BC Cancer Protocol Summary for Therapy for High Risk Gestational Trophoblastic Neoplasia (GTN) using Etoposide, Methotrexate, Leucovorin (Folinic Acid), DACTINomycin, Cyclophosphamide and vinCRIStine

Protocol Code GOTDEMACO

Tumour Group Gynecology

Contact Physician Dr. Anna Tinker

ELIGIBILITY:

- High risk Gestational Trophoblastic Neoplasm (GTN) as determined using the modified World Health Organization (WHO) Prognostic Scoring System as adapted by the International Federation of Gynecology and Obstetrics (FIGO).
 - Patients with a Risk Score of ≥7 are considered high risk
 - o The risk score is calculated as follows:

| Risk Factor | SCORE | | | |
|---|------------------|----------------------------------|----------------------------------|------------------|
| | 0 | 1 | 2 | 4 |
| Age (years) | <40 | >40 | | |
| Antecedent pregnancy | Mole | Abortion | Term | |
| Interval (months)* | <4 | 4-6 | 7-12 | >12 |
| Pre-treatment serum beta hCG (mIU/mL)** | <10 ³ | 10 ³ -10 ⁴ | 10 ⁴ -10 ⁵ | >10 ⁵ |
| Sites of metastases | Lung | Spleen and kidney | GI tract | Brain and liver |
| Number of metastases | - | 1-4 | 5-8 | >8 |
| Largest tumour size (including uterus) (cm) | | 3-4 | 5 | |
| Prior failed chemotherapy | - | - | Single drug | ≥2 drugs |

^{*} interval (in months) between end of antecedent pregnancy and start of chemotherapy

 Contact BC Cancer for recommendations if more than 1 year post normal pregnancy and not currently pregnant

EXCLUSIONS:

• Low to moderate risk GTN (prognostic score less than seven). These patients should be treated on the BC Cancer low risk protocol GOTDLRA (risk score of 0 to 4) and moderate risk protocol GOTDMR (risk score of 5 or 6), respectively.

^{**} use the immediate pre-treatment beta hCG - not the peak hCG during pregnancy or prior to uterine evacuation

TESTS:

- If the GTN was not confirmed histopathologically (e.g. only products of conception were identified at the time of the dilation and curettage) then a pelvic ultrasound within 5-7 days of treatment initiation is required to rule out a previously undetected viable intra-uterine pregnancy.
- **Baseline:** CBC & diff, beta hCG tumour marker, sodium, potassium, creatinine, Alk Phos, ALT, GGT, LDH, Bilirubin, chest X-ray, CT of brain, CT abdomen and pelvis
- If the beta hCG tumour marker is > 100,000 mIU/mL check TSH
- **Before each treatment (Day 1):** CBC & diff, beta hCG tumour marker, electrolytes panel, creatinine, Alk Phos, ALT, GGT, LDH, Bilirubin
- Before each treatment (Day 8): CBC & diff, creatinine

PREMEDICATIONS:

Antiemetic protocol for high-moderate emetogenic chemotherapy (see protocol SCNAUSEA).

If history of etoposide hypersensitivity:

- hydrocortisone 100 mg IV prior to etoposide
- diphenhydrAMINE 50 mg IV prior to etoposide

PREHYDRATION:

Not routinely necessary.

TREATMENT:

| Day | Drug | Dose | BC Cancer Administration Guideline | | |
|-----|------------------------------------|------------------------|--|--|--|
| | etoposide | 100 mg/m² | IV in NS 250 to 1000 mL over 45 minutes to 1 hour 30 minutes (Use non-DEHP bag and tubing with 0.2 micron in-line filter) | | |
| 1 | DACTINomycin | 0.5 mg | IV push | | |
| | methotrexate 300 mg/m ² | | IV in NS 500 mL over 12 hours | | |
| | | | | | |
| | etoposide 100 mg/m² | | IV in NS 250 to 1000 mL over 45 minutes to 1 hour 30 minutes (Use non-DEHP bag and tubing with 0.2 micron in-line filter) | | |
| 2 | 2 DACTINomycin 0.5 n | 0.5 mg | IV push | | |
| | leucovorin (folinic acid) | 15 mg q12h for 4 doses | PO Start 24 hours after start of Day 1 methotrexate infusion | | |
| _ | | | | | |
| 8 | vinCRIStine | 0.8 mg/m ² | IV in NS 50 mL over 10 minutes | | |
| | cyclophosphamide | 600 mg/m ² | IV in NS 250 to 500 mL over 30 minutes | | |

