

Regimen Monograph

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A - Regimen Name

CISPGEMCPACL Regimen

CISplatin-Gemcitabine-PACLitaxel

Disease Site Genitourinary - Testis

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 Salvage therapy for relapsed/refractory germ cell tumours

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B - Drug Regimen

PACLitaxel	80 mg /m ²	IV	Days 1 and 8
gemcitabine	800 mg /m ²	IV	Days 1 and 8
CISplatin	50 mg /m ²	IV	Days 1 and 8

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C - Cycle Frequency**REPEAT EVERY 21 DAYS**

For up to 8 cycles unless disease progression or unacceptable toxicity occurs

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D - Premedication and Supportive Measures

Antiemetic Regimen: Moderate

Other Supportive Care:

Also refer to [CCO Antiemetic Summary](#)

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

Paclitaxel: Patients should be pretreated with a corticosteroid as well as an antihistamine and a H2 blocker: For example:

- DEXAMETHASONE 20mg PO 12 & 6 hours or 20mg IV 30 minutes before paclitaxel
- DIPHENHYDRAMINE 50mg IV 30 minutes before paclitaxel
- RANITIDINE 50mg IV 30 minutes before paclitaxel

GCSF support was used in clinical trials

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J - Administrative Information

Approximate Patient Visit	5 hours
Pharmacy Workload (average time per visit)	43.811 minutes
Nursing Workload (average time per visit)	49.833 minutes

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K - References

Cisplatin, gemcitabine and paclitaxel drug monographs, Cancer Care Ontario.

Necchi A, Nicolai N, Mariani L, et al. Combination of paclitaxel, cisplatin, and gemcitabine (TPG) for multiple relapses or platinum-resistant germ cell tumors: long-term outcomes. Clin Genitourin Cancer 2014;12(1):63-69.

Nicolai N, Necchi A, Gianni L, et al. Long-term results of a combination of paclitaxel, cisplatin and gemcitabine for salvage therapy in male germ-cell tumours. BJU Int. 2009 Aug;104(3):340-6.

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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.

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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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