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
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.