First Name: Ira
Last Name: Dunkel
Institution: Memorial Sloan-Kettering Cancer Center
Address 1: 1275 York Avenue
Address 2: Box 185
City: New York
State/Province: NY
Country: USA
Zip: 10065
Phone: 212-639-2153
Alternate Phone:
Email: dunkeli@mskcc.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: Long-term outcomes in survivors of retinoblastoma (the New York cohort): a comparison of survivors versus non-cancer controls
Planned research population (eligibility criteria): CCSS SIBLINGS who were greater than 18 years of age and completed the baseline questionnaire.
Proposed specific aims: 1. To describe the long-term health status of adult survivors of childhood retinoblastoma. 2. To describe the quality of life and psychosocial functioning of adult survivors of childhood retinoblastoma. 3. To determine whether long-term survivors of retinoblastoma will have poorer general medical outcomes and/or a higher incidence of psychosocial dsyfunction compared to healthy controls.
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.:

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

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

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

Late mortality:
Second Malignancy:

Health Behaviors

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

Psychosocial

Insurance: Primary
Marriage: Primary
Education: Primary
Employment: Primary
Other:
If other, please specify:

Medical conditions

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

Medications

Describe medications:

Pregnancy and offspring: Primary
Family History:

Psychologic/Quality of Life

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

Chronic conditions (CTCAE v3): Primary
Health status: Primary

Demographic

Age: Correlative Factors
Race: Correlative Factors
Sex: Correlative Factors
Others:
If others, please specify:

Cancer treatment

Chemotherapy:
Radiation therapy:
Surgery:

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.: Yuelin Li, Memorial Sloan-Kettering Cancer Center, liy12@mskcc.org
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:

Other general comments: