Received: 6.10.2010 First Name: Elizabeth Last Name: Wells

Institution: Children's National Medical Center

Address 1: 111 Michigan Ave, NW

Address 2: Neurology City: Washington State/Province: DC Country: USA Zip: 20010

Phone: 202-476-5973

Alternate Phone: 202-476-2120 Email: ewells@cnmc.org

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: Neurologic and Neurosensory Deficits in Long-term Survivors of Childhood Brain Tumors: Occurrence of New Deficits and Effects of Aging in Occurrence as Assessed in 2007 Survey

Planned research population (eligibility criteria): Survivors of primary central nervous systems in the original cohort.

Proposed specific aims: 1. Describe the type of neurologic and neurosensory deficits occurring in long-term survivors of childhood brain tumors and compare the incidence of these deficits to sibling controls. 2. Determine type and incidence of new deficits which have arisen since baseline assessment and the 2007 follow-up. 3. Assess factors associated with occurrence and later development of neurologic/neurosensory deficits including: tumor type, gender, age at treatment, present age, therapy (radiation therapy dose, location, etc., chemotherapy), occurrence of secondary malignancy, disease state, habits (smoking, etc.). 4. Explore possibility of utilizing genetic material available in cohort to determine molecular (genetic) parameters which may predispose to neurologic deficits (still to be determined).

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?:

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.: We stated no to biologic samples, but

may revisit this after we discuss with the chronic disease committee

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

Second Malignancy: Secondary Chronic Disease: Primary

Psychology / Neuropsychology: Secondary

Genetics:

Cancer Control:

Epidemiology / Biostatistics:

To describe the anticipated scope of the study, please indicate the specific CCSS data to be included as <u>outcome</u> (primary or secondary) or <u>correlative factors</u>. (Check all that apply)

Late mortality: Correlative Factors

Second Malignancy:

Health Behaviors

Tobacco: Correlative Factors Alcohol: Correlative Factors

Physical activity: Correlative Factors

Medical screening:

Other:

If other, please specify:

Psychosocial

Insurance:
Marriage:
Education:
Employment:

Other:

If other, please specify:

Medical conditions

Hearing/Vision/Speech: Primary

Hormonal systems: Correlative Factors Heart and vascular: Correlative Factors

Respiratory:

Digestive:
Surgical procedures:
Brain and nervous system: Primary
Other:
If other, please specify:
Medications
Describe medications: methotrexate, vincristine, cis-platin, ccnu.decadron
Pregnancy and offspring: Family History: Correlative Factors
Psychologic/Quality of Life
BSI-18:
SF-36:
CCSS-NCQ:
PTS:
PTG:
Other:
If other, please specify:
Chronic conditions (CTCAE v3): Health status:
Demographic
Age: Correlative Factors
Race: Correlative Factors
Sex: Correlative Factors
Others: Correlative Factors
If others, please specify: tumor location, radiation dose and sector
Cancer treatment
Chemotherapy: Correlative Factors
Radiation therapy: Correlative Factors
Surgery: Correlative Factors
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?

Buccal cell DNA:
Peripheral blood:
Lymphoblastoid cell lines:
Second malignancy pathology samples:
Other requiring collection of samples:
If other, please explain: this needs to be discussed

Other general comments: