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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: BMI in survivors and siblings in cohort 1 vs 2: differences and risk factors  
Planned research population (eligibility criteria): Phase 1 and 2 CCSS survivors and siblings  
Proposed specific aims: 1. Describe the BMI status distribution in survivors cohort 2 and compare to the general population 2. Compare distribution of BMI status in survivors from cohort 1 vs 2 by diagnosis 3. Identify treatment and demographic risk factors for BMI > 30 and < 18.5 in survivors from cohort 2  
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: Primary  
Psychology / Neuropsychology:  
Genetics:

Cancer Control:  
Epidemiology / Biostatistics:

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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: Correlative Factors  
Alcohol: Correlative Factors  
Physical activity: Correlative Factors  
Medical screening:  
Other:  
If other, please specify:

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Psychosocial

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

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Medical conditions

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

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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:  
SF-36:  
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: This would be a follow up paper to the 2005 Cancer manuscript: Body Mass Index in Long-Term Adult Survivors of Childhood Cancer but this time comparing cohort 1 and 2. This was discussed with Dr Oeffinger and Sklar at the 2013 CCSS meeting