

First Name: David  
Last Name: Buchbinder  
Institution: CHOC Children's Hospital / UC Irvine  
Address 1: 1201 W. La Veta Avenue  
Address 2:  
City: Orange  
State/Province: CA  
Country:  
Zip: 92868  
Phone: 714-509-8744  
Alternate Phone:  
Email: [dbuchbin@uci.edu](mailto:dbuchbin@uci.edu)

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Requirements to submit AOI:

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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

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Project Title: Psychosocial Concerns Among Siblings of Childhood Cancer Survivors

Planned research population (eligibility criteria): The proposed analyses will utilize data which was collected in 2007 via a 28-page self-report follow-up questionnaire. We will utilize data collected from healthy adult (greater than or equal to 18 years of age at time of responding to follow-up questionnaire) siblings of childhood cancer survivors (n = 2,152) and their matched survivors.

Proposed specific aims: Aim 1: To describe the psychosocial concerns of siblings of childhood cancer survivors with a focus on siblings' future health concerns, fertility concerns, cancer risk, and insurability. Aim 2: To identify the siblings' demographic factors, and health-related factors associated with psychosocial concerns (future health, fertility, cancer risk, insurability concerns) amongst siblings of survivors. Aim 3: To identify the siblings' matched survivor diagnosis, treatment-related, and health-related factors associated with psychosocial concerns (future health, fertility, cancer risk, insurability concerns) amongst siblings of survivors.

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?:

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Does this project require contact of CCSS study subjects for . . .

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Additional self-reported information: No

Biological Samples: No

Medical record data: No

If yes to any of the above, please briefly describe.:

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

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Second Malignancy:

Chronic Disease:

Psychology / Neuropsychology: Primary

Genetics:

Cancer Control:

Epidemiology / Biostatistics:

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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)

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Late mortality: Correlative Factors

Second Malignancy: Correlative Factors

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Health Behaviors

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Tobacco:

Alcohol:

Physical activity:

Medical screening:

Other:

If other, please specify:

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Psychosocial

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Insurance:

Marriage: Correlative Factors

Education: Correlative Factors

Employment: Correlative Factors

Other: Correlative Factors

If other, please specify: These will be sibling marital status, educational attainment, and employment status.

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Medical conditions

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Hearing/Vision/Speech:

Hormonal systems:

Heart and vascular:

Respiratory:

Digestive:

Surgical procedures:

Brain and nervous system:

Other: Correlative Factors

If other, please specify: We will look at survivor health status and the presence of chronic health conditions.

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Medications

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Describe medications:

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Pregnancy and offspring:

Family History:

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Psychologic/Quality of Life

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BSI-18: Correlative Factors

SF-36: Correlative Factors

CCSS-NCQ:

PTS:

PTG:

Other:

If other, please specify: We will look at the impact of survivor adverse HRQOL and psychological distress.

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Chronic conditions (CTCAE v3):

Health status:

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Demographic

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Age: Correlative Factors

Race: Correlative Factors

Sex: Correlative Factors

Others: Correlative Factors

If others, please specify: These will be sibling age, race, and sex. We would also like to look at sibling age and sex in relationship to the survivor age and sex.

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Cancer treatment

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Chemotherapy: Correlative Factors

Radiation therapy: Correlative Factors

Surgery: Correlative Factors

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Anticipated sources of statistical support

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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

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If yes, which of the following?

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Buccal cell DNA:

Peripheral blood:

Lymphoblastoid cell lines:

Second malignancy pathology samples:

Other requiring collection of samples:

If other, please explain:

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Other general comments: The outcomes of interest will be sibling concerns regarding future health, fertility, cancer risk, and insurability.