Name Tara Brinkman

Institution St. Jude Children's Research Hospital

Address 262 Danny Thomas Place, MS 740

Memphis, TN, 38105

United States

Phone Number 9015955683

Alternate Phone Number

Email Address 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

Memory Problems in Aging Adult Survivors of Childhood Cancer

Planned research population (eligibility criteria)

Survivors who completed CNS Vital Signs (CNS VS) and CCSS NCQ. Siblings may be used for prevalence estimates in Aim 1. Aim 3 will require completion of NCQ at two time points.

Proposed specific aims

- 1. Estimate the prevalence and identify patterns of performance-based (CNS VS) and self-reported (NCQ) memory problems in a large cohort of adult survivors of pediatric cancer
- 2. Identify mechanisms (i.e., diagnosis/ treatment exposures, lifestyle factors, chronic health conditions) associated with patterns (i.e., performance-based only, self-report only, performance-based & self-report) of memory problems in adult survivors
- 3. Identify the contribution of performance-based neurocognitive skills (memory, attention, processing speed, executive function) to self-reported memory decline
- 4. Estimate the impact of patterns of memory impairment on functional outcomes (including ability to participate in social roles/activities and complete activities of daily living)

If yes, 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? (Select all that apply)

	Primary	Secondary
Second Malignancy		
Chronic Disease		✓
Psychology/Neuropsychology	✓	
Genetics		
Cancer Control		
Epidemiology/Biostatistics		

Outcomes or Correlative Factors

	Primary	Secondary	Correlative Factors
Late Mortality			
Second Malignancy			

Health Behaviors

	Primary	Secondary	Correlative Factors
Tobacco			✓
Alcohol			✓
Physical Activity			✓
Medical Screening			
Other			

If other, please specify

Psychosocial

	Primary	Secondary	Correlative Factors
Insurance			✓
Marriage			✓
Education			✓
Employment			✓
Other			

If other, please specify

Medical Conditions

	Primary	Secondary	Correlative Factors
Hearing/Vision/Speech			✓
Hormonal Systems			
Heart and Vascular			✓
Respiratory			✓
Digestive			
Surgical Procedures			
Brain and Nervous System			✓
Other			

Medications

Describe medications

Psychologic/Quality of Life

	Primary	Secondary	Correlative Factors
BSI-18			✓
SF-36			
CCSS-NCQ	✓		
PTS			
PTG			
Other			

If other, please specify

CNS Vital Signs

Other

	Primary	Secondary	Correlative Factors
Pregnancy and Offspring			
Family History			
Chronic Conditions (CTCAE v3)			✓
Health Status			

Demographic

	Primary	Secondary	Correlative Factors
Age			✓
Race			✓
Sex			✓
Other			

If other, please specify

Cancer Treatment

	Correlative Factors
Chemotherapy	✓
Radiation Therapy	✓
Surgery	✓

Anticipated Sources of Statistical Support

CCSS Statistical Center	Yes
Local Institutional Statistician	No

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

Other General Comments

Agree

I agree to share this information with St. Jude

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