

## **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 : **Trends in life expectancy and quality-adjusted life expectancy among childhood cancer survivors**

Planned research population (eligibility criteria) :  
**Original and Expansion Cohort participants**

Proposed specific aims :

**1. Estimate the cumulative effect of disease- and treatment-related mortality risks on survivor life expectancy by treatment era**

**2. Assess the cumulative effect of late-effects on quality-adjusted life expectancy by treatment era**

**3. Explore diagnosis and/or treatment exposure-specific variations in survivor life expectancy and quality-adjusted life expectancy**

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? :  
**This project will not require non-CCSS fund to complete. However, we may seek external funding to support the project.**

***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 : **Secondary**

Psychology / Neuropsychology : **Secondary**

Genetics :

Cancer Control :

Epidemiology / Biostatistics : **Primary**

**Section: Outcomes or Correlative Factors**

Late mortality : **Primary**

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 :

Surgical procedures :

Brain and nervous system :

Other :

If other, please specify :

**Group: Medications**

Describe medications :

**Group: Psychologic/Quality of Life**

BSI-18 :

SF-36 : **Primary**

CCSS-NCQ :

PTS :

PTG :

Other :

If other, please specify :

**Group: Other**

Pregnancy and offspring :

Family history :

Chronic conditions (CTCAE v3) : **Primary**

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. :

**For the analysis on life expectancy, we will request excess mortality risk estimates from the CCSS Statistical Center. We will incorporate these risk estimates into a simulation model to project LE.**

**For the analysis on quality-adjusted life expectancy, we will request individual-level CCSS data. We will then use this data to develop a simulation model capable of projecting QALE.**

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 :