

## Bendamustine - First Line - Indolent Non-Hodgkin's Lymphoma and Mantle Cell Lymphoma

(This form should be completed before the first dose is dispensed.)

## 1. Patient Profile

- \* Surname: .....
- \* Given Name: .....
- \* OHIN: ..... \* Chart Number: .....
- \* Postal Code: .....
- \* Height (cm): ..... \* Weight (kg): .....
- \* BSA (m<sup>2</sup>): ..... \* Gender:  Male  Female  Other
- \* Date of Birth: .....  
Day    Month    Year
- \* Site: .....
- \* Attending Physician (MRP- Most Responsible Physician): .....
- Requested Prior Approval  Yes    \* Patient on Clinical Trial  Yes  No
- Other (specify): .....
- Specify Arm:  
 Standard of care arm                       Experimental arm  
 Blinded / Unknown

## Request prior approval for enrolment

- \* Justification for Funding
- .....

## 2. Eligibility Criteria

The patient meets all of the following criteria:

Bendamustine is used in combination with rituximab in the first line setting in patients with indolent CD20 positive non-Hodgkin's lymphoma or mantle cell lymphoma  Yes

The patient has an ECOG performance status of less than or equal to 2

Yes

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### 3. Funded Dose

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Bendamustine 90mg/m<sup>2</sup> on Days 1 and 2 of a 28-day cycle to a maximum of 6 cycles (combination therapy).

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### 4. Notes

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- a. Bendamustine is not funded if used as a single agent.
- b. Patients who receive first line rituximab bendamustine would be eligible for rituximab maintenance provided that the maintenance rituximab funding criteria are met.

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### Supporting Documents

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To ensure reimbursement of your claim, both the completed enrolment form and a copy of the required documentation (where applicable) must be submitted through CCO e-Claims.

Signature of Attending Physician (MRP- Most Responsible Physician): .....

.....  
Day    Month    Year