; NR, not reached; PFS, progression-free survival.

*Genomic abnormalities by FISH cytogenetics were categorized according to Döhner hierarchical classification.
Purine analog refractoriness had no impact on PFS, with median PFS of 44.6 (n = 86; 95% CI: 31.7-57.4) and 44.0 months (n = 108; 95% CI: 35.1-60.7) in patients who were and were not refractory at study enrollment, respectively (HR: 0.983; 95% CI: 0.685-1.413).

Median PFS was 40.6 months (95% CI: 25.4-44.6) in the del(17p) subgroup (n = 63) and 60.7 months (95% CI: 36.4-NE) in the del(11q) subgroup (n = 50; Figure 2B). Median PFS was 42.5 months (95% CI: 31.7-56.2) in patients with neither del(11q) nor del(17p) (n = 81). In an exploratory analysis that combined data from patients with del(17p) and those with TP53 mutation, median PFS was 40.6 months (95% CI: 27.5-44.1) in the subgroup with del(17p) and/or TP53 mutation (n = 104) and 56.9 months (95% CI: 27.5-NE) in the del(11q) subgroup (n = 33) (Figure S2). Median PFS was not reached (95% CI: 42.5-NE) in patients with neither del(11q) nor del(17p) (n = 58). Patients with complex karyotype (n = 39) had a median PFS of 40.8 months (95% CI: 24.4-67.7) and patients without a complex karyotype (n = 114) had a median PFS of 44.6 months (95% CI: 37.9-61.0) (HR: 1.279; 95% CI: 0.801-2.044) (Figure 2C). Patients with mutated (n = 36) and unmutated (n = 98) IGHV had similar PFS (median 48.4 [95% CI: 35.6-60.8] vs 49.7 [95% CI: 40.2-NE] months; HR: 1.208; 95% CI: 0.741-1.971) (Figure 2D).

Patients without TP53 mutation (n = 75) had a median PFS of 56.9 months (95% CI: 36.4-NE), longer than the median PFS of 40.7 months (95% CI: 25.4-44.6) for patients with TP53 mutations (n = 79) (HR: 1.731; 95% CI: 1.156-2.593) (Figure 2E). Median PFS was 40.6 months (95% CI: 17.6-56.2) for patients with both del(17p) and TP53 mutation (n = 38), 40.7 months (95% CI: 25.4-57.3) for those with either del(17p) or TP53 mutation (n = 48), and 56.9 months (95% CI: 36.4-NE) for those with neither del(17p) nor TP53 mutation (n = 68; Figure 2F).

### 3.3 Overall response

At extended follow-up, the cumulative ORR was 91% for ibrutinib. The proportion of patients with a best response of CR/CRI increased over time to 11% with current follow-up (Figure S3).

### 3.4 Overall survival

With up to 6 years of post-randomization follow-up, median OS was 67.7 months (95% CI: 61.0-NE) in the ibrutinib arm and 65.1 months (95% CI: 50.6-NE) in the ofatumumab arm, irrespective of the extensive (68%) crossover to ibrutinib (HR: 0.810; 95% CI: 0.602-1.091) (Figure S4A). To compare OS outcomes between treatment arms accounting for the extensive crossover from ofatumumab to ibrutinib treatment, OS was analyzed with censoring at time of crossover and was generally decreased over time for patients remaining on ibrutinib therapy, with few exceptions (hypertension and bruising). Over up to 71 months of treatment, commonly reported grade ≥3 hematologic AEs included neutropenia (25%), thrombocytopenia (10%), and anemia (9%). Commonly reported grade ≥3 nonhematologic AEs included pneumonia (21%), hypertension (9%), urinary tract infection (7%), diarrhea (7%), and atrial fibrillation (6%). Overall, 19 patients (10%) in the ibrutinib arm experienced major hemorrhage. The prevalence of any grade ≥3 AEs with ibrutinib decreased after the first year and remained stable thereafter, with rates of 62%, 48%, 46%, 46%, 48%, and 32% during years zero to one, one to two, two to three, three to four, four to five, and five to six, respectively.

In line with the overall prevalence, the prevalence of most grade ≥3 AEs of clinical interest with ibrutinib decreased over time (Figure 3). With up to 6 years of follow-up, grade ≥3 infections occurred in 87 patients (45%) in the ibrutinib arm (Figure S7), occurring at a median time of onset of 7.0 months (range: 0.0 to 63.0). The most frequent categories/types of grade ≥3 infections included pneumonia (21%), urinary tract infection (7%), cellulitis (4%), sepsis (4%), and lung infection (4%). The prevalence of grade ≥3 atrial fibrillation was 4%, 0%, 2%, 2%, 1%, and 0% over years zero to one, one to two, two to three, three to four, four to five, and five to six, respectively (Figure 3). The prevalence of all-grade atrial fibrillation was 6%, 5%, 8%, 8%, 4%, and 4% over years zero to one, one to two, two to three, three to four, four to five, and five to six, respectively. Overall,
FIGURE 3  Prevalence of grade ≥3 AEs of clinical interest over time for ibrutinib arm (ITT population). Prevalence was determined by the proportion of patients with a given AE (existing event or new onset of an event) during each yearly interval. Multiple onsets of the same AE term within a specific yearly interval were counted once, and the same AE term continuing across several yearly intervals was counted in each of the intervals. Congestive heart failure and peripheral neuropathy were defined per terms included in the SMQ categories. AE, adverse event; ITT, intention-to-treat; RBC, red blood cells.

24 patients (12%) experienced atrial fibrillation of any grade, with a median duration of 10.0 days (range: 1-1323) for the events (Table S2). Most patients who experienced atrial fibrillation had one or more relevant risk factors (22/24, 92%), such as a history of cardiovascular disease (10/24, 42%), hypertension (9/24, 38%), and/or baseline body mass index >25 kg/m² (11/24, 46%). Most received concomitant medications (19/24, 79%), including antithrombotic medications in eight of 24 patients (33%). The prevalence of grade ≥3 hypertension was 4%, 8%, 7%, 7%, 8%, and 11% over years zero to one, one to two, two to three, three to four, four to five, and five to six, respectively (Figure 3). A total of 41 patients (21%) experienced hypertension of any grade (Figure S8), occurring at a median time of onset of 13.8 months (range: 1.0-54.0) and a median duration of 16.6 months (range: 0.1-64.5) for the events (Table S3). Most patients who experienced hypertension had relevant risk factors (28/41, 68%) of either a history of hypertension or related disorders (21/41, 51%) or a history of other risk factors, such as diabetes mellitus, hyperlipidemia, renal disease, smoking, or thyroid disease (19/41, 46%). Most patients who experienced hypertension were treated with concomitant antihypertensive medication during the treatment-emergent period (26/41, 63%). Other AEs of clinical interest with ibrutinib included peripheral neuropathy (all grade: n = 26 [13%]; grade ≥3: n = 1 [0.5%]), congestive heart failure (all grade: n = 9 [5%]; grade ≥3: n = 5 [3%]), and ventricular arrhythmia (all grade: n = 2 [1%]; no grade ≥3 events).

Among patients in the ibrutinib arm, 17% (34/195) and 65% (126/195) of patients required dose reductions and dose holds, respectively, to manage AEs. Among the 34 patients with dose reductions due to AEs, eight (24%) ultimately discontinued because of an AE, of which 4 discontinuations (12%) were due to the same AE that led to dose reduction. Of the 126 patients with a dose held because of AEs, 20 patients (16%) ultimately discontinued ibrutinib because of an AE, of which 13 discontinuations (10%) were due to the same AE that led to the dose hold. The frequency of AEs that led to ibrutinib discontinuation remained consistent over time, and occurred in 6%, 3%, 4%, 4%, 6%, and 4% of patients during years zero to one, one to two, two to three, three to four, four to five, and five to six, respectively. Over the course of follow-up with a median treatment duration of 41 months, 32 of 195 patients (16%) discontinued ibrutinib due to AEs, including 11 grade 5 events of sepsis (n = 3), pneumonia (n = 2), unknown cause (sudden death, n = 2), lower respiratory tract infection (n = 1), lung infection (n = 1), cardiac arrest (n = 1), terminal bowel cancer (n = 1), subdural hematoma (n = 1), and serious burn and ensuing complications (n = 1).

