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: Nonmelanoma Skin Cancer in Survivors of Childhood and Adolescent Cancer: An update from the Childhood Cancer Survivor Study
Planned research population (eligibility criteria) :
CCSS participants of any age
Proposed specific aims :
1. Identify the overall incidence of NMSC and incidence of developing multiple NMSC in the CCSS study group.
2. Identify risk factors for development of NMSC in CCSS participants.
3. Compare the age-adjusted incidence of NMSC in CCSS participants based on era of primary cancer treatment.
4. Assess the impact of attained age on risk of skin cancer in the original patient cohort.
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 :
Section: Outcomes or Correlative Factors

Late mortality:
Second Malignancy: Primary

Group: Health Behaviors
Tobacco: Correlative Factors
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: Primary
If other, please specify: 1. Non-melanoma skin cancer history (basal cell and cutaneous squamous cell carcinoma. 2. History of underlying genetic disease 3. History of bone marrow transplantation

Group: Medications
Describe medications:
1. Use of chronic immunosuppressive medications.
2. Use of voriconazole

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:
If other, please specify: Region of US where cancer treatment was delivered

**Group: Cancer treatment**
Chemotherapy: Correlative Factors
Radiation therapy: Correlative Factors
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:
I agree to share this information with St. Jude: Yes