

Form record received

International Workshop on Waldenstrom's Macroglobulinemia <pattersonkent@outlook.com>

Sun 7/14/2024 3:40 PM

To:Patterson, Christopher <Christopher_Patterson@DFCI.HARVARD.EDU>

External Email - Use Caution

Record saved to database with ID: 112

Form ID: 1

Form title: Abstract Submission

Form name: Abstract_Submission

Submitted at: 2024-07-14 15:38:59

Submitter IP: 185.16.81.128

User-ID: 0

Username: -

User full name: -

Submitter provider: Unknown

Submitter browser: Mozilla/5.0 (Macintosh; Intel Mac OS X 10_15_7) AppleWebKit/605.1.15 (KHTML, like Gecko) Version/16.4.1 Safari/605.1.15

Submitter operating system: mac

First Name: Viera

Last Name: Sandecka

Email: vierkabj@yahoo.com

Registration Type: Delegate

Abstract Title: Obinotuzumab Plus Bendamustine In Patients With Relapsed/Refractory Waldenstrom's Macroglobulinemia: The Experience Of One Center.

Special Instructions: Background:

Waldenstrom's Macroglobulinemia (WM) belongs to the group of indolent B-Non-Hodgkin's lymphomas, characterized by a mostly slowly progressing clinical course and recurrent relapses (Grunenberg and Buske 2019). Fortunately, recent years have seen considerable progress in establishing new treatment concepts. We present the results of a evaluating the combination of obinotuzumab plus bendamustine in relapsed/refractory (R/R) WM.

Patients and Methods:

All nine patients had measurable, relapsed WM with CD20 expression and required treatment. The goal was to determine efficacy- overall response rate (ORR), progression free survival (PFS), overall survival (OS) and safety of this treatment combination. During the induction phase, patients received obinotuzumab plus bendamustine for 8 cycles, followed by a maintenance phase with obinotuzumab alone for ≤ 2 years. Three patients with R/R WM were treated with the induction combination, and 6 patients participated in the maintenance phase.

Results:

The median age was 70 years (range 48- 86) and the median number of prior treatments was 2 (range 1- 3). ORR was 100% at the end of induction phase; with 33,3% of patients achieving partial response; 33,3% of patients achieving very good partial response and 33,3% of patients achieving complete response. With a median follow-up of 90 months, median PFS and median OS were not reached. It is important to emphasize that all R/R WM patients treated with obinotuzumab plus bendamustine were pretreated with the median PFS after first line treatment of 72 months (95% confidence interval [CI], 12.7-168.2) and median PFS after second line treatment of 34 months (95% CI, 4.1- 48.5). At least one adverse events (AE) of any grade occurred in all nine treated patients. The most frequent AEs of any grade were neutropenia (55,5%), infection (33,3%) and thrombocytopenia (22,2%). The AE of grade 3 was herpes-zoster infection, occurred in one patient.

Conclusion:

Our modest findings show that the combination of obinotuzumab plus bendamustine is effective in R/R WM patients. Even if the median follow-up is long enough (median 90 months), the apparent lack of number patients could give new meaning to targeting of this treatment.

Conference: IWWM12

Authors:

Viera Sandecka (1,2), Zdenek Adam (1,2), Marta Krejci (1,2), Ivanna Boichuk (1,2), Martin Stork (1,2), Zdenek Kral (1,2), Jiri Mayer (1,2), Ludek Pour (1,2).

Affiliations:

(1) Department of Internal Medicine, Hematology and Oncology, University Hospital, Brno, Czech Republic

(2) Masaryk University Faculty of Medicine, Brno, Czech Republic

Title:

Obinotuzumab Plus Bendamustine In Patients With Relapsed/Refractory Waldenstrom's Macroglobulinemia: The Experience Of One Center.

Background:

Waldenstrom's Macroglobulinemia (WM) belongs to the group of indolent B-Non-Hodgkin's lymphomas, characterized by a mostly slowly progressing clinical course and recurrent relapses (Grunenberg and Buske 2019). Fortunately, recent years have seen considerable progress in establishing new treatment concepts. We present the results of a evaluating the combination of obinotuzumab plus bendamustine in relapsed/refractory (R/R) WM.

Patients and Methods:

All nine patients had measurable, relapsed WM with CD20 expression and required treatment. The goal was to determine efficacy- overall response rate (ORR), progression free survival (PFS), overall survival (OS) and safety of this treatment combination. During the induction phase, patients received obinotuzumab plus bendamustine for 8 cycles, followed by a maintenance phase with obinotuzumab alone for ≤ 2 years.

Three patients with R/R WM were treated with the induction combination, and 6 patients participated in the maintenance phase.

Results:

The median age was 70 years (range 48- 86) and the median number of prior treatments was 2 (range 1-3). ORR was 100% at the end of induction phase; with 33,3% of patients achieving partial response; 33,3% of patients achieving very good partial response and 33,3% of patients achieving complete response. With a median follow-up of 90 months, median PFS and median OS were not reached. It is important to emphasize that all R/R WM patients treated with obinotuzumab plus bendamustine were pretreated with the median PFS after first line treatment of 72 months (95% confidence interval [CI], 12.7-168.2) and median PFS after second line treatment of 34 months (95% CI, 4.1- 48.5). At least one adverse events (AE) of any grade occurred in all nine treated patients. The most frequent AEs of any grade were neutropenia (55,5%), infection (33,3%) and thrombocytopenia (22,2%). The AE of grade 3 was herpes-zoster infection, occurred in one patient.

Conclusion:

Our modest findings show that the combination of obinotuzumab plus bendamustine is effective in R/R WM patients. Even if the median follow-up is long enough (median 90 months), the apparent lack of number patients could give new meaning to targeting of this treatment.