3.7 | Response to next-line treatment following ibrutinib discontinuation

Best response to next-line CLL therapy was reported for 27 patients after discontinuation of ibrutinib randomized therapy. Reasons for discontinuation in these patients included PD (n = 22), AE (n = 3), and investigator decision (n = 2). Responses were observed in 10 of these 27 patients upon next-line treatment, including patients who had discontinued ibrutinib due to PD (eight of 22), AE (one of three), and physician decision (one of two). Next-line treatment regimens with responders were venetoclax single agent (five responders of seven treated), idelalisib + rituximab (two of six), high-dose methylprednisolone + monoclonal antibody (one of one, alemtuzumab), and investigational agent(s) (two of three).

4 | DISCUSSION

Targeted therapy with ibrutinib for relapsed CLL has changed the natural history of this disease, as first demonstrated in the trial described herein. This 6-year update of the RESONATE study confirmed the robust efficacy of ibrutinib in patients with relapsed or refractory CLL. The median PFS of 44.1 months in the ibrutinib arm is substantially superior to the median PFS in the comparator arm of ofatumumab (8.1 months) and to that of other treatments available at the time of study initiation. The efficacy of ibrutinib is confirmed in patients with...
high-risk clinical and genomic features, including TP53 alteration (deletion/mutation), unmutated IGHV status, complex karyotype, and ATM aberrations (as shown by del(11q)). Analysis of OS is confounded by the high number of patients (n = 133) who crossed over to receive ibrutinib, with nonprogressing ofatumumab arm patients having the most indolent disease, and with ibrutinib use prolonging survival of the patients who crossed over. Despite this, when censored for crossover, OS was significantly improved in patients treated with ibrutinib although the difference in the median OS for ibrutinib vs ofatumumab was small (2.6 months). While a larger difference was observed using the RPSFT method, the analysis was confounded by the design of the study, which permitted early and extensive crossover, and the underlying assumption of this method that there is a constant risk of death throughout the study for both treatment arms. As expected, analysis of OS outcomes by timing of relapse suggests that patients who experienced PD earlier in the course of ibrutinib treatment (within the first one or two years) had shorter OS, as compared to those who experienced PD at any time in the first four years. Ibrutinib was well tolerated; the main reason for treatment discontinuation in this extended follow-up was PD rather than drug-related toxicity. The frequency and timing of Richter's transformation in ibrutinib-treated patients in the RESONATE study are consistent with emerging data from clinical studies with novel agents and historical data with chemotherapy-based regimens. Notably, the patient population in the RESONATE study was characterized by a high prevalence of factors previously identified as associated with increased risk of Richter's transformation, including genomic risk factors and heavy prior treatment burden. Most cases of Richter's transformation were identified early in the course of ibrutinib treatment, with almost half of cases occurring in the first year and all but two of the remaining cases occurring during years 2 and 3.

