NCI Community Oncology Research Program

EAQ211

ECOG-ACRIN cancer research group

For AYA Survivors of Hodgkin/Non-Hodgkin Lymphoma

Please Enroll Your Eligible Patients!

EAQ211 Available Through ECOG-ACRIN Cancer Research Group

Social Genomic Mechanisms of Health Disparities among Adolescent and Young Adult (AYA) Survivors of Hodgkin and Non-Hodgkin Lymphoma

Patient Population

See Section 3 for complete eligibility criteria

Selection of Sites: Enrollment is open at NCORP sites, and effective with Addendum #4, select non-NCORP sites (refer to the protocol cover page)

- *Note: when AYAs self-refer via procedures outlined in the protocol (effective with Add. #5), UM staff will sequentially assign each self-referred AYA on a rotating basis to one of 5 NCORP sites serving as entry portals for registering self-referred participants (see protocol for details)*

Selection of Participants:

- Must be ≥ 18 years of age at the time of registration
- Must have been between the ages of 15-39 at the time of their first primary cancer diagnosis of Hodgkin or Non-Hodgkin lymphoma
- Must have completed therapy (with a complete response, per clinician determination) at the time of registration
- Patient's last date of prior systemic therapy for first primary diagnosis for Hodgkin/Non-Hodgkin lymphoma must have been within 3 years prior to registration
  - ◊ Systemic therapy refers to all anti-cancer therapy, including but not limited to chemotherapy, IV/oral targeted medications, or radiation, and administered via a clinical trial or standard approach
- Must have an ECOG PS 0-3
- Must be English speaking (note: sites cannot translate the associated QOL forms)
- Must not be receiving active therapy for Hodgkin/Non-Hodgkin lymphoma
- Must have internet access through a computer, tablet, or smartphone
- Must have an email address

Methodology Plan

See Section 5 for methodology details

- Longitudinal study of 2,000 AYA lymphoma survivors recruited within 3 years following completion of treatment for Hodgkin/Non-Hodgkin lymphoma
- **Blood specimens** will be collected at baseline and subsequently every 6 months over 2 years (at 6, 12, 18, and 24 months; +/- 14 days)
- **Patient-reported outcome measures** will be administered via online/mobile survey at the same time points (at 6, 12, 18 and 24 months; +/- 14 days)
  - ◊ Information from survey data will be captured via EASEE-PRO
  - ◊ Health-Related QOL (including physical well-being) will be assessed via the PROMIS-29 v2
  - ◊ Symptoms and late effects will be assessed via the self-administered Charlson Co-morbidity index and the NCI Common Toxicity Criteria for AEs
  - ◊ Social-environmental risk factors are assessed by:
    - The Accountable Health Communities Health-Related Social Needs Screening Tool
    - The Attachment Subscale of the Social Provisions Scale
    - The Behavioral Risk Factor Surveillance System Adverse Childhood Experiences module
    - The Perceived Discrimination Scale
  - ◊ Individual resilience factors will be assessed by:
    - DUFSS
    - Mental Health Continuum Short Form
    - PROMIS Short Form
    - Indicators of socioeconomic status, assets, and employment/school status will be collected via survey

Study Chair:
Bradley J. Zebrack, PhD

Study Co-Chair:
Steven Cole, PhD

Patient Enrollment (Oncology Patient Enrollment Network [OPEN])

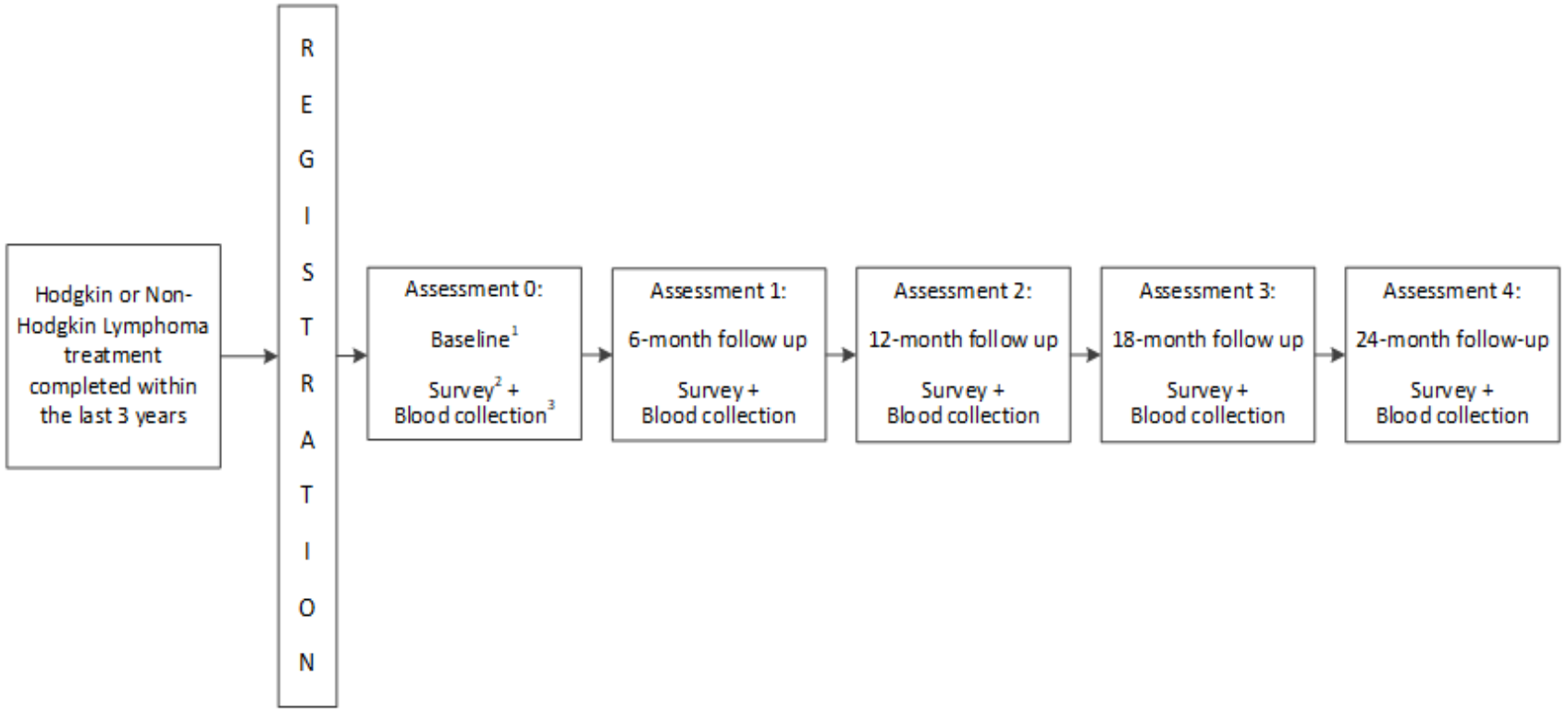
https://open.ctsu.org/open 1-888-823-5923

Protocol Information (ECOG-ACRIN Operations – Boston)

http://ecog-acrin.org (Member Login) 1-857-504-2900

EAQ211

Schema



1. All time points have a window of +/- 14 days.
2. Please see Section 5.3 for survey administration instructions for all timepoints
3. Please see Section 6 for blood collection instructions for all timepoints