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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: HPV-related SMNs in childhood cancer survivors
Planned research population (eligibility criteria): Whole cohort
Proposed specific aims: 1. Describe the cumulative incidence of HPV-related cancers as SMN (oropharyngeal, penile, vaginal, vulvar, anal and cervical) in childhood cancer survivors. 2. Compare the risk of HPV-related SMN in childhood cancer survivors as compared to among age and sex matched population controls (SEER data).
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?:

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.: 

What CCSS Working Group(s) would likely be involved? (Check all that apply)

Second Malignancy: Primary
Chronic Disease:
Psychology / Neuropsychology:
Genetics:
To describe the anticipated scope of the study, please indicate the specific CCSS data to be included as **outcome** (primary or secondary) or **correlative factors**. (Check all that apply)

**Late mortality:**
Second Malignancy: Primary

**Health Behaviors**

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

**Psychosocial**

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

**Medical conditions**

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

**Medications**

Describe medications:
Pregnancy and offspring:
Family History:

Psychologic/Quality of Life

BSI-18:
SF-36:
CCSS-NCQ:
PTS:
PTG:
Other:
If other, please specify:

Chronic conditions (CTCAE v3):
Health status:

Demographic

Age: Correlative Factors
Race: Correlative Factors
Sex: Correlative Factors
Others:
If others, please specify:

Cancer treatment

Chemotherapy: Correlative Factors
Radiation therapy: Correlative Factors
Surgery: Correlative Factors

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?

Buccal cell DNA:
Peripheral blood:
Lymphoblastoid cell lines:
Second malignancy pathology samples:
Other requiring collection of samples:
If other, please explain:

Other general comments: