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: Disparities in Adherence to Screening Guidelines in Hodgkin's Lymphoma Survivors

Planned research population (eligibility criteria):
Hodgkin's Lymphoma survivors

Proposed specific aims:
1. Describe adherence to recommended screening guidelines among survivors of Hodgkin's Lymphoma by race, ethnicity, and socioeconomic factors
2. Identify disparities that exist in adherence to recommended screening
3. Determine whether racial/ethnic, socioeconomic, other demographic or disease related characteristics predict risk of noncompliance with screening
4. Identify populations at risk for noncompliance to develop future targeted interventions in these groups.
5. Assess whether disparities have changed over time using current and past questionnaires

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

**Section: Outcomes or Correlative Factors**

Late mortality: Primary
Second Malignancy: Primary

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

**Group: Psychosocial**
Insurance: Correlative Factors
Marriage: Correlative Factors
Education: Correlative Factors
Employment: Correlative Factors
Other: Correlative Factors
If other, please specify: SES, Race-Ethnicity, Education, Demographics

**Group: Medical Conditions**
Hearing/Vision/Speech:
Hormonal systems: Primary
Heart and vascular: Primary
Respiratory: Primary
Digestive: Primary
Surgical procedures:
Brain and nervous system:
Other: Primary
If other, please specify: Skin, Breast, Thyroid, Diabetes

**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: Secondary
- Family history:
- Chronic conditions (CTCAE v3): Primary
- Health status:

**Group: Demographic**
- Age: Correlative Factors
- Race: Primary
- 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.
- 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 project will take place under the mentorship of Dr. Tara Henderson, Dr. Paul Nathan, and Dr. Kevin Oeffinger who have participated in the inception of the proposal.
  - I agree to share this information with St. Jude: Yes