Studies have shown that patients with del(17p) CLL treated with standard chemioimmunotherapy regimens have poor outcomes. The 5-year follow-up data from the relapsed/refractory cohort of the phase 2 PCYC-1102/1103 study with single-agent ibrutinib showed a lower PFS rate in patients with del(17p) than in patients without del(17p). The 3-year follow-up of the RESONATE study reported the effectiveness of ibrutinib in patients regardless of del(17p). In this final analysis of RESONATE with 6 years of follow-up, we confirm no significant difference in PFS for ibrutinib-treated patients with or without del(17p). However, when data from patients with TP53 mutation were pooled with those with del(17p) in an exploratory analysis, we observed a lower PFS rate in the subgroup of patients with del(17p) and/or TP53 mutation than in patients without either of these abnormalities. Consistent with findings reported with earlier RESONATE follow-up, the subset of ibrutinib-treated patients with both del(17p) and TP53 mutation trended to have shorter PFS relative to patients without either of these high-risk features. In contrast to the earlier report from RESONATE, PFS outcomes in patients with either del(17p) or TP53 mutation (but not both) were similar to that of patients with both del(17p) and TP53 mutation. The small number of patients in these subgroups and the retrospective genetic/molecular analysis defining many of the subgroups may be potential limitations in the interpretation of these results. Development of acquired mutations in the B-cell receptor signaling pathway has been observed in patients relapsing during ibrutinib treatment and together with other as yet undefined mechanisms contributing to relapse may be associated with unstable genomes. Answers to these questions are being actively pursued in ongoing studies.

Unmutated IGHV and del(11q) have an adverse prognostic impact on patients with CLL. The negative impact from these markers has been substantially abrogated in the context of ibrutinib therapy, as shown in this study and other studies. This is confirmed by the current results from RESONATE in which patients with del(11q) CLL appeared to have no decrement in benefit from continuous ibrutinib therapy. This is reflected in the median PFS for this group (n = 50), which was 60.7 months and appeared to be longer than the median PFS of the overall ibrutinib cohort. In addition, median PFS with ibrutinib treatment was similar between patients with mutated and unmutated IGHV. Thus, our follow-up data suggest that IGHV mutational status and del(11q) have no impact on the long-term outcomes with ibrutinib treatment.

The ORR with ibrutinib remains high with longer follow-up (91%) and was similar to rates reported with a median follow-up of 44 months (91%),. The CR/CRi rates have increased over time with ibrutinib treatment, approaching a plateau beyond 3.5 years (9% and 11% with median 44 and 64 months of follow-up, respectively). This is consistent with deepening of response with continuous ibrutinib therapy for responding patients and is similar to the improved responses observed over time with longer follow-up in patients with relapsed/refractory CLL in the HELIOS study and in the phase 2 PCYC-1102/1103 study. Notably, the proportion of patients who reported clinically meaningful improvement in FACIT-F score was greater with ibrutinib than ofatumumab, and the improvement was maintained over time.

Assessment of long-term safety data is an important factor with continuous therapy, especially given the extended progression-free efficacy conferred by ibrutinib treatment. This update confirms that ibrutinib is well tolerated, and the prevalence of most grade ≥3 AEs of clinical interest decreased with each subsequent year of treatment, apart from hypertension, in which prevalence remained relatively consistent from year to year. During prolonged follow-up, the overall prevalence of treatment-emergent atrial fibrillation and hypertension of any grade were 12% and 21%, respectively. These rates were similar to those reported earlier in RESONATE with a median 44-month follow-up (11% and 20%, respectively). In addition, most of the patients who developed atrial fibrillation or hypertension had relevant risk factors for developing these complications. It is important to note that the incidence of major hemorrhage with continued treatment was consistently low over the extended follow-up period. In addition, substantial proportions of ibrutinib-treated patients who experienced dose reductions or dose holds due to AEs were able to restart or remain on ibrutinib treatment (15% and 25% of these patients, respectively, remained on ibrutinib until study closure). There
were no new or concerning AE trends or unexpected events noted with extended follow-up, and the safety profile appears consistent with earlier reports.

To conclude, these 6-year follow-up results from the RESONATE study confirm the robust and durable efficacy of ibrutinib with extended treatment in patients with relapsed or refractory CLL. The improvement in outcomes is evident in all risk groups, including patients with high-risk clinical and genomic features, and the safety analyses established that continuous treatment with ibrutinib can be delivered long-term to provide benefit to patients with CLL.

