Follow-up results of a phase II study of ibritumomab tiuxetan radioimmunotherapy in patients with relapsed or refractory low-grade, follicular, or transformed B-cell non-Hodgkin's lymphoma and mild thrombocytopenia

Russell Schilder  
*Fox Chase Comprehensive Cancer Center*

Arturo Molina  
*Biogen Idec*

Nancy Bartlett  
*Washington University School of Medicine in St. Louis*

Thomas Witzig  
*Mayo Clinic*

Leo Gordon  
*Northwestern University*

**Recommended Citation**

Schilder, Russell; Molina, Arturo; Bartlett, Nancy; Witzig, Thomas; Gordon, Leo; Murray, James; Spies, Stewart; Wang, Hua; Wiseman, Gregory; and White, Christine, ”Follow-up results of a phase II study of ibritumomab tiuxetan radioimmunotherapy in patients with relapsed or refractory low-grade, follicular, or transformed B-cell non-Hodgkin's lymphoma and mild thrombocytopenia.” Cancer Biotherapy & Radiopharmaceuticals. 19,4. 478-481. (2004).  
http://digitalcommons.wustl.edu/open_access_pubs/3129
See next page for additional authors

Follow this and additional works at: http://digitalcommons.wustl.edu/open_access_pubs
Authors
Russell Schilder, Arturo Molina, Nancy Bartlett, Thomas Witzig, Leo Gordon, James Murray, Stewart Spies, Hua Wang, Gregory Wiseman, and Christine White
Follow-Up Results of a Phase II Study of Ibritumomab Tiuxetan Radioimmunotherapy in Patients with Relapsed or Refractory Low-Grade, Follicular, or Transformed B-Cell Non-Hodgkin’s Lymphoma and Mild Thrombocytopenia

Russell Schilder,1 Arturo Molina,2 Nancy Bartlett,3 Thomas Witzig,4 Leo Gordon,5 James Murray,6 Stewart Spies,2 Hua Wang,2 Gregory Wiseman,4 and Christine White2

1Fox Chase Comprehensive Cancer Center, Philadelphia, PA
2Biogen Idec, San Diego, CA
3Washington University, St. Louis, MO
4Mayo Clinic, Rochester, MI
5Northwestern University, Chicago, IL
6M.D. Anderson Cancer Center, Houston, TX

ABSTRACT

This report presents updated time-to-event variables from a multicenter phase II trial of reduced-dose 90Y ibritumomab tiuxetan in patients with relapsed or refractory low-grade, follicular, or transformed B-cell non-Hodgkin’s lymphoma (NHL) and mild thrombocytopenia (platelet counts of 100 to 149 × 10^9 platelets/L). Patients received a single course of ibritumomab tiuxetan radioimmunotherapy, with 90Y ibritumomab tiuxetan administered at 0.3 mCi/kg (compared to a standard dose of 0.4 mCi/kg). In 30 patients, the overall response rate was 83%, with complete responses (confirmed [CR] and unconfirmed [CRu]) of 47%. Median follow-up time is currently 36.5 months (range: 7.5–54.9 months). Median duration of response was 11.5 months (range: 1.0–53.9 months), median time to progression was 9.4 months (range: 1.7–54.8 months), and median time to next lymphoma therapy was 14.6 months (range: 2.3–54.9 months). Median overall survival time has not yet been reached. Long-term responses, defined as time to progression of 12 months or greater, have been seen in 14 of 30 patients (47%) overall, and 12 of 14 CR/CRu patients (86%). Toxicities were primarily hematologic and reversible. No additional long-term adverse events have been observed in the follow-up period, and treatment did not preclude subsequent lymphoma therapies.

Key words: ibritumomab tiuxetan, radioimmunotherapy, non-Hodgkin’s lymphoma, therapy, thrombocytopenia

INTRODUCTION

The clinical course of low-grade or follicular B-cell non-Hodgkin’s lymphomas (NHL) is characterized by initial responsiveness to therapy and...
Radioimmunotherapy with ibritumomab tiuxetan is an effective treatment option for patients with relapsed or refractory low-grade, follicular, or transformed B-cell NHL. It was recognized early in the clinical development of ibritumomab tiuxetan that baseline platelet count, an indicator of bone-marrow reserve, correlated with the risk of hematologic toxicity. Early in the phase I/II study, investigators noted that patients with baseline mild thrombocytopenia treated with 90Y ibritumomab tiuxetan at a dose of 0.4 mCi/kg were at an increased risk of grade 4 hematologic toxicity, compared to patients with normal baseline platelet counts. Thereafter, patients with mild thrombocytopenia, defined as a baseline platelet count of 100 to 149 × 10^9 platelets/L, received a reduced dose of 90Y-ibritumomab tiuxetan of 0.3 mCi/kg. Results from this study were first reported in 2002. Patients received a single course of treatment, in which 90Y ibritumomab tiuxetan radioimmunotherapy in 30 patients with relapsed or refractory low-grade, follicular, or transformed B-cell NHL and mild thrombocytopenia (platelet counts of 100 to 149 × 10^9 platelets/L). Results from this study were first reported in 2002. Patients received a single course of treatment, in which 90Y ibritumomab tiuxetan was administered at a dose of 0.3 mCi/kg. The overall response rate was 83% using International Workshop response criteria. Toxicities were primarily hematologic, and were transient and reversible. This report provides updated data on response and survival outcomes from an additional 2 years of follow-up time.

