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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: The health system use and cost implications of amputation vs. limb salvage surgery among pediatric lower-extremity bone tumor survivors  
Planned research population (eligibility criteria): Survivors of lower extremity bone tumors  
Proposed specific aims: (1) To estimate and compare the 1-year, 5-year, 10-year and 25-year health systems use and costs of amputation vs. limb salvage surgery among pediatric cancer survivors. (2) To examine year of surgery, demographics (state of residence, age at surgery, gender, years of education at surgery, year of surgery) and clinical (primary and secondary diagnosis, presence of secondary cancers, other health morbidities, treatment received prior to and following surgery) factors as potential correlates of treatment modality selection and estimated health systems use and costs among pediatric lower-extremity bone tumor survivors. (3) To compare short-term and long-term health systems use and costs to existing estimates of mortality and morbidity outcomes among pediatric lower-extremity bone tumor survivors by selected surgical modality.  
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:

Genetics:

Cancer Control: Primary

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:

Alcohol:

Physical activity:

Medical screening:

Other:

If other, please specify:

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Psychosocial

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Insurance: Correlative Factors

Marriage:

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:

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: Correlative Factors  
SF-36: Correlative Factors  
CCSS-NCQ:  
PTS:  
PTG:  
Other:  
If other, please specify:

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Chronic conditions (CTCAE v3): Correlative Factors  
Health status: Correlative Factors

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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: