Coordinating Center Report

CCSS Investigator Meeting

May 27, 2022



Childhood Cancer Survivor Study



An NCI-funded Resource

Outline



- Provide a staffing overview
- Discuss the core activities of the CCSS Coordinating Center
- Update regarding the myLTFU participant portal
- Review priorities for the next 5 years

Overview and Staffing

Primary role: coordinate the daily activities of the study

3 main teams within the Coordinating Center:

- Project coordination, recruitment, survey administration, biologic specimen and medical record collection, coding
 - Led by Shani Alston, CCSS Project Coordinator
- Phone-based recruitment and data collection, participant tracing
 - Led by Dayton Rinehart, Call Center Coordinator
- Survey creation, database design, data cleaning, myLTFU portal development
 - Led by Chris Vukadinovich, Director of Data and Systems

CCS

Serve as the central liaison for all support facilities, committees, working groups, collaborating sites, and external investigators

Coordinate Follow-Up Surveys

- Administer questionnaires to 22,000+ active participants every 2 3 years
- Paper, online, and phone survey completion options
- Format, mail, process, code, transcribe, scan
- Reminders (lots of reminders!)
 - Email, text messages, phone calls, mailings

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Follow-Up 7 Survey Status

Participants Invited		Survivors (N = 17,552)		
	<u> </u>	% of Invited	% of Completed	
Total Completed	11,745	66.9%		
Completed Online	8,008	45.6%	68.1%	
Completed by Mail/Paper	3,028	17.3%	25.8%	
Completed by Phone	709	4.0%	6.0%	
Non-Responder	5,500	31.3%		
Refused Survey	92	0.5%		
Refused Further Contact	215	1.2%		
	Su	Survey Period: 8/19 – 12/21		

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Core Activities

External investigator-initiated studies (ancillary studies)

- Project coordination
- Recruitment
- Data collection
- Regulatory requirements

Subsequent neoplasm confirmation

Obtain pathology reports/records to histologically confirm reported conditions

Biologic specimen collection

- Oragene (saliva) kits
- Blood sample for participants with a subsequent malignant neoplasm
- Somatic tumor tissue for subsequent malignant neoplasms
- Blood sample for participants with a chronic health condition/matched controls

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Collection of DNA: Oragene

- Completed kits go to CCSS Molecular Biology Support Center in Cincinnati
- Front-end \$25 Target gift card

Kits Sent	Overall S (n = 19		Overall Siblings (n = 4,002)	
	N	% of Sent	N	% of Sent
Returned	10,409	54%	1,643	41%
Refused	1,486	7%	325	8%
Pending	5,165	27%	1,408	35%
In Tracing	1,636	9%	581	14%
Deceased	503	3%	45	1%

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Core Activities

Participant tracing

Data checks and quality control for survey data

Maintain participant and researcher websites

Participant communication and education in coordination with the CCSS Education Committee (M. Hudson, Chair)

- Newsletters 3 4 times per year
- Study results
- Respond to requests and questions
- Liaison with Participant Advisory Committee

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Sample Participant Newsletter

CCSS

///// Long-Term Follow-Up Study

Winter 2021

ITEU

A NOTE FROM US

As 2021 begins with the roll out of an effective vaccine against COVID-19, we can finally see a glimmer of light at the end of the tunnel. The pandemic has impacted our lives, work, relationships, and mental health. As a childhood cancer survivor. you may have been challenged by worries about becoming sick, difficulties in seeing health care providers, and isolation from your usual support systems.

During this stressful period, completing your LTFU survey may not feel like a pressing priority. But your survey response, amidst a global health crisis, is critically important to help us learn how the pandemic is affecting the health and wellbeing of childhood cancer survivors.

The information you provide will advance clinicians' and researchers' understanding of the pandemic's effect on all aspects of your health, including your physical, emotional, and social wellbeing.

This is a unique opportunity to "give back" to the survivorship community and help others at a time when we have all, at times, felt helpless

Please complete your survey, if you have not done so. If you didn't receive the survey or need another copy, please contact us at 1-800-775-2167 or Itfu@stjude.org.

For those of you who have already completed it, know that we are grateful. Your responses will continue to

support childhood cancer survivorship research and our eforts to keep survivors healthy.

Greg Armstrong, MD Principal investigator

RESEARCH RESULTS Pain and survivorship

Nearly one-third of LTFU Study participants reported moderate to severe pain more than 10 years after completing treatment. For many, pain significantly affects their daily life.



This is the first longitudinal study of pain a mong childhood cancer survivors. Thanks to the LTFU Study, researchers can keep moving the field forward." Cynthia Karlson, PhD, is a pediatric psychologist at University of Mississippi and a childhood cancersurvivor.

The study also identified factors that were associated with pain in daily life, · Demographic: Female sex, minority race/ethnicity, older age at diagnosis

· Cancer-related: Sarcomas/bone tumors, platinum-based chemotherapy, amputation surgery · Medical: Chronic conditions

· Psychological: Depression, anxiety, fatigue

who led the research team.

than 10 years after treatment:

• 20% of survivors reported that pain

significantly interferes with their

to severe pain

daily activities

such as:

Predictors of pain "Although some of these predictors, like a survivor's sex or cancer diagnosis, can't be changed, the study highlights lifestyle factors that are very effective for reducing and managing pain," Dr. Karlson says. "We can manage pain by decreasing stress and anxiety, and getting good sleep. When we're more stressed or not sleeping well, our pain is worse." continued on page 3

LTFU Study siblings

LTFL

///// Long-Term Follow-Up Stud

While cancer research usually focuses on the patient, a cancer diagnosis affects the entire family. This is

especially true for siblings.

Having a sibling treated for cancer during childhood is stressful. Family and school routines can change when parents must focus time and attention on the child with cancer. Parents' ability to work may also be affected, resulting in further hardships for some families. This upheaval may cause a range of emo-

tions in siblings, such as worry, sadness. guilt, and even anger. But how are siblings doing many years after this experience? Do these feelings

persist? And how do they affect siblings' health? We asked LTFU Study siblings to share their experiences.

The siblings

Jack. Jack was in the fourth grade when his sister was diagnosed with cancer in 1975. His lost my best friend." parents traveled from northeastern Massachusetts to Boston for his sister's treatment. "I remember getting bounced around a lot when my parents were at the hospital, going to neighbors or relatives," he says. "It was difficult at times. Childhood cancer is a completely different ballgame now-my sister is one of the first ones who lived." Dennis. Dennis' brother was diagnosed with cancer in 1985 at four years old; Dennis was five and the fifth of seven siblings.

Winter 2022 A NOTE FROM US

They lived in Wyoming, which lacked a

pediatric cancer treatment center, so

his mom took his brother to Denver for

there; sometimes we went, other times

our grandmother came out to support

was five, he had a stroke that left him

and I were really close, so in a way, I

with severe permanent disabilities. He

Kelly. Kelly's sister had cancer in 1977

five, and she had a relapse in 1984. Kelly

remembers her friend's mother telling

coming off the bus happy, laughing with

my best friend, happy to see my friend's

continued on page

mom, and then the crushing feeling. I

remember the exact spot in the yard

where I was told."

her that her sister was in the hospital.

"Those moments you never forget-

when Kelly was three and her sister

us," Dennis says, "Then when my brother

treatment. "She would spend months

Your health information is influencing

childhood cancer survivorship care around the world! The LTFU Study's annual Investigator Meeting was held virtually in June 2021 because of COVID-19. While I

missed seeing colleagues in person, it provided the opportunity for 600 researchers from 167 hospitals in 35 countries to attend and learn about the latest advances in survivorship care after childhood cancer.

These advances are based on data you have provided. Thank you for completing the most recent survey during the last two years.

