Regimen Monograph

Regimen Name | Drug Regimen | Cycle Frequency | Premedication and Supportive Measures | Administrative Information | References | Other Notes | Disclaimer

A - Regimen Name

MOGA Regimen

Mogamulizumab

Disease Site

Hematologic
Lymphoma - T-cell

(Mycosis fungoides or Sézary syndrome)

Intent

Palliative

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.

Rationale and Uses

For the treatment of patients with relapsed or refractory mycosis fungoides (MF) or Sézary syndrome (SS) (stage IB to IV), who had treatment failure after at least 1 prior course of systemic treatment. Patients must **NOT** have active or untreated CNS metastases.

Supplementary Public Funding	**mogamulizumab** New Drug Funding Program (Mogamulizumab - Relapsed or Refractory Mycosis Fungoides or Sezary Syndrome) (NDFP Website)

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B - Drug Regimen

Cycle 1:

mogamulizumab	1 mg /kg	IV	Days 1, 8, 15, 22

Cycle 2 and onwards:

mogamulizumab	1 mg /kg	IV	Days 1 and 15

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C - Cycle Frequency

REPEAT EVERY 28 DAYS

Continue until disease progression or unacceptable toxicity, whichever comes first

Patients with a global complete response can continue treatment for up to 12 months or until disease progression, whichever comes first

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D - Premedication and Supportive Measures

Antiemetic Regimen: Low

Also refer to CCO Antiemetic Recommendations.

Screen for hepatitis B virus in all cancer patients starting systemic treatment. Refer to the hepatitis B virus screening and management guideline.

Patients at risk of tumour lysis syndrome should have appropriate prophylaxis and be monitored closely.

Premedications (prophylaxis for infusion reactions):

- Consider premedication (e.g. diphenhydramine and acetaminophen) before the first infusion.
- Give premedication for subsequent infusions if an infusion reaction has occurred.

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J - Administrative Information

Approximate Patient Visit	1.5 hours

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K - References

CADTH reimbursement recommendation: Mogamulizumab (relapsed or refractory mycosis fungoides or Sézary syndrome). Journal of Canadian Health Technologies 2022;2(8) (published online).

Kim YH, Bagot M, Pinter-Brown L, et al. Mogamulizumab versus vorinostat in previously treated cutaneous T-cell lymphoma (MAVORIC): an international, open-label, randomised, controlled phase 3 trial. Lancet Oncol. 2018 Sep;19(9):1192-204.

May 2024 Modified Rationale/uses and Cycle frequency sections; added NDFP form

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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.

The information set out in the drug monographs, regimen monographs, appendices and symptom management information (for health professionals) contained in the Drug Formulary (the "Formulary") is intended for healthcare providers and is to be used for informational purposes only. The information is not intended to cover all possible uses, directions, precautions, drug interactions or adverse effects of a particular drug, nor should it be construed to indicate that use of a particular drug is safe, appropriate or effective for a given condition. The information in the Formulary is not intended to constitute or be a substitute for medical advice and should not be relied upon in any such regard. All uses of the Formulary are subject to clinical judgment and actual prescribing patterns may not follow the information provided in the Formulary.

The format and content of the drug monographs, regimen monographs, appendices and symptom management information contained in the Formulary will change as they are reviewed and revised on a periodic basis. The date of last revision will be visible on each page of the monograph and regimen. Since standards of usage are constantly evolving, it is advised that the Formulary not be used as the sole source of information. It is strongly recommended that original references or product monograph be consulted prior to using a chemotherapy regimen for the first time.

Some Formulary documents, such as the medication information sheets, regimen information sheets and symptom management information (for patients), are intended for patients. Patients should always consult with their healthcare provider if they have questions regarding any information set out in the Formulary documents.

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