

## Regimen Monograph

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## A - Regimen Name

# CRBPETOP Regimen

CARBOplatin-Etoposide

**Disease Site** Skin - Merkel Cell

**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.

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**B - Drug Regimen****CARBO**platin

AUC 5

IV

Day 1

Adjust Carboplatin dose to AUC target (using Calvert formula) as outlined in the "Other Notes" section.

**etoposide**100 mg /m<sup>2</sup>

IV

Days 1 to 3

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Until disease progression or unacceptable toxicity, usually up to 6 cycles due to cumulative carboplatin toxicity

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**Antiemetic Regimen:** Moderate + NK1 antagonist (Carboplatin AUC  $\geq$  5) (D1)  
Low (D2-3)

**Other Supportive Care:**

Also refer to [CCO Antiemetic Recommendations](#).

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Approximate Patient Visit	Day 1: 2 hours; Day 2-3: 1 hour
Pharmacy Workload (average time per visit)	13.782 minutes
Nursing Workload (average time per visit)	42.500 minutes

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Carboplatin, etoposide drug monographs, Cancer Care Ontario.

Davis MP, Miller EM, Rau RC, et al. The use of VP16 and cisplatin in the treatment of Merkel cell carcinoma. J Dermatol Surg Oncol. 1990 Mar;16(3):276-8.

McAfee WJ, Morris CG, Mendenhall CM, et al. Merkel cell carcinoma: treatment and outcomes. Cancer 2005; 104(8):1761-4.

Pectasides D, Pectasides M, Psyrri A, et al. Cisplatin-based chemotherapy for merkel cell carcinoma of the skin. Cancer Invest 2006;24(8):780-5.

Redmond III J, Perry J, Sowray P, et al. Chemotherapy of disseminated merkel-cell carcinoma. American Journal of Clinical Oncology 1991;14(4):305-7.

Tai PTH, Yu E, Winquist E, et al., Chemotherapy in neuroendocrine/ Merkel cell carcinoma of the skin: case series and 8 Journal of Skin Cancer review of 204 cases. Journal of Clinical Oncology 2000;18(12):2493-9.

**June 2019** Updated emetic risk category

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## L - Other Notes

### Calvert Formula:

**DOSE (mg) = target AUC X (GFR + 25)**

- AUC = product of serum concentration (mg/mL) and time (min)
- GFR (glomerular filtration rate) expressed as measured Creatinine Clearance or estimated from Serum Creatinine (by Cockcroft and Gault method or Jelliffe method)

Calvert AH, Newell DR, Gumbrell LA, et al, Carboplatin dosage: Prospective evaluation of a simple formula based on renal function. J Clin Oncol, 1989; 7: 1748-1756

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## M - Disclaimer

### ***Regimen Abstracts***

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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.

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 [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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