**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: **Cost-effectiveness of the Children’s Oncology Group Colorectal Cancer Screening Guidelines for Childhood Cancer Survivors**  
Planned research population (eligibility criteria): Childhood cancer survivors  
Proposed specific aims:  
1. Examine the efficacy and cost-effectiveness of the COG Guidelines colorectal cancer screening strategies  
2. Explore alternative colorectal cancer screening schedules that might be more cost-effective  
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 :
Section: Outcomes or Correlative Factors

Late mortality: Primary
Second Malignancy: Primary

Group: Health Behaviors
Tobacco:
Alcohol:
Physical activity:
Medical screening: Correlative Factors
Other:
If other, please specify:

Group: Psychosocial
Insurance: Correlative Factors
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:
CCSS-NCQ:
PTS:
PTG:
Other:
If other, please specify:

Group: Other
Pregnancy and offspring:
Family history: Correlative Factors
Chronic conditions (CTCAE v3):

Health status:

**Group: Demographic**
- Age: Correlative Factors
- Race: Correlative Factors
- Sex: Correlative Factors
- Other: Correlative Factors

If other, please specify: Age at diagnosis of the primary malignancy; age at death (and cause of death); age at last survey

**Group: Cancer treatment**
- Chemotherapy: Correlative Factors
- Radiation therapy: Correlative Factors
- Surgery:

**Section: Anticipated Sources of Statistical Support**
- CCSS Statistical Center: Yes
- Local institutional statistician: Yes

If local, please provide the name(s) and contact information of the statistician(s) to be involved.

The principal investigator will execute the statistical analyses under the supervision of Dr. Smita Bhatia.

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 is a revised submission after my mentor, Dr. Smita Bhatia, consulted with the CCSS senior investigator(s).

I agree to share this information with St. Jude: Yes