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ELIGIBILITY

CONCURRENT
COMPONENTS
OF VISIT

DISPOSITION

Indolent B-Cell
Lymphoma, 5 years
post treatment and
NED or
co-management
with oncologist at
any time post
treatment

SURVEILLANCE

- Annual history and physical examination with full nodal survey
- Annual CBC with differential, CMP, lipid panel, and vitamin D 25-OH
- Imaging: Consider CT² chest, abdomen, and/or pelvis (see [Appendix A](#) for recommended schedule)



Return to primary treating physician
• **Primary Oncologist** to discuss Goal Concordant Care (GCC) with patient, or if clinically indicated, with Patient Representative³

Continue survivorship monitoring

MONITORING
FOR LATE
EFFECTS

- Consider:
- Annual cardiovascular screening (see [Survivorship – Adult Cardiovascular Screening](#) algorithm)
 - Women who received radiation to the chest should receive annual breast cancer surveillance at age 25, or 8 years after radiation, whichever occurs later, with annual mammogram and breast MRI (bilateral) with contrast (see [Breast Cancer Screening algorithm](#))
 - Annual thyroid-stimulating hormone (TSH) and free T4 if prior radiation to the neck or the chest
 - Annual skin examination
- Colorectal cancer screening (see [Colorectal Cancer Screening](#) algorithm)
 - Bone health education and screening via DEXA scan starting at age 40
 - Monitor for neuropathy symptoms
 - Check immunoglobulin levels as clinically indicated
 - Annually for patients whose prior levels showed continued persistent deficiencies post treatment
 - Every 6 months for patients with a history of recurrent infections

Refer or
consult as
indicated

RISK REDUCTION/
EARLY DETECTION

PSYCHOSOCIAL
FUNCTIONING

CHRONIC HEALTH
MAINTENANCE

See [Page 2](#)

NED = no evidence of disease
CMP = complete metabolic panel
DEXA = dual-energy x-ray absorptiometry

¹ This algorithm contains the following subtypes: follicular, marginal zone, small lymphocytic, lymphoplasmacytic, nodular lymphocyte predominant (NLP) Hodgkin lymphomas, Castleman’s disease, Rosai-Dorfman Syndrome
² Consider surveillance CT scans with contrast, if not contraindicated, and CT scans without and with contrast only for the initial CT study
³ GCC should be initiated by the **Primary Oncologist**. If Primary Oncologist is unavailable, Primary Team/Attending Physician to initiate GCC discussion and notify Primary Oncologist. Patients, or if clinically indicated, the Patient Representative should be informed of therapeutic and/or palliative options. GCC discussion should be consistent, timely, and re-evaluated as clinically indicated. The Advance Care Planning (ACP) note should be used to document GCC discussion. Refer to [GCC home page](#) (for internal use only).

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RISK REDUCTION/
EARLY DETECTION

Patient education, counseling, and screening:

- Lifestyle risk assessment¹
- Cancer screening²
- Vaccinations³ as appropriate
 - Annual influenza vaccination, COVID complete vaccination and boosters, Tdap, shingles vaccine (if not already given), and other adult vaccines based on age and immune status (see [NCCN Guidelines: Survivorship: Immunizations and Infections](#))
- HPV vaccination as clinically indicated (see [HPV Vaccination algorithm](#))
- Screening for Hepatitis B and C as clinically indicated (see [Hepatitis B Virus \(HBV\) Screening and Management](#) and [Hepatitis C Virus \(HCV\) Screening](#) algorithms)

PSYCHOSOCIAL
FUNCTIONING

Assess for:

- Distress management (see [Distress Screening and Psychosocial Management algorithm](#))
- Sleep disorder and consider referral to Sleep Clinic
- Access to primary health care
- Relationship issues
- Employment status/financial issues
- Patient related outcomes

CHRONIC HEALTH
MAINTENANCE

- Confirm primary care provider (PCP) or recommend establishing care with a local PCP
- PCP responsible for assessment and management of non-cancer chronic health conditions (hyperlipidemia, diabetes, hypertension, *etc.*) or refer as indicated

Refer or consult
as indicated

¹ See [Physical Activity, Nutrition, Obesity Screening and Management](#), and [Tobacco Cessation Treatment](#) algorithms; ongoing reassessment of lifestyle risks should be a part of routine clinical practice

² Includes [breast](#), [cervical](#), [colorectal](#), [liver](#), [lung](#), [pancreatic](#), [prostate](#), and [skin](#) cancer screening

³ Based on [American Society of Clinical Oncology \(ASCO\) guidelines](#). For COVID information, see [CDC COVID vaccination guidelines](#).

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APPENDIX A: Schedule for Surveillance of Relapse with CT Scans¹

Low risk for relapse: limited stage disease, low tumor burden any stage

- Every 6 months for 2 years then yearly until year 5 **and**
- As indicated by new signs or symptoms suggesting relapse

High risk for relapse: high tumor burden meeting Group d'Etude des Lymphomes Folliculaires (GELF) criteria²

- Every 4 months for 2 years then yearly until year 10 **and**
- As indicated by new signs or symptoms suggesting relapse

¹ Consider surveillance CT scans with contrast, if not contraindicated, and CT scans without and with contrast only for the initial CT study

² High tumor burden defined as