**Section: Contact Information**

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**Section: Project Requirements and Description**

**Group: 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: Adherence to and Effectiveness of Survivorship Care Guidelines in CCSS  
Planned research population (eligibility criteria): All CCSS participants, however this may change based upon the data system we receive funding to support linkage to (e.g., public vs. private insurance).  
Proposed specific aims:  
Aim 1: To utilize key population- and healthcare system-based data sources (e.g., insurance claims data) to determine the patterns and predictors of COG guideline utilization in CCSS participants.  
Aim 2: To utilize key population- and healthcare system-based data sources (e.g., insurance claims data) to evaluate whether receiving COG guideline-based care improves morbidity, quality of life, and mortality among CCSS participants.  
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? : Institutional developmental funds, foundation grants (St. Baldrick’s, Alex’s Lemonade Stand).  

**Group: Does this project require contact of CCSS study subjects for:**  
Additional self-reported information: No  
Biological samples: No  
Medical record data: Yes  
If yes to any of the above, please briefly describe. : The study will need to link to claims data (e.g. Medicaid, Kaiser Permanente) for the
purposes of confirming receipt of screening tests (e.g. mammography, echocardiograms).

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

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

**Section: Outcomes or Correlative Factors**

Late mortality: Secondary 
Second Malignancy: Secondary

**Group: Health Behaviors**

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

**Group: Psychosocial**

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

**Group: Medical Conditions**

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

**Group: Medications**

Describe medications:

**Group: Psychologic/Quality of Life**

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

**Group: Other**
Pregnancy and offspring:
Family history: Correlative Factors
Chronic conditions (CTCAE v3): Secondary
Health status: Secondary

**Group: Demographic**
Age: Correlative Factors
Race: Correlative Factors
Sex: Correlative Factors
Other:
If other, please specify:

**Group: Cancer treatment**
Chemotherapy: Correlative Factors
Radiation therapy: Correlative Factors
Surgery: Correlative Factors

**Section: Anticipated Sources of Statistical Support**
CCSS Statistical Center:
Local institutional statistician: Yes
If local, please provide the name(s) and contact information of the statistician(s) to be involved.:
Yutaka Yasui
Will this project utilize CCSS biologic samples?: No
If yes, which of the following?: Other requiring collection of samples
If other, please explain:

**Section: Other General Comments**
Other General Comments:
We plan to use treatment exposures in CCSS to identify expected screening evaluations. Then we will link to claims data to track surveillance evaluations obtained. Lastly, we will look at associations with outcomes again using CCSS outcomes data.
I agree to share this information with St. Jude: Yes