ACKNOWLEDGMENTS

The authors thank the patients who participated in this study and their supportive families, as well as the investigators, subinvestigators, and coordinators at each of the study sites. The authors thank Jerry Ping of Pharmacyclics LLC, an AbbVie Company, for his contributions to the statistical analyses and Joris Diels of Janssen and Suzy Van Sanden of Janssen-Cigli Ltd for their contributions to the crossover-adjusted overall survival analysis using the rank-preserving structural failure time method. John C. Byrd is supported by the Four Winds Foundation, D. Warren Brown Foundation, Mr. and Mrs. Michael Thomas, Sullivan CLL Research Foundation, and R35 CA197734. Editorial support was provided by Melanie Sweetlove, MSc, and was funded by Pharmacyclics LLC, an AbbVie Company.

CONFLICT OF INTEREST

T.M. has received honoraria from Janssen, AbbVie, Gilead, Alexion, Novartis, and Roche; and has been in a consulting role for MorphoSys and Sunesis. J.R.B. has received honoraria from Janssen and Teva; has been in a consulting role for AbbVie, Acerta, Astellas, BeiGene, Genentech/Roche, Gilead, Juno/Celgene, Kite, Loxo, Novartis, Pfizer, Pharmacyclics LLC, an AbbVie Company, Redx, Sun, Sunesis, TG Therapeutics, and Verastem; has received research funding from Gilead, Loxo, Sun, and Verastem; and has served on data safety monitoring committees for MorphoSys and Invecentx. S.O. has been in a consulting role for Amgen, Astellas, Celgene, GlaxoSmithKline, Janssen Oncology, Aptose Biosciences, Vaniam Group, AbbVie, Alexion, Verastem, and Eisai; has received research funding from Kite, Regeneron, and Acerta; and has been in a consulting role and received research funding from Gilead, Pharmacyclics LLC, an AbbVie Company, TG Therapeutics, Pfizer, and Sunesis. J.C.Ba. has received honoraria from Janssen; has been in a consulting role for Genentech, Gilead, Bayer, Pharmacyclics LLC, an AbbVie Company, AbbVie, AstraZeneca, and Sandoz; and has received research funding from Pharmacyclics LLC, an AbbVie Company, Oncertnal Therapeutics, and AbbVie. P.M.B. has been in a consulting role for Pharmacyclics LLC, an AbbVie Company, and AbbVie, and has received research funding from Pharmacyclics LLC, an AbbVie Company. N.M.R. has received honoraria from and has been in a consulting role for Bristol-Myers Squibb (BMS), Genentech, AbbVie, Gilead, AstraZeneca, and Celgene; has received research funding from BMS and Merck; and has been a paid speaker for Gilead and Celgene. S.C. has received honoraria from Pharmacyclics LLC, an AbbVie Company, and Janssen; has been in a consulting role for AbbVie, Celgene, Genentech, Pharmacyclics LLC, an AbbVie Company, Janssen, Novartis, Astellas, and AstraZeneca; has received research funding from AbbVie, Acerta, Celgene, Gilead, Janssen, Pharmacyclics LLC, an AbbVie Company, and Takeda; has provided expert testimony for Genentech; has received travel, accommodations and other expenses from AbbVie, BeiGene, Celgene, Genentech, Janssen, and Pharmacyclics LLC, an AbbVie Company; and has other relationships with BeiGene. C.S.T. has received honoraria from Janssen and Pharmacyclics LLC, an AbbVie Company, and research funding from Janssen. S.P.M. has received honoraria from and has been in a consulting role for Roche, AbbVie, Janssen, Gilead, and GlaxoSmithKline; has received research funding from Roche, AbbVie, and Janssen; and has been a paid speaker for Roche, AbbVie, Janssen, and Gilead. U.J. has received honoraria and research funding from Janssen, and has been in a consulting role for Janssen. T.J.K. has been in a consulting role for AbbVie, Genentech-Roche, Gilead, Pharmacyclics LLC, an AbbVie Company, and Celgene and has received research funding from AbbVie, Genentech-Roche, Pharmacyclics LLC, an AbbVie Company, and Oncertnal. C.M. has been in a consulting role for Pharmacyclics LLC, an AbbVie Company, Janssen, and AbbVie. M.M. has received honoraria from and has been in a consulting role for AbbVie, Janssen, Gilead, Roche, and AstraZeneca; has been a paid speaker for AbbVie, Janssen, and Gilead; has received research funding from Roche; and has received travel, accommodations, and expenses reimbursement from AbbVie, Janssen, and AstraZeneca. J.A.B. has received honoraria from Janssen; has been in a consulting role for Janssen; has received research funding from Gilead. TG Therapeutics, Pharmacyclics LLC, an AbbVie Company, and BeiGene; has been a paid speaker for Gilead, TG Therapeutics, Pharmacyclics LLC, an AbbVie Company, Novartis, and Janssen; and has received travel, accommodations, and expenses reimbursement from Gilead, TG Therapeutics, Pharmacyclics LLC, an AbbVie Company, Novartis, and Janssen. J.C.By. has been in a consulting role for Janssen; has reported research funding from Pharmacyclics LLC, an AbbVie Company, Gilead, TG Therapeutics, and BeiGene; and has been a paid speaker for Pharmacyclics LLC, an AbbVie Company, Gilead, TG Therapeutics, Novartis, and Janssen; and received travel, accommodations, and expenses reimbursement from Pharmacyclics LLC, an AbbVie Company, Gilead, TG Therapeutics, Novartis, and Janssen; and TG Therapeutics. P.H. has received honoraria from, has been in a consulting role for, and has received research funding from Janssen, Pharmacyclics LLC, an AbbVie Company, and AbbVie; and has received travel, accommodations, and expenses reimbursement from Janssen and AbbVie. S.D. is an employee of Pharmacyclics LLC, an AbbVie Company, and owns stock in AbbVie, Celgene, GlaxoSmithKline, and Exelixis. A.S. is an employee of Pharmacyclics LLC, an AbbVie Company, and owns stock in AbbVie. J.P.D. is an employee of Pharmacyclics LLC, an AbbVie Company, and owns stock in AbbVie and was formerly employed by and owned stock in CTI BioPharma. J.W. has been in a consulting role for Pharmacyclics LLC, an AbbVie Company, and Janssen;
and has received research funding from Pharmacyclics LLC, an AbbVie Company, Janssen, AbbVie, Morphosys, Karyopharm, and Loxo.

AUTHORSHIP CONTRIBUTIONS

J.C.By., J.R.B., and P.H. designed the study in collaboration with the study sponsor; T.M., J.R.B., S.O., J.C.Ba., P.M.B., N.M.R., S.C., C.S.T., S.P.M., U.J., T.J.K., C.M., M.M., J.A.B., J.C. Byrd, P.H., and J.W. collected data; S.D., A.S., and J.P.D. confirmed the accuracy of the data, interpreted the data, and compiled it for analysis; S.D. performed statistical analysis. All authors had access to the data and were involved in the interpretation of data. All authors contributed to the manuscript preparation and approved the final version of the manuscript for submission. T.M. wrote the first draft of the manuscript.

References

30. ESMO Guidelines Committee. eUpdate – Chronic lymphocytic leukaemia treatment recommendations. Available at: https://www.esmo.


SUPPORTING INFORMATION

Additional supporting information may be found online in the Supporting Information section at the end of this article.
Author/s:
Munir, T; Brown, JR; O'Brien, S; Barrientos, JC; Barr, PM; Reddy, NM; Coutre, S; Tam, CS; Muligan, SP; Jaeger, U; Kipps, TJ; Moreno, C; Montillo, M; Burger, JA; Byrd, JC; Hillmen, P; Dai, S; Szoke, A; Dean, JP; Woyach, JA

Title:
Final analysis from RESONATE: Up to six years of follow-up on ibrutinib in patients with previously treated chronic lymphocytic leukemia or small lymphocytic lymphoma

Date:
2019-10-13

Citation:

Persistent Link:
http://hdl.handle.net/11343/245222

File Description:
published version

License:
CC BY-NC-ND