**METHODS**

A detailed description of the design and methods of this single-arm, multicenter phase II study has been previously reported. Briefly, patients age 18 or above with histologically confirmed relapsed or refractory low-grade, follicular, or transformed CD20+ NHL and mild thrombocytopenia, defined as a baseline platelet count 100 to 149 × 10^9 platelets/L were eligible for enrollment. Patients had to have bidimensionally measurable disease, with at least 1 lesion measuring 2.0 cm or greater in a single dimension. Additional eligibility criteria included less than 25% bone marrow involvement; baseline World Health Organization (WHO) performance status of 0, 1, or 2; no prior myelofibrosis; and no prior radioimmunotherapy or anti-CD20 therapy. Patients had to have otherwise adequate hematologic and hepatic function and provide written, informed consent.

On day 1 of therapy, patients received an i.v. infusion of rituximab 250 mg/m^2 followed by an imaging dose of 111In ibritumomab tiuxetan 5 mCi (185 MBq). Provided biodistribution was adequate on gamma-camera images, approximately 1 week later patients received a second infusion of rituximab 250 mg/m^2, followed by 90Y ibritumomab tiuxetan 0.3 mCi/kg (11 MBq/kg), up to a maximum dose of 32 mCi (1.2 GBq).

**Statistical Design**

The primary study endpoint was overall response rate (ORR), with complete response (CR/CRu) rate, partial response (PR) rate, time to progression (TTP), duration of response (DR), and time to next NHL therapy (TTNT) evaluated as secondary endpoints. Response rates were determined using International Workshop NHL response criteria.

**RESULTS**

The study enrolled a total of 30 patients between May 1998 and August 1999, with a current fol-
Low-up through August 2003, for a median follow-up time of 36.5 months (range: 7.5–54.8 months). All 30 patients enrolled were evaluable for response and toxicity. Baseline patient characteristics are listed in Table 1. Patients had received a median of 2 prior therapies for NHL (range: 1–9). Nineteen (19) patients (63%) were resistant to at least 1 prior chemotherapy, and 14 patients (47%) were resistant to their last chemotherapy. Resistance was defined as a failure to achieve a complete or partial response, or disease progression, within 6 months.

Overall, 25 of 30 patients (83%) responded to therapy, with 11 CR (37%), 3 CRu (10%) and 11 PR (37%) by International Workshop response criteria.6 The median duration of response was 11.5 months (range: 1.0–53.9 months). The median time to progression for all patients was 9.4 months (range: 1.7–54.8 months) (Fig. 1). Overall, responses have been durable. Long-term responses, defined as time to progression of 12 months or greater, has been seen in 14 of 30 patients (47%) overall, including 12 of 14 patients (86%) who achieved a CR or CRu (the 2 patients who relapsed within 12 months were true CRs, not CRus). Median TTP in the 14 patients achieving long-term responses of 12 months or greater was 24.6 months (range: 12.1–54.8 months), and median TTP in all CR/CRu patients (independent of CR/CRu duration) was 24.5 months (range: 4.9–54.8 months). Two (2) patients had long-term partial responses of 12.6 and 17.3 months, respectively. These 2 patients had a residual mass on computed tomography (CT) scan that did not meet CRu criteria but was more likely fibrosis/scar than viable tumor, accounting for the long duration of response. Functional imaging with positron emission tomography (PET) was not done, but may have better defined these residual masses. Two (2) of the patients have remained in continuous, complete remission for 41 and 54.8 months, respectively, following ibritumomab tiuxetan radioimmunotherapy.

Median overall survival time has not yet been reached (data not shown). Overall median TTNT was 14.6 months (range: 2.3–54.8 months). A total of 24 patients received treatment subsequent to ibritumomab tiuxetan radioimmunotherapy, including rituximab, chemotherapy, and other therapies. In the 13 patients who received rituximab as their first NHL treatment following ibritumomab tiuxetan radioimmunotherapy, clinical responses were observed in 6 patients, 3 patients did not respond to therapy, and clinical outcome was unknown in the remaining 4 patients.

