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: Patterns of motor and sensory neuropathy and associated outcomes in long term survivors of childhood cancer

Planned research population (eligibility criteria):

The study population will consist of all CCSS cases (diagnosed 1970-1999) who were alive and completed the CCSS Baseline survey as well as the sibling comparison group. A subset analysis will require that cases also complete at least one Follow-up survey.

Proposed specific aims:

Aim 1: To determine the prevalence and patterns of motor and sensory neuropathy in CCSS cases overall, and by diagnosis in comparison to the sibling group.

Aim 2: To determine patient and treatment characteristics associated with motor and sensory neuropathy in CCSS cases.

Aim 3: To determine the association of sensory and motor neuropathy with the following:

- Sedentary levels of physical activity
- Use of pharmacologic (i.e. analgesics, neuromodulators, muscle relaxants) and non-pharmacologic (i.e. physical therapy) therapeutic interventions
- Psychosocial outcomes (i.e. anxiety, depression)
- Employment among adult CCSS cases, including employment status, sick days

per year, and income

Aim 4: To determine longitudinal patterns in the severity of motor and sensory neuropathy in the subset of CCSS cases who have at least one completed Follow-up Survey

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 : **Yes**

If yes to any of the above, please briefly describe. : Yes, chemotherapy type and cumulative dose

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

Second Malignancy:

Chronic Disease : Primary

Psychology / Neuropsychology : **Secondary**

Genetics:

Cancer Control : **Secondary** Epidemiology / Biostatistics :

Section: Outcomes or Correlative Factors

Late mortality:

Second Malignancy:

Group: Health Behaviors

Tobacco: Alcohol:

Physical activity: Secondary

Medical screening:
Other: Secondary

If other, please specify: physical therapy

Group: Psychosocial Insurance: Secondary

Marriage:

Education : Secondary
Employment : Secondary

Other:

If other, please specify:

Group: Medical Conditions

Hearing/Vision/Speech : Hormonal systems : Heart and vascular :

Respiratory:
Digestive:

Surgical procedures:

Brain and nervous system: Primary

Other:

If other, please specify: **Group: Medications**Describe medications:

analgesics, neuromodulators, muscle relaxants

Group: Psychologic/Quality of Life

BSI-18: Secondary

SF-36:

CCSS-NCQ:

PTS: PTG: Other:

If other, please specify:

Group: Other

Pregnancy and offspring:

Family history:

Chronic conditions (CTCAE v3): Primary

Health status:

Group: Demographic

Age: Correlative Factors

Race: Correlative Factors

Sex: Correlative Factors

Other: Correlative Factors

If other, please specify:

Group: Cancer treatment

Chemotherapy : Correlative Factors
Radiation therapy : Correlative Factors

Surgery:

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:

I am starting my 2nd year of pediatric hematology-oncology fellowship and will have 2 dedicated research years without clinical responsibilities to focus on this project. I will be working with Dr. Nina Kadan-Lottick as a mentor (Nina.kadan-lottick@yale.edu).

I agree to share this information with St. Jude: Yes