First Name: Tara Last Name: Brinkman Institution: SJCRH

Address 1: 262 Danny Thomas Place

Address 2: City: Memphis State/Province: TN

Country: Zip: 38105

Phone: 901-595-5683 Alternate Phone:

Email: tara.brinkman@stjude.org

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: Social and behavioral phenotypes of adolescent survivors of childhood cancer Planned research population (eligibility criteria): Adolescents (aged 12-17 years) in the baseline expansion cohort with data from the behavior problem index

Proposed specific aims: Aim 1. To identify social and behavioral phenotypes through examination of symptom patterns and comorbidities (i.e. clusters of co-occurring symptoms) Aim 2. To investigate the association between sociodemographic characteristics and treatment late effects (e.g., scarring/disfigurement, sensory impairment, pain, health status) and adolescent social and behavioral phenotypes. Aim 3. To compare adolescent social and behavioral problems in the expansion cohort (diagnosed between 1987-1999) to the original CCSS cohort (diagnosed between 1970-1896) as well as to siblings from the original cohort.

Will the project require non-CCSS funding to complete?: No

If ves, 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:
Chronic Disease:
Psychology / Neuropsychology: Primary
Genetics:
Cancer Control:
Epidemiology / Biostatistics:
To describe the anticipated scope of the study, please indicate the specific CCSS data to be included as <u>outcome</u> (primary or secondary) or <u>correlative factors</u> . (Check all that apply)
Late mortality:
Second Malignancy:
Health Behaviors
Tobacco:
Alcohol:
Physical activity:
Medical screening:
Other:
If other, please specify:
Psychosocial
Insurance:
Marriage:
Education: Secondary
Employment:
Other: Secondary
If other, please specify: Household income
Medical conditions
Hearing/Vision/Speech: Primary
Hormonal systems:
Heart and vascular:
Respiratory:
Digestive:
Surgical procedures: Primary
Brain and nervous system: Primary
Other: Primary
If other, please specify: Scarring/disfigurement; body mass index; perceived physical health status; perceived as disabled
Medications

Describe medications:
Pregnancy and offspring: Family History:
Psychologic/Quality of Life
BSI-18:
SF-36:
CCSS-NCQ:
PTS:
PTG:
Other: Primary
If other, please specify: Behavior Problem Index
Chronic conditions (CTCAE v3):
Health status:
Demographic
Age: Secondary
Race: Secondary
Sex: Secondary
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: