

Regimen Monograph

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A - Regimen Name

RITU(IT) Regimen

Rituximab (intrathecal)

Disease Site Hematologic - Leukemia - Acute Lymphoblastic (ALL)
Hematologic - Lymphoma - Non-Hodgkin's High Grade
Hematologic - Lymphoma - Non-Hodgkin's Intermediate Grade

Intent Curative

Regimen Category **Evidence-informed :**

Regimen is considered appropriate as part of the standard care of patients; meaningfully improves outcomes (survival, quality of life), tolerability or costs compared to alternatives (recommended by the Disease Site Team and national consensus body e.g. pan-Canadian Oncology Drug Review, pCODR). Recommendation is based on an appropriately conducted phase III clinical trial relevant to the Canadian context OR (where phase III trials are not feasible) an appropriately sized phase II trial. Regimens where one or more drugs are not approved by Health Canada for any indication will be identified under Rationale and Use.

This **Regimen Abstract** is an **abbreviated** version of a Regimen Monograph and contains only top level information on usage, dosing, schedule, cycle length and special notes (if available). Information in regimen abstracts is accurate to the extent of the ST-QBP regimen master listings, and has not undergone the full review process of a regimen monograph. Full regimen monographs will be published for each ST-QBP regimen as they are developed.

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B - Drug Regimen

Note: Different rituximab products are NOT INTERCHANGEABLE.

[riTUXimab](#)

25 to 40 mg

IT

once or twice weekly

(This drug is not currently publicly funded for this regimen and intent)

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C - Cycle Frequency

FOR UP TO 8 INJECTIONS

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D - Premedication and Supportive Measures

Antiemetic Regimen: Minimal

Other Supportive Care:

Also refer to [CCO Antiemetic Recommendations](#).

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J - Administrative Information

Pharmacy Workload (average time per visit) 15.075 minutes

Nursing Workload (average time per visit) 65 minutes

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K - References

Jaime-Pérez JC, Rodríguez-Romo LN, González-Llano O, et al. Effectiveness of intrathecal rituximab in patients with acute lymphoblastic leukaemia relapsed to the CNS and resistant to conventional therapy. *Br J Haematol.* 2009 Mar;144(5):794-5.

Rubenstein JL, Fridlyand J, Abrey L, et al. Phase I Study of Intraventricular Administration of Rituximab in Patients With Recurrent CNS and Intraocular Lymphoma. *J Clin Oncol* 2007;25:1350-6.

Schulz H, Pels H, Schmidt-Wolf I, et al. Intraventricular treatment of relapsed central nervous system lymphoma with the anti-CD20 antibody rituximab. *Haematologica* 2004; 88:753-4.

August 2020 Updated interchangeability information in Drug Regimen section

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M - Disclaimer

Regimen Abstracts

A Regimen Abstract is an abbreviated version of a Regimen Monograph and contains only top level information on usage, dosing, schedule, cycle length and special notes (if available). It is intended for healthcare providers and is to be used for informational purposes only. It is not intended to constitute or be a substitute for medical advice, and all uses of the Regimen Abstract are subject to clinical judgment. Such information is provided on an "as-is" basis, without any representation, warranty, or condition, whether express, or implied, statutory or otherwise, as to the information's quality, accuracy, currency, completeness, or reliability, and Cancer Care Ontario disclaims all liability for the use of this information, and for any claims, actions, demands or suits that arise from such use.

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Regimen Monographs

Refer to the [New Drug Funding Program](#) or [Ontario Public Drug Programs](#) websites for the most up-to-date public funding information.

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