Beginning in 2022, you will receive the next LTFU Study survey. It will be a little shorter than the previous survey and, for the first time, will ask about your experiences accessing health care, the quality of the care you receive, and the impact of the cost of your care. Look for your survey soon

This issue of the LTFU Update focuses on siblings of childhood cancer survivors. Siblings in the study share their experiences, reminding us that childhood cancer can have lasting effects on the whole family. You'll also learn about LTFU research on sibling health and find resources that support families that are affected by childhood cancer. Thank you to all of the siblings and sur-

vivors supporting the LTFU Study. Your dedication to research is helping improve health outcomes for survivors and their families

> Greg Armstrong, MD Principal investigator

> > Childhood Cancer Survivor Study

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myLTFU Participant Portal - Background

Goals:

- Implement a flexible, mobile participant portal that can remotely collect self-report and sensor-based data
- Scalable and adaptable system architecture to reliably curate large quantities of data
- Automate study processes
- Dynamic coordinator interface/dashboard for the study team
- Foster participant engagement and communication
- Establish a pool of participants available for mHealth and intervention studies

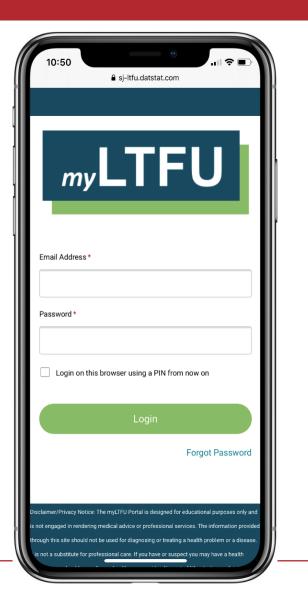
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Utilize the DatStat Connect platform to create a mobile-friendly web portal accessible by smartphone, tablet, or PC

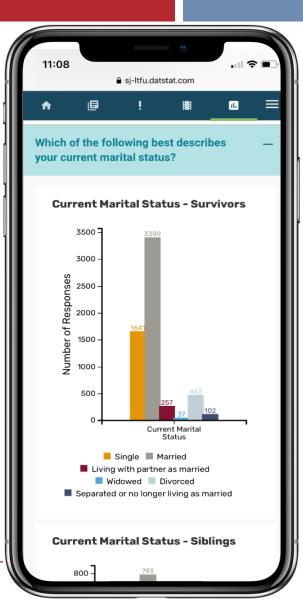
- Participants can bookmark site and save shortcut/icon to mobile device
- Administer online consents, forms, and surveys
- Upload files to the study team
- Schedule messaging (emails and texts) and data collection activities
- Provide participant resources
 - Recommended websites; newsletters; brief study results/updates; live survey results
- Coordinator portal

myLTFU Sample Screenshots





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Follow-Up Survey
COVID-19 Resources
The COVID-19 pandemic has presented new challenges and worries for all of us. Although we do not yet know how cancer survivors may be affected by this virus in comparison with the general population, please know that we are working to provide you with up-to-date resources to help you stay informed. To this end, we have created the COVID-19 Resources tab.
Communication
The mul TELL sustain will cand amails and taxts



Invited participants to the portal with the Follow-Up 7 survey

Over 13,400 participants have activated their account

Used myLTFU to rapidly collect data regarding experiences during COVID-19

Four randomized, controlled intervention trials now funded that utilize myLTFU as the primary mode for interaction with participants

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Priorities – Next 5 Years

- Administer two follow-up surveys to the cohort
- Conduct an in-home assessment of 1000 participants to examine frailty/ aging
- Administer online cognitive assessment to the cohort
- Conduct National Death Index search
- Continue subsequent neoplasm adjudication
- Continue collecting blood and tumor tissue specimens
- Coordinate recruitment and data collection for ancillary studies
- Continue participant engagement efforts while minimizing burden

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