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 :