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: Prevalence and Pattern of Prescription Psychoactive Medication Use in Adolescent Survivors of Childhood Cancer: A Report from the Childhood Cancer Survivor Study

Planned research population (eligibility criteria):

(1) All cancer survivors and siblings <18 years of age whose parents completed the Baseline survey, and (2) survivors and siblings from the Original Cohort who completed the 2003 or 2007 Follow-up survey.

Proposed specific aims:

We propose to examine prevalence and patterns of psychoactive medication use in adolescent survivors of childhood cancer. Previous CCSS publications by Brinkman et al, examined this outcome only in survivors >=18 years old and, thus, information on use of such medications by children is undocumented. We are particularly interested in whether use has increased or decreased over the decades from 1970-79, 1980-89, and 1990-99.

Specific Aim 1: To estimate the prevalence of psychoactive medication use in a large and geographically diverse cohort of adolescent survivors of childhood cancer

Specific Aim 2: To identify demographic, diagnosis, treatment and psychological

factors associated with psychoactive medication use in adolescent survivors of childhood cancer

Specific Aim 3: To examine associations between psychoactive medication use and academic functions, as reflected through special education services

Specific Aim 4: In the original cohort, to examine the impact of psychoactive mediation use during adolescence and psychologic and quality of life outcomes during adulthood

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:

Psychology / Neuropsychology: Primary

Genetics:

Cancer Control:

Epidemiology / Biostatistics:

Section: Outcomes or Correlative Factors

Late mortality:

Second Malignancy:

Group: Health Behaviors

Tobacco : Alcohol :

Physical activity:
Medical screening:

Other:

If other, please specify:

Group: Psychosocial

Insurance: Correlative Factors

Marriage:

Education: Correlative Factors

Employment:

Other: Correlative Factors

If other, please specify: Household income

Group: Medical Conditions

Hearing/Vision/Speech : Hormonal systems :

Heart and vascular:

Respiratory : Digestive :

Surgical procedures:

Brain and nervous system: Correlative Factors

Other:

If other, please specify:

Group: Medications

Describe medications:

Outcomes: Psychoactive medications - (1) antidepressants, (2) anxiolytics/sedatives/hypnotics, (3) anticonvulsants, (4) non-opioid analgesics, (5) opioids, (6) muscle relaxants, (7) neuroleptics, and (8) stimulants.

Group: Psychologic/Quality of Life

BSI-18 : Correlative Factors
SF-36 : Correlative Factors

CCSS-NCQ : Correlative Factors

PTS : Correlative Factors
PTG : Correlative Factors
Other : Correlative Factors

If other, please specify: **Brief Problem Inventory**

Group: Other

Pregnancy and offspring:

Family history:

Chronic conditions (CTCAE v3) : Health status : Correlative Factors

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:

Local institutional statistician: Yes

If local, please provide the name(s) and contact information of the statistician(s) to be involved. :

Dr Kumar Srivastava's statistical team at St. Jude Children's Research Hospital

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: