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: Infertility, assisted reproductive technology utilization and pregnancy outcomes in childhood cancer survivor population: A CCSS and SART CORS (Society for Assisted Reproductive Technology Clinic Outcome Reporting Study) data linkage study

Planned research population (eligibility criteria): All CCSS participants.

Proposed specific aims:
1. Determine utilization rate of ART in childhood cancer survivors.
2. Compare pregnancy and live birth rates in childhood cancer survivors who pursue ART to general and specific IVF populations.
3. Compare pregnancy outcomes for childhood cancer survivors pursuing ART (congenital anomalies, gestational age and birth weight) to outcomes in general and specific IVF populations.

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

**Section: Outcomes or Correlative Factors**

Late mortality:
Second Malignancy:

**Group: Health Behaviors**

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

**Group: Psychosocial**

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

**Group: Medical Conditions**

Hearing/Vision/Speech:
Hormonal systems: **Correlative Factors**
Heart and vascular:
Respiratory:
Digestive:
Surgical procedures: **Correlative Factors**
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: Primary
Family history:
Chronic conditions (CTCAE v3): Correlative Factors
Health status:

**Group: Demographic**
Age: Correlative Factors
Race: Correlative Factors
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: