#### Regimen Monograph

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

# A - Regimen Name

# **DEXAPOMASELI** Regimen

Dexamethasone-Pomalidomide-Selinexor

Disease Site Hematologic

Multiple Myeloma

**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 treatment of relapsed or refractory multiple myeloma

Supplementary dexamethasone

Public Funding ODB - General Benefit (dexamethasone) (ODB Formulary )

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## **B** - Drug Regimen

dexamethasone

40 mg

PO

Days 1, 8, 15, 22

In elderly patients, the dexamethasone dose should be reduced (i.e. to 20 mg once weekly).

**pomalidomide** 

4 mg

PO

Days 1 to 21

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

Pomalidomide may only be prescribed and dispensed by physicians and pharmacists registered with a controlled distribution program. Patients must also be registered and meet all conditions of the program.

selinexor

40 mg

PO

Days 1, 8, 15, 22

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

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## C - Cycle Frequency

## **REPEAT EVERY 28 DAYS**

Until disease progression or unacceptable toxicity

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## **D** - Premedication and Supportive Measures

## Antiemetic Regimen: Moderate

(In the BOSTON study (selinexor + bortezomib + dexamethasone), patients received a 5-HT3 receptor antagonist ± other antiemetics (e.g. olanzapine or NK1 RA) prior to and during treatment, and as needed after treatment.)

Also refer to <u>CCO Antiemetic Recommendations</u>.

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

## Other Supportive Care:

- Patients at risk of tumour lysis syndrome should have appropriate prophylaxis and be monitored closely.
- Prophylactic antithrombotics, such as low dose aspirin, low molecular weight heparins or warfarin, are recommended.
- Patients should maintain adequate fluid and caloric intake during selinexor treatment.
   Consider IV hydration for patients at risk of dehydration.

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#### K - References

Bahlis NJ, Sutherland H, White D, et al. Selinexor plus low-dose bortezomib and dexamethasone for patients with relapsed or refractory multiple myeloma. Blood.2018;132(24):2546-2554.

Bortezomib drug monograph. Ontario Health (Cancer Care Ontario).

Schiller GJ, Lipe BC, Bahlis NJ, et al. Selinexor-based triplet regimens in patients with multiple myeloma previously treated with anti-CD38 monoclonal antibodies. Clin Lymphoma Myeloma Leuk 2023 Sep;23(9):e286-e296.e4. doi: 10.1016/j.clml.2023.06.001.

Selinexor drug monograph. Ontario Health (Cancer Care Ontario).

White D, Schiller GJ, Madan S, et al. Efficacy and safety of once weekly selinexor 40 mg versus 60 mg with pomalidomide and dexamethasone in relapsed and/or refractory multiple myeloma. Front Oncol 2024 May 17;14:1352281. doi: 10.3389/fonc.2024.1352281.

May 2025 new ST-QBP regimen

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#### M - Disclaimer

#### Regimen Abstracts

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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.

#### Regimen Monographs

Refer to the <u>New Drug Funding Program</u> or <u>Ontario Public Drug Programs</u> 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.

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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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