**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: **Psychosocial and Neurocognitive Outcomes in Survivors of Hodgkin Lymphoma**

Planned research population (eligibility criteria):  
**Survivors with a primary diagnosis of Hodgkin Lymphoma and all siblings who completed FU2 or FU5.**

Proposed specific aims:  
1) **Describe the self-reported psychosocial and neurocognitive outcomes of survivors of a primary diagnosis of childhood Hodgkin Lymphoma in reference to sibling controls.**  
2) **Identify demographic and treatment factors that predict risks for poor psychosocial and neurocognitive outcomes.**  
3) **Determine if potential late effects, such as pulmonary or cardiac features, are predictive of risk for poor psychosocial and neurocognitive outcomes in survivors of childhood Hodgkin Lymphoma.**

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

**Section: Outcomes or Correlative Factors**

- Late mortality:
- Second Malignancy:

**Group: Health Behaviors**

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

**Group: Psychosocial**

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

**Group: Medical Conditions**

- Hearing/Vision/Speech:
- Hormonal systems:
- Heart and vascular: Correlative Factors
- Respiratory: Correlative Factors
- Digestive:
- Surgical procedures:
- Brain and nervous system: Primary
- Other:
  - If other, please specify:

**Group: Medications**
Describe medications:

**Group: Psychologic/Quality of Life**
BSI-18: Primary
SF-36: Primary
CCSS-NCQ: Primary
PTS:
PTG:
Other:
If other, please specify:

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

**Group: Demographic**
Age: Correlative Factors
Race: Correlative Factors
Sex: Correlative Factors
Other:
If other, please specify:

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

**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: