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 : **Long-term Outcomes of Pelvic Sarcoma Based on Initial Treatment**
Planned research population (eligibility criteria) :
Five-year childhood cancer survivors (baseline cohort) diagnosed with sarcoma of the pelvis at age <21.
Proposed specific aims :
S.A. 1: To describe the difference in mortality among childhood pelvic sarcoma survivors treated with A) surgery alone vs. B) radiotherapy alone vs. C) surgery and radiotherapy.

S.A. 2: To describe the difference in late effects* among childhood pelvic sarcoma survivors treated with A) surgery alone vs. B) radiotherapy alone vs. C) surgery and radiotherapy.
***Late effects include functional outcomes (e.g. physical function, fertility, bowel/bladder), recurrence/second neoplasm, and CTCAE chronic health conditions.**

S.A. 3: To identify risk factors (in addition to surgery and radiotherapy) for the above primary (i.e. mortality) and secondary outcomes among childhood pelvic sarcoma survivors.
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 : **Secondary**
Chronic Disease : **Primary**
Psychology / Neuropsychology :
Genetics :
Cancer Control :
Epidemiology / Biostatistics : **Secondary**

Section: Outcomes or Correlative Factors

Late mortality : **Primary**
Second Malignancy : **Secondary,Correlative Factors**

Group: Health Behaviors

Tobacco : **Correlative Factors**
Alcohol : **Correlative Factors**
Physical activity : **Secondary**
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 : **Secondary**
Heart and vascular : **Secondary**
Respiratory :
Digestive : **Secondary**
Surgical procedures : **Secondary,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 : **Secondary**
Family history :
Chronic conditions (CTCAE v3) : **Secondary**
Health status : **Secondary**

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 :