

CCSS Application of Intent

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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	Immune Deficits in Survivors of Childhood Cancer
Planned research population (eligibility criteria)	1) Survivors with thyroid or meningioma SMN 2) Survivors matched by treatment exposures, and with no SMN and no grade 3/4 CHC (collected prospectively as part of the CCSS supplemental blood banking recruitment study)
Proposed specific aims	1) To characterize the prevalence and type of immune deficits in long-term radiation-exposed survivors of childhood cancer with meningioma or thyroid SMN, compared to matched survivors with no SMN 2) To determine host and disease-related factors that are associated with immune deficits in long-term survivors of childhood cancer 3) To identify genetic and molecular mechanisms underlying immune deficits in long-term survivors of childhood cancer
Will the project require non-CCSS funding to complete?	Yes
What would be the anticipated source(s) and timeline(s) for securing funding?	NCI R01 (MPI with Kavita Dhodapkar, Emory and Smita Bhatia, UAB)

Additional self-reported information No

Biological samples No

Medical record data No

If yes to any of the above, please briefly describe.

WORKING GROUPS

Second Malignancy Primary

Chronic Disease

Psychology /
Neuropsychology

Genetics Secondary

Cancer Control

Epidemiology /
Biostatistics

OUTCOMES/CORRELATIVE

FACTORS

Late Mortality

Second Malignancy Primary

HEALTH BEHAVIORS

Tobacco

Alcohol

Physical Activity

Medical Screening

Other

If other, please specify

PSYCHOSOCIAL

Insurance Correlative Factors

Marriage Correlative Factors

Education Correlative Factors

Employment Correlative Factors

Other

If other, please specify

MEDICAL CONDITIONS

Hearing/Vision/Speech

Hormonal Systems

Heart and Vascular

Respiratory	
Digestive	
Surgical Procedures	
Brain and Nervous System	
Other	
If other, please specify	
MEDICATIONS	
Describe medications	
PSYCHOLOGIC/QOL	
BSI-18	
SF-36	
CCSS-NCQ	
PTS	
PTG	
Other	
If other, please specify	
OTHER	
Pregnancy and Offspring	
Family History	Correlative Factors
Chronic Conditions (CTCAE v3)	Correlative Factors
Health Status	Correlative Factors
DEMOGRAPHIC	
Age	Correlative Factors
Race	Correlative Factors
Sex	Correlative Factors
Other	Correlative Factors
If other, please specify	
CANCER TREATMENT	
Chemotherapy	Correlative Factors
Radiation Therapy	Correlative Factors
Surgery	
<u>STATISTICAL SUPPORT</u>	CCSS Statistical Center Local institutional statistician
If local, please provide the name(s) and contact information of the	Undecided, if local will be Melissa Richard, PhD

statistician(s) to be involved.

Will this project utilize CCSS biologic samples?

Yes

If yes, which of the following?

Peripheral blood

If other, please explain

Some being collected prospectively under supplemental funding protocol (controls without grade 3/4 CHC and additional survivors with SMN history)

Agree

I agree to share this information with St. Jude