Rationale

- Germ cell tumors (GCT) represent the most common malignancy affecting adolescent and young adult men in both Europe and the United States. [1]
- Up to 30% of pts with advanced GCT are not cured with first-line chemotherapy and require salvage treatment, consisting of either conventional-dose chemotherapy (CDCT) or high-dose chemotherapy (HDCT).
- The optimal initial salvage approach (HDCT vs. CDCT) is not clear; retrospective studies consistently report superior outcomes with HDCT but the one prior RCT (IT-94) found no improvement with HDCT. [2]
- IT-94 had several shortcomings including:
  - Only 1 high-dose cycle in the HCT arm (most effective regimens use 2 or 3)
  - Exclusion of pts with incomplete response to first line chemotherapy (only allowed relapsing pts)
  - >25% of pts assigned to HDCT arm never received the high-dose cycle
- The current TIGER study aims to determine whether HDCT or CDCT is the optimal initial salvage approach.
- Collaboration between Alliance, EORTC, ECOG, SWOG, COG, ANZUP
- Open in 13 countries across 3 continents (North America, Europe and Australia)

Alliance A031102: A Randomized Phase III Trial Comparing Conventional-Dose Chemotherapy Using Paclitaxel, Ifosfamide, and Cisplatin (TIP) with High-Dose Chemotherapy Using Mobilizing Paclitaxel Plus Ifosfamide Followed By High-dose Carboplatin and Etoposide (TI-CE) as First Salvage Treatment Of Germ Cell Tumors (TIGER)

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Endpoints

Rationale

Endpoints

Study Schema

Treatment Plan

Key Eligibility Criteria

Follow Up

Primary
Overall Survival (OS)
Sample size = 420 patients (expected deaths = 232)
- 168 patients from US Cooperative Groups; 252 patients from EORTC
- Cure rate model: 35% of patients will be cured with TIP (median OS 1.5yrs)
- Power 81%, 1-sided type I error rate = 0.05 where TI-CE will reduce the hazard by 29%
  ($\theta = 0.71$)
  - Efficacy & futility analyses regularly per Alliance policy
  - Formal toxicity analysis for early stopping (90% CI for one-sided difference in treatment-related mortality > 16%)

Secondary
- Progression-free survival (PFS)
- Favorable response rate
  - CR and PR-negative markers
  - Treatment-related mortality
  - Toxicities by CTCAE v4.0
  - Validation of the IPFSG risk model [3]

Correlative Studies
- Quality of life assessment of TIP vs. TI-CE using EORTC QLQ-C30 and the QLQ-TC26
- Pharmacogenomics: correlation of SNP rs1649942 (associated with platinum response) with PFS in the entire study population
- Tumor whole exome analysis assessing for the association of genetic alterations with OS
  - DNA Repair genes (TP53) [6]
  - PI3K/RAS/RAF genes [7]

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Study Schema

N=420

1:1

Primary Endpoint
Overall Survival

TIP (n=210)

TI-CE (n=210)

Stratification Factors:
1. International Prognostic Factor Study Group (IPFSG) risk class
2. Continent of enrollment

Please use the headings above to navigate through the different sections of the poster

TAP TO RETURN TO KIOSK MENU
**Alliance A031102: A Randomized Phase III Trial Comparing Conventional-Dose Chemotherapy Using Paclitaxel, Ifosfamide, and Cisplatin (TIP) with High-Dose Chemotherapy Using Mobilizing Paclitaxel Plus Ifosfamide Followed By High-dose Carboplatin and Etoposide (TI-CE) as First Salvage Treatment Of Germ Cell Tumors (TIGER)**

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### Treatment Plan

**ARM A (CDCT): TIP**

<table>
<thead>
<tr>
<th>Drugs and Doses</th>
<th>Infusion Time</th>
<th>Days</th>
</tr>
</thead>
<tbody>
<tr>
<td>Paclitaxel 250mg/m²</td>
<td>24 hours</td>
<td>1</td>
</tr>
<tr>
<td>Ifosfamide 1500mg/m²</td>
<td>1-2 hours</td>
<td>2-5</td>
</tr>
<tr>
<td>Mesna</td>
<td>1-2 hours</td>
<td>2-5</td>
</tr>
<tr>
<td>Cisplatin 25mg/m²</td>
<td>0.5 – 1 hour</td>
<td>2-5</td>
</tr>
</tbody>
</table>

- 4 cycles given every 21 days
- Exact infusion time at discretion of investigator and institution
- Levofloxacin 500mg daily given as prophylaxis from day 7 to 13 of each cycle

**ARM B (HDCT): TI-CE**

<table>
<thead>
<tr>
<th>Cycle #</th>
<th>Cycle Duration</th>
<th>Drugs and Doses</th>
<th>Days</th>
</tr>
</thead>
<tbody>
<tr>
<td>1-2</td>
<td>14 days</td>
<td>Paclitaxel 200mg/m²</td>
<td>1</td>
</tr>
<tr>
<td>3, 4, 5</td>
<td>21 days</td>
<td>Etoposide 400mg/m²/d</td>
<td>1-3</td>
</tr>
</tbody>
</table>

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Key Eligibility Criteria

- Male
- Age ≥ 14 years
- ECOG 0 - 2
- Histologic confirmation of GCT diagnosis
- Evidence of progressive disease after one line of cisplatin-based chemotherapy including:
  - Biopsy of new/growing mass consistent with non-teratomatous GCT
  - Rising HCG or AFP
  - New or enlarging lesions in the setting of elevated (even if not rising) HCG or AFP
- Received 1 prior line of cisplatin-based chemo (3 – 6 cycles)
- No prior treatment with TIP or HDCT
- GFR ≥ 50ml/min by 2 methods or ≥ 70ml/min by 1 method
- ANC ≥ 1,500/mm3, PLT ≥ 100,000/mm3
- T bilirubin ≤ 2.5 x ULN and AST/ALT ≤ 2.5 x ULN
- Negative serology for HIV, HTLV, Hep B, and Hep C
- No secondary somatic malignancy arising from teratoma
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