Regimen Monograph

 Regimen Name
 Drug Regimen
 Cycle Frequency
 Premedication and Supportive Measures
 Administrative Information
 References

 Other Notes
 Disclaimer

A - Regimen Name

CISPETOP(PO) Regimen			
Disease Site	Lung Non-Small Cell Small Cell		
Intent	Adjuvant 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.		
	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.		
Supplementary Public Funding	<u>etoposide</u> ODB - General Benefit (etoposide - oral capsules)		

back to top

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CISPETOP(PO)

B - Drug Regimen			
<u>CISplatin</u>	75 mg /m²	IV	Day 1
<u>etoposide</u>	200 mg /m²	PO	Days 1 to 3

Alternative Schedule: Etoposide 100 mg/m² IV day 1 then 200 mg/m² PO days 2 to 3

back to top

C - Cycle Frequency

REPEAT EVERY 21 DAYS

For a usual total of 4 to 6 cycles unless disease progression or unacceptable toxicity occurs

back to top

Antiemetic Regimen:	High (Cisplatin ≥70 mg/m2)
	No routine prophylaxis for etoposide PO

Other Supportive Care:

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

Standard regimens for Cisplatin premedication and hydration should be followed. Refer to local guidelines.

back to top

J - Administrative Information

Approximate Patient Visit	3-4 hours
Pharmacy Workload (average time per visit)	42.128 minutes
Nursing Workload (average time per visit)	49.167 minutes

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back to top

K - References

Cisplatin and etoposide drug monographs, Cancer Care Ontario.

Belani CP, Lee JS, Socinski MA, et al. Randomized phase III trial comparing cisplatin-etoposide to carboplatin-paclitaxel in advanced or metastatic non-small cell lung cancer. Ann Oncol. 2005;16(7):1069-75.

Bonomi P, Kim KM, Fairclough D et al. Comparison of survival and quality of life in advanced nonsmall cell lung cancer patients treated with two dose levels of paclitaxel combined with cisplatin versus etoposide with cisplatin: results of an Eastern Cooperative Oncology Group trial. J Clin Oncol 2000; 18: 623–631.

Klastersky J, Sculier JP, Dabouis G, et al. A randomized trial of two platinum combinations in patients with advanced non-small cell lung cancer: a preliminary report. European Organization for the Research and Treatment of Cancer--Lung Cancer Working Party. Semin Oncol. 1990 Feb;17(1 Suppl 2):20-4.

Sundstrom S, Bremnes RM, Kaasa S, et al. Cisplatin and etoposide regimen is superior to cyclophosphamide, epirubicin, and vincristine regimen in small-Cell lung cancer: results from a randomized phase III trial with 5 years' follow-up. J Clin Oncol 2002;20:4665-72.

PEBC Advice Documents or Guidelines

- <u>Adjuvant Systemic and Radiation Therapy for Stage I to IIIA Completely Resected Non–Small-Cell Lung Cancers: ASCO-CCO Clinical Practice Guideline Update</u>
- <u>Systemic Therapy for Small-Cell Lung Cancer: ASCO-OH(CCO) Guideline</u>

November 2023 Added PEBC guideline link

back to top

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,

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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.

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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back to top