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 : **Loss of Sight, Hearing and Peripheral Sensation and its Effect on HRQOL and Social Functioning in Childhood Cancer Survivors Across Three Decades of Diagnosis**
Planned research population (eligibility criteria) :
Survivors of childhood cancers who self-report loss of sight, loss of hearing and loss of peripheral sensation; survivors diagnosed with cancers between 1970 and 1999
Proposed specific aims :
Aim 1: We propose to estimate the prevalence of young adult survivors of childhood cancer with the late effects of vision loss, hearing loss and loss of peripheral sensation
Aim 2: We would like to examine the impact of these late effects on HRQOL, loneliness, social functioning and life achievement.
Aim 3: Assess the long term mortality of survivors with vision loss, hearing loss and loss of peripheral sensation
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? :

Group: Does this project require contact of CCSS study subjects for:

Additional self-reported information : **No**

Biological samples : **No**
Medical record data : **No**
If yes to any of the above, please briefly describe. :

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

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

Section: Outcomes or Correlative Factors

Late mortality : **Primary**
Second Malignancy :

Group: Health Behaviors

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

Group: Psychosocial

Insurance : **Correlative Factors**
Marriage : **Correlative Factors**
Education : **Correlative Factors**
Employment : **Correlative Factors**
Other : **Correlative Factors**
If other, please specify : **living arrangement, income**

Group: Medical Conditions

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

Group: Medications

Describe medications :

Group: Psychologic/Quality of Life

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

Group: Other

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

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 : **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**
If yes, which of the following? :
If other, please explain :

Section: Other General Comments

Other General Comments : **This is proposed as a collaboration among Pinki Prasad, Emily Tonorezos and David Freyer**