Regimen Monograph

 Regimen Name
 Drug Regimen
 Cycle Frequency
 Premedication and Supportive Measures
 Dose Modifications
 Adverse

 Effects
 Interactions
 Drug Administration and Special Precautions
 Recommended Clinical Monitoring
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 Information
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A - Regimen Name

AVD+NIVL Regimen

DOXOrubicin-vinBLAStine-Dacarbazine-Nivolumab

Disease Site Hematologic

Lymphoma - Hodgkin

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.

Rationale and Uses

First-line treatment of advanced stage (Ann Arbor stage III or IV) Hodgkin lymphoma

(Refer to the NDFP eligibility form for detailed funding criteria.)

Supplementary

<u>nivolumab</u>

Public Funding Ne

New Drug Funding Program (Nivolumab - In Combination with Chemotherapy for Previously Untreated Advanced Stage Hodgkin Lymphoma) (NDFP Website)

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B - Drug Regimen			
DOXOrubicin	25 mg /m²	IV	Days 1, 15
<u>vinBLAStine</u>	6 mg /m²	IV	Days 1, 15
dacarbazine	375 mg /m²	IV	Days 1, 15
nivolumab ^{1,2}	3 mg /kg	IV (max 240 mg)	Days 1, 15

¹Dosing based on NDFP funding criteria.

C - Cycle Frequency

REPEAT EVERY 28 DAYS

Up to a maximum of 6 cycles, unless disease progression or unacceptable toxicity

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D - Premedication and Supportive Measures

Antiemetic Regimen: Moderate

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

²Administer nivolumab after AVD as per institutional guidelines. back to top

Nivolumab pre-medications (prophylaxis for infusion reaction):

- Routine pre-medication is not recommended.
- May consider pre-medication with antipyretics and H1-receptor antagonists if an IR has occurred in the past.

Also refer to the CCO guideline for detailed description of <u>Management of Cancer Medication-Related Infusion Reactions</u>.

Other Supportive Care:

- Screen for hepatitis B virus in all cancer patients starting systemic treatment. Refer to the <u>hepatitis B virus screening and management</u> guideline.
- G-CSF was allowable in the clinical trial (Herrera et al).
- Refer to the CCO guideline for detailed description of Immune-mediated toxicities and their management.

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J - Administrative Information

Approximate Patient Visit 2 hours

Pharmacy Workload (average time per visit) 41.773 minutes

Nursing Workload (average time per visit) 62.417 minutes

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K - References

Canada's Drug Agency. Reimbursement Recommendation: Nivolumab. Canadian Journal of Health Technologies. August 2025

Herrera AF, LeBlanc M, Castellino SM, et al. Nivolumab+AVD in advanced-stage classic Hodgkin's lymphoma. N Engl J Med 2024 Oct 17;391(15):1379-89.

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

December 2025 Updated Rationale/Uses, Supplementary Public Funding, Drug Regimen and Cycle Frequency sections

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M - Disclaimer

Regimen Abstracts

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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.

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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