NCI Community Oncology Research Program

EAQ211

ECOG-ACRIN cancer research group

For AYA Survivors of Hodgkin/Non-Hodgkin Lymphoma

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 Details

Selection of Sites: NCORP sites

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 1 year 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

- This is a longitudinal study of 2,000 AYA lymphoma survivors recruited within 1 year 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 will be 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

All Sites: Oncology Patient Enrollment Network (OPEN) https://open.ctsu.org/open

Protocol Information

ECOG-ACRIN Operations-Boston: 857-504-2900, http://ecog-acrin.org (Member Login)

Please Enroll Your Eligible Patients!

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