**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: **Cardiometabolic outcomes among survivors of childhood and young adult cancer**

Planned research population (eligibility criteria):  
1. Survivors of childhood and young adult cancer from the original and expanded CCSS cohorts.

Proposed specific aims:  
1. Describe the prevalence of cardiometabolic disease including: hypertension, hyperlipidemia, obesity, and diabetes mellitus.  
2. Examine treatment-related risk factors for cardiometabolic outcomes and resolve ongoing debate, such as the role of dexamethasone.

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: **Primary**
Psychology / Neuropsychology: **Primary**
Genetics: **Primary**
Cancer Control: **Secondary**
Epidemiology / Biostatistics: **Secondary**

**Section: Outcomes or Correlative Factors**

Late mortality: **Primary**
Second Malignancy: **Secondary**

**Group: Health Behaviors**

Tobacco: **Primary**
Alcohol: **Secondary**
Physical activity: **Correlative Factors**
Medical screening: **Other**
Other: **Other**
If other, please specify: **Other**

**Group: Psychosocial**

Insurance: **Primary**
Marriage: **Primary**
Education: **Primary**
Employment: **Secondary**
Other: **Secondary**
If other, please specify: **Secondary**

**Group: Medical Conditions**

Hearing/Vision/Speech: **Primary**
Hormonal systems: **Primary**
Heart and vascular: **Primary**
Respiratory: **Primary**
Digestive: **Primary**
Surgical procedures: **Primary**
Brain and nervous system: **Primary**
Other: **Primary**
If other, please specify: **Primary**

**Group: Medications**

Describe medications: **Anti-hypertensives**
**Lipid-lowering medications**
Metformin, insulin, and other anti-diabetes drugs

**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: **Primary**
  - If other, please specify: **Body Mass Index (height and weight)**

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

**Section: Anticipated Sources of Statistical Support**
- CCSS Statistical Center:
- Local institutional statistician: **Yes**
  - If local, please provide the name(s) and contact information of the statistician(s) to be involved.
    - **Chaya Moskowitz, PhD**
    - moskowc1@mskcc.org
- 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: