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 : **EMPOWER-II**
Planned research population (eligibility criteria) :
1) Female survivors treated with chest radiation (>= 20 Gy) who are 27 - 49 years of age and have not had a mammogram AND a breast MRI in the preceding two years.

2) Primary care physicians of participants.

Proposed specific aims :
Determine the efficacy of an intervention, targeted for both female survivors and their primary care physicians, on the rates of recommended breast cancer screening (mammogram + breast MRI) compared with an attention control.
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; renewal submission date March 5, 2016

Group: 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. :
As in the EMPOWER Study, we will collect information regarding breast cancer screening from the participants at baseline and at 12-months. We will also collect faxed reports of the breast cancer screening tests (mammogram, MRI, ultrasound) completed in the study period.

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

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

Section: Outcomes or Correlative Factors

Late mortality :
Second Malignancy : **Secondary**

Group: Health Behaviors

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

Group: Psychosocial

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

Group: Medical Conditions

Hearing/Vision/Speech :
Hormonal systems :
Heart and vascular :
Respiratory :
Digestive :
Surgical procedures :
Brain and nervous system :
Other :

If other, please specify :

Group: Medications

Describe medications :

Group: Psychologic/Quality of Life

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

Group: Other

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

Group: Demographic

Age : **Secondary**
Race : **Secondary**
Sex :
Other :
If other, please specify :

Group: Cancer treatment

Chemotherapy :
Radiation therapy :
Surgery :

Section: Anticipated Sources of Statistical Support

CCSS Statistical Center : **Yes**
Local institutional statistician : **Yes**
If local, please provide the name(s) and contact information of the statistician(s) to be involved. :
Chaya Moskowitz, PhD - MSKCC
Will this project utilize CCSS biologic samples? : **No**
If yes, which of the following? :
If other, please explain :

Section: Other General Comments

Other General Comments :
This will be a competing renewal R01 with MPI of Kevin Oeffinger and Jennifer Ford.