Repeat every 14 days until 3 cycles post fall of beta hCG tumour marker to below 5 mlU/mL

POST-HYDRATION:

 1000 mL D5W-1/2NS with 20 mEq Potassium Chloride and 100 mEq Sodium Bicarbonate/L at 200 mL/h IV for 20 hours after the end of the methotrexate infusion. Hydration infusion may be interrupted for administration of Day 2 chemotherapy.

DOSE MODIFICATIONS:

1. Hematological

On treatment day:

| ANC (x10 ⁹ /L) | Platelets (x10 ⁹ /L) | etoposide dose | DACTINomycin dose | methotrexate dose | cyclophosphamide dose |
|------------------------------|------------------------------------|------------------------|---------------------------------|-------------------|--------------------------|
| greater than or equal to 1.0 | greater than or equal to 100 | 100% | 100% | 100% | 100% |
| 0.7 to less than 1.0 | 75 to less than100 | 100% | 66% | 100% | 100% |
| less than 0.7* | less than 75 | 50 mg x 7 days only | Hold until next cycle, then 66% | 100% | 100% |

^{*}unless monocytes greater than 10% of total WBC, then give 66% of DACTINomycin dose

- 2. **Renal dysfunction:** Dose modification of methotrexate may be required. Refer to BC Cancer Drug Manual.
- 3. **Neurotoxicity:** vinCRIStine only:

| Toxicity | Dose Modification |
|------------------------------|-------------------|
| Dysesthesias, areflexia only | 100% |
| Abnormal buttoning, writing | 67% |
| Motor neuropathy, moderate | 50% |
| Motor neuropathy, severe | omit |

- 4. **Hepatic dysfunction**: Dose modification of all drugs may be required. Refer to BC Cancer Drug Manual.
- 5. Third space fluids (ascites, pleural effusions): See Precaution statement, below.
- 6. **Stomatitis:** decrease DACTINomycin to 0.4 mg

PRECAUTIONS:

- 1. **Neutropenia**: Fever or other evidence of infection must be assessed promptly and treated aggressively. Filgrastim (G-CSF) may be needed to maintain white count.
- 2. **Extravasation**: DACTINomycin and vinCRIStine causes pain and tissue necrosis if extravasated. Refer to BC Cancer Extravasation Guidelines.
- 3. **Renal Toxicity**: Nephrotoxicity is possible with high (greater than 1 g/m²) doses of methotrexate. The risk of renal failure can be minimised by brisk diuresis and alkalinization of the urine with sodium bicarbonate. Encourage oral hydration.
- 4. **Third Space Fluids**: Evacuation of third space fluids e.g., ascites, pleural effusions, prior to treatment is recommended. Consider methotrexate drug level monitoring and dose reduction if drainage cannot be achieved.

Call Dr. Anna Tinker or tumour group delegate at (604) 877-6000 or 1-800-663-3333 with any problems or questions regarding this treatment program.

References

- Alazzam M1, Tidy J, Osborne R, et al. Chemotherapy for resistant or recurrent gestational trophoblastic neoplasia. Cochrane Database Syst Rev 2016 Jan 13;1:CD008891. 10.1002/14651858.CD008891.pub3.
- 2. Alifrangis C, Agarwal R, Short D, et al. EMA/CO for high-risk gestational trophoblastic neoplasia: good outcomes with induction low-dose etoposide-cisplatin and genetic analysis. J Clin Oncol 2013;31(2):280-6.
- 3. Turan T, Karacay O, Tulunay G, et al. Results with EMA/CO (etoposide, methotrexate, actinomycin D, cyclophosphamide, vincristine) chemotherapy in gestational trophoblastic neoplasia. Int J Gynecol Cancer 2006;16(3):1432-8.