**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**: Socioeconomic and Rural/Urban Differences in Adverse Outcomes Among Childhood Cancer Survivors

**Planned research population (eligibility criteria)**:

The study population will include survivors in the expansion CCSS cohort who had complete information for socioeconomic status and zip code in the baseline questionnaire.

**Proposed specific aims**:

- **Aim #1**: To evaluate differences in overall and cause-specific mortality by household income
- **Aim #2**: Evaluate differences in overall and cause-specific mortality by rurality
- **Aim #3**: Assess effect modification of the relationship between mortality and rurality by household income

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:
Genetics:
Cancer Control:
Epidemiology / Biostatistics: Primary

**Section: Outcomes or Correlative Factors**

Late mortality: Primary
Second Malignancy: Secondary

**Group: Health Behaviors**

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

**Group: Psychosocial**

Insurance: Correlative Factors
Marriage:
Education: Correlative Factors
Employment: Correlative Factors
Other:
If other, please specify:

**Group: Medical Conditions**

Hearing/Vision/Speech:
Hormonal systems:
Heart and vascular: Secondary
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:
CCSS-NCQ:
PTS:
PTG:
Other:
If other, please specify:

**Group: Other**

Pregnancy and offspring:
Family history:
Chronic conditions (CTCAE v3): **Secondary**
Health status: **Secondary**

**Group: Demographic**

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

**Group: Cancer treatment**

Chemotherapy:
Radiation therapy:
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:
This proposed analysis would be mentored/supervised by Dr. Smita Bhatia.
I agree to share this information with St. Jude: **Yes**