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: Subsequent neoplasms among childhood cancer survivors exposed to chemotherapy and not radiation

Planned research population (eligibility criteria):

Cohort members treated between 1970 and 1999, treated with chemotherapy but not therapeutic radiation for their childhood malignancy (N=7313)

Proposed specific aims:

- 1. Describe the cumulative incidence of subsequent neoplasms (SN) and the risk for subsequent malignant neoplasms (SMN).
- 2. Identify chemotherapeutic exposures associated with SMN risk, based on:
- a. cumulative dose of chemotherapeutic agent delivered
- b. combinations of chemotherapeutic agents
- c. timing of the occurrence of the SMN from exposure to particular agents
- 3. Describe specific SN patterns and risk factors based on therapeutic and clinical factors with consideration for SNs not considered to be radiation-related vs. those considered to be radiation-related.

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: Primary

Chronic Disease:

Psychology / Neuropsychology:

Genetics:

Cancer Control:

Epidemiology / Biostatistics : Secondary

Section: Outcomes or Correlative Factors

Late mortality:

Second Malignancy: Primary

Group: Health Behaviors

Tobacco: 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:

If other, please specify:

Group: MedicationsDescribe medications:

Group: Psychologic/Quality of Life

BSI-18 : SF-36 :

CCSS-NCQ:

PTS : PTG : Other :

If other, please specify:

Group: Other

Pregnancy and offspring:

Family history:

Chronic conditions (CTCAE v3):

Health status:

Group: Demographic

Age: Correlative Factors

Race: Correlative Factors

Sex: Correlative Factors

Other:

If other, please specify:

Group: Cancer treatment

Chemotherapy: Correlative Factors

Radiation therapy:

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: