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Requirements to submit AOI:

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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

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Project Title: Psychosocial and physical factors influencing fatigue in survivors of childhood Hodgkin Lymphoma.  
Planned research population (eligibility criteria): Survivors of Hodgkin Lymphoma.  
Proposed specific aims: Aim 1: To investigate the association between pain, emotional distress, and physical functioning on levels of fatigue in survivors of childhood Hodgkin Lymphoma. Aim 2: To investigate the influence of survivor demographics and treatment characteristics on the association between pain, emotional distress, and physical functioning on levels of fatigue in survivors of childhood Hodgkin Lymphoma.  
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?:

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Does this project require contact of CCSS study subjects for . . .

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Additional self-reported information: No  
Biological Samples: No  
Medical record data: No  
If yes to any of the above, please briefly describe.:

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

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Second Malignancy:  
Chronic Disease:

Psychology / Neuropsychology: Primary  
Genetics:  
Cancer Control:  
Epidemiology / Biostatistics: Secondary

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To describe the anticipated scope of the study, please indicate the specific CCSS data to be included as outcome (primary or secondary) or correlative factors. (Check all that apply)

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Late mortality:  
Second Malignancy:

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#### Health Behaviors

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Tobacco:  
Alcohol:  
Physical activity: Primary  
Medical screening:  
Other:  
If other, please specify:

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#### Psychosocial

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Insurance:  
Marriage:  
Education:  
Employment:  
Other:  
If other, please specify:

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#### Medical conditions

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Hearing/Vision/Speech:  
Hormonal systems:  
Heart and vascular:  
Respiratory:  
Digestive:  
Surgical procedures:  
Brain and nervous system:  
Other:  
If other, please specify:

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#### Medications

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Describe medications:

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Pregnancy and offspring:  
Family History:

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Psychologic/Quality of Life

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BSI-18: Primary  
SF-36: Primary  
CCSS-NCQ:  
PTS:  
PTG:  
Other:  
If other, please specify:

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Chronic conditions (CTCAE v3):  
Health status:

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Demographic

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Age: Correlative Factors  
Race: Correlative Factors  
Sex: Correlative Factors  
Others:  
If others, please specify:

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Cancer treatment

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Chemotherapy: Correlative Factors  
Radiation therapy: Correlative Factors  
Surgery: Correlative Factors

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Anticipated sources of statistical support

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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

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If yes, which of the following?

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Buccal cell DNA:  
Peripheral blood:

Lymphoblastoid cell lines:  
Second malignancy pathology samples:  
Other requiring collection of samples:  
If other, please explain:

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Other general comments: Data will come from the “2003 Follow-up Survey” and the “Sleep Survey” which includes the FACIT, Pittsburg Sleep Index, and Epworth Sleepiness scale.

Dr. Crabtree will be the mentor, and assume the role of PI for this study as Amanda Rach who proposed the study is a graduate student in the Department of Psychology at the University of Memphis