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: Esophageal strictures after childhood cancer therapy: Experience from three childhood cancer survivor cohorts

Planned research population (eligibility criteria):
Survivors who finished at least one questionnaire that is more resent than the original baseline questionnaire. The same criteria will apply to the siblings included.

Proposed specific aims:
Describe the pattern of esophageal stricture among survivors in the CCSS, the Adult Life after Childhood Cancer in Scandinavia (ALiCCS) study, and the St Jude Lifetime Cohort study (SJLife).

Investigate associations between specific treatment exposures and the risk of esophageal strictures.
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 : Primary

Psychology / Neuropsychology :

Genetics :

Cancer Control :

Epidemiology / Biostatistics :

**Section: Outcomes or Correlative Factors**

Late mortality : Correlative Factors

Second Malignancy :

**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: *Primary*

Surgical procedures: *Secondary*

Brain and nervous system:

Other:

If other, please specify:

**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:

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: 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: 
As noted under the aims of the study, this study is intended as a joint publication from the CCSS, ALiCCS, and SJLife studies.