**Section: Contact Information**

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**Section: Project Requirements and Description**

**Group: Requirements to submit AOI**
A comprehensive review of previously published data has been completed. : Yes 
The specific aims are clear and focused. : Yes 
The investigator has appropriate experience and expertise to develop the concept proposal; if not, has identified a mentor or senior co-investigator. : Yes 
The investigator agrees to develop an initial draft of the concept proposal within 6 weeks of approval of the AOI and to finalize the concept proposal within 6 months. : Yes 
Project Title: Late Outcomes in Survivors of Childhood Cancer Exposed to Platinum Based Chemotherapy: A Report from the Childhood Cancer Survivor Study (CCSS) 
Planned research population (eligibility criteria): Survivors of childhood cancer enrolled in the Childhood Cancer Survivor Study (CCSS) 
Proposed specific aims: 
The study aims to characterize and estimate the prevalence of late outcomes among survivors of childhood cancer exposed to platinum-based chemotherapy. 

1)Estimate the prevalence of chronic health conditions among survivors treated with platinum compounds in the Childhood Cancer Survivor Study (CCSS) cohort compared to survivors treated with non-platinum-based therapy. 

2)Describe the prevalence of subsequent neoplasms (benign and malignant) among survivors treated with platinum compounds in the CCSS cohort compared to survivors treated with non-platinum-based therapy. 

3)Confirm the prevalence of chronic health conditions and subsequent neoplasms identified in CCSS survivors exposed to platinum-based compounds among survivors in the St. Jude Lifetime (SJLIFE) cohort study exposed to platinum-based compounds. 

Will the project require non-CCSS funding to complete? : No 
If yes, what would be the anticipated source(s) and timeline(s) for securing funding? : 

**Group: Does this project require contact of CCSS study subjects for:** 
Additional self-reported information: No 
Biological samples: No
Medical record data: No
If yes to any of the above, please briefly describe:

**Group: What CCSS Working Group(s) would likely be involved? (Check all that apply)**

- Second Malignancy: Secondary
- Chronic Disease: Primary
- Psychology / Neuropsychology:
- Genetics:
- Cancer Control:
- Epidemiology / Biostatistics:

**Section: Outcomes or Correlative Factors**

- Late mortality: Secondary
- Second Malignancy: Primary

**Group: Health Behaviors**

- Tobacco: Correlative Factors
- Alcohol:
- Physical activity:
- Medical screening:
- Other:
  If other, please specify:

**Group: Psychosocial**

- Insurance:
- Marriage:
- Education:
- Employment:
- Other:
  If other, please specify:

**Group: Medical Conditions**

- Hearing/Vision/Speech:
- Hormonal systems:
- Heart and vascular:
- Respiratory:
- Digestive:
- Surgical procedures:
- Brain and nervous system:
- Other: Primary
  If other, please specify: We will use the CTCAE grading

**Group: Medications**

Describe medications:

**Group: Psychologic/Quality of Life**

BSI-18:
SF-36:
CCSS-NCQ:
PTS:
PTG:
Other:
If other, please specify:

**Group: Other**
Pregnancy and offspring: Primary
Family history:
Chronic conditions (CTCAE v3): Primary
Health status:

**Group: Demographic**
Age: Correlative Factors
Race: Correlative Factors
Sex: Correlative Factors
Other: Correlative Factors
If other, please specify: Height, Weight, Body mass index (kg/m2)

**Group: Cancer treatment**
Chemotherapy: Correlative Factors
Radiation therapy: Correlative Factors
Surgery: Correlative Factors

**Section: Anticipated Sources of Statistical Support**
CCSS Statistical Center: Yes
Local institutional statistician:
If local, please provide the name(s) and contact information of the statistician(s) to be involved.
Will this project utilize CCSS biologic samples?: No
If yes, which of the following?:
If other, please explain:

**Section: Other General Comments**
Other General Comments:
I agree to share this information with St. Jude: Yes