

## COG Phase III Ewing Sarcoma

### Protocol Title

*AEWS1031 A Phase III Randomized Trial of Adding Vincristine-topotecan-cyclophosphamide to Standard Chemotherapy in Initial Treatment of Non-metastatic Ewing Sarcoma*

### COG

Children's  
Oncology Group

### Principal Investigator

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### Study Coordinator

Wendy Bissette, RN, BS

### Status

Active

### Key Inclusion Criteria\*

- Age  $\leq$  50 years at diagnosis
- Patients with newly diagnosed, biopsy confirmed, extracranial, non-metastatic Ewing sarcoma or PNET of bone or soft tissue
  - For the purpose of this study, chest wall tumors with ipsilateral pleural effusions, ipsilateral positive pleural fluid cytology or ipsilateral pleural based secondary tumor nodules will be considered localized disease
  - Patients with regional node involvement, based on clinical suspicion confirmed by pathologic documentation are considered to be non-metastatic
  - Tumors arising in the bony skull (extra-dural) are considered to be extracranial
- Adequate renal, liver, and cardiac function

### Key Exclusion Criteria\*

- Prior chemotherapy or radiation therapy
- Evidence of metastatic disease
- Patients whose tumors arise in the intra-dural soft tissue
- Pregnant or breastfeeding females

### Therapies

#### Chemotherapy

- |                    |               |
|--------------------|---------------|
| ▪ Vincristine      | ▪ Ifosfamide  |
| ▪ Doxorubicin      | ▪ Etoposide   |
| ▪ Cyclophosphamide | ▪ Mesna       |
| ▪ Topotecan        | ▪ Dexrazoxane |

#### Radiation

#### Surgery

### Referrals

*Please contact for referrals and/or study information:*

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\* Additional criteria apply.