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Project Requirements and Description

Requirements to submit AOI

Requirements to submit AOI (all answers must be "yes" to proceed)

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 Promoting Resilience in Childhood, Adolescent and Young Adult Survivors

Planned research population (eligibility criteria)

Survivors of Pediatric or AYA cancers (not currently undergoing cancer treatment)

Proposed specific aims

Aim 1: Determine the effectiveness of two digital resilience interventions (SMART-3RP-S and SMART-3RP-A) in improving resilience (primary outcome) among posttreatment CAYA survivors. We will also compare the effects of both digital interventions on coping, emotional distress (anxiety/depression, uncertainty

intolerance), physical health (physical functioning), and quality of life (secondary outcomes).

Aim 2: To explore moderators (e.g., age, race/ethnicity, language, zip code, cancer/medical history, baseline distress) of the intervention effect on resilience at 6 and 12 months (Aim 2a); To assess if improvements in resilience are associated with improvements in healthcare use at 12-months (Aim 2b).

Aim 3: To assess intervention reach, engagement, acceptability, fidelity, sustainability, and cost-effectiveness.

Will the project require non-CCSS funding to complete? Yes

If yes, what would be the anticipated source(s) and timeline(s) for securing funding?

NCI funding

Does this project require contact of CCSS study subjects for:

Additional self-reported information	Yes
Biological samples	No
Medical record data	Yes

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

healthcare use information

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

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

Outcomes or Correlative Factors

Late Mortality	
Second Malignancy	Correlative Factors

Health Behaviors

Tobacco	
Alcohol	
Physical Activity	Correlative Factors
Medical Screening	Correlative Factors

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	Correlative Factors

If other, please specify

any other chronic medical conditions will be collected and potentially examined as moderators

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Medications

Describe medications

n/a

Psychologic/Quality of Life

BSI-18	
SF-36	
CCSS-NCQ	
PTS	
PTG	

Other	Primary

If other, please specify

PROMIS measures for Anxiety, depression, Intolerance of Uncertainty-12, MOCS-A for coping, Quality of Life (maybe assess with SF36), Resilience

Other

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

Demographic

Age	
Race	
Sex	
Other	Secondary

If other, please specify

These will be examined as moderators (and we hope to oversample)

Cancer Treatment

Chemotherapy	
Radiation Therapy	
Surgery	

Anticipated Sources of Statistical Support

CCSS Statistical Center	No
Local Institutional Statistician	Yes

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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