At this time, 7 of 30 patients enrolled in this study have died; 6 deaths were the result of disease progression, and 1 death was the result of other causes. In the original report, 1 patient with a history of chronic alkylator exposure developed acute myelogenous leukemia 20 months after

Table 1. Patient Characteristics

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>No.</th>
<th>%</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sex</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>18</td>
<td>60</td>
</tr>
<tr>
<td>Female</td>
<td>12</td>
<td>40</td>
</tr>
<tr>
<td>Median age (yrs.), range</td>
<td>61</td>
<td>(29–85)</td>
</tr>
<tr>
<td>Histology</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Small lymphocytic lymphoma</td>
<td>2</td>
<td>7</td>
</tr>
<tr>
<td>Follicular</td>
<td>25</td>
<td>83</td>
</tr>
<tr>
<td>Transformed</td>
<td>3</td>
<td>10</td>
</tr>
<tr>
<td>Disease stage</td>
<td></td>
<td></td>
</tr>
<tr>
<td>III</td>
<td>3</td>
<td>10</td>
</tr>
<tr>
<td>IIIv</td>
<td>27</td>
<td>90</td>
</tr>
<tr>
<td>Degree of bone marrow involvement</td>
<td></td>
<td></td>
</tr>
<tr>
<td>0 to &lt;5%</td>
<td>10</td>
<td>33</td>
</tr>
<tr>
<td>5% to &lt;20%</td>
<td>12</td>
<td>40</td>
</tr>
<tr>
<td>20% to &lt;25%</td>
<td>8</td>
<td>27</td>
</tr>
<tr>
<td>No. of extranodal disease sites</td>
<td></td>
<td></td>
</tr>
<tr>
<td>0 or 1</td>
<td>24</td>
<td>80</td>
</tr>
<tr>
<td>2 or more</td>
<td>6</td>
<td>20</td>
</tr>
<tr>
<td>Tumor size</td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt;5 cm</td>
<td>16</td>
<td>53</td>
</tr>
<tr>
<td>5 to &lt;7 cm</td>
<td>9</td>
<td>30</td>
</tr>
<tr>
<td>7 to &lt;10 cm</td>
<td>3</td>
<td>10</td>
</tr>
<tr>
<td>10 cm or more</td>
<td>2</td>
<td>7</td>
</tr>
<tr>
<td>Splenomegaly</td>
<td>7</td>
<td>23</td>
</tr>
<tr>
<td>Median no. of prior chemotherapy (range)</td>
<td>2 (1–9)</td>
<td></td>
</tr>
</tbody>
</table>

Figure 1. Time to progression (n = 30); median time to progression = 9.4 months.
study entry. With longer follow-up, there have been no additional cases of secondary malignancies or other long-term toxicities in this patient cohort.

**DISCUSSION**

In this phase II trial, 30 patients with relapsed or refractory low-grade, follicular, or transformed B-cell NHL and mild thrombocytopenia received a single course of radioimmunotherapy, with $^{90}$Y ibritumomab tiuxetan administered at a reduced dose of 0.3 mCi/kg. Overall, 83% of patients responded to therapy, with 14 patients (47%) achieving a CR or CRu. Responses have been durable, with 14 of 30 patients overall, and 12 of 14 CR/CRu patients, having a time to disease progression in excess of 12 months. Median TTP for all patients was 9.4 months, with a median TTNT of 14.6 months. In the 14 patients with long-term responses, median TTP was 24.6 months. At this time, median overall survival time has not yet been reached.

With respect to safety, toxicities were primarily hematologic, but were transient and reversible. With extended follow-up, no long-term adverse effects were noted, consistent with data previously indicating that treatment with ibritumomab tiuxetan radioimmunotherapy is not associated with an increased risk of secondary malignancies. Recent reports have indicated that subsequent NHL therapies, such as chemotherapy, immunotherapy, and even high-dose therapy with stem-cell or bone-marrow transplant, can be safely administered to patients who have previously received $^{90}$Y ibritumomab tiuxetan radioimmunotherapy.

In this study, 24 of 30 patients received one or more subsequent NHL therapies.

**CONCLUSION**

Persistent mild thrombocytopenia in patients with relapsed or refractory NHL is indicative of impaired bone-marrow reserves, either from prior chemotherapy, lymphomatous bone marrow involvement, or both. For these patients, treatment with reduced-dose $^{90}$Y ibritumomab tiuxetan is an effective option, producing a high rate of response and durable remissions. Therapy is generally well tolerated, and does not preclude the use of other subsequent lymphoma therapies.

**ACKNOWLEDGMENTS**

The authors wish to thank Christine Gutheil for her editorial assistance and Jessica Olson for her technical assistance.

**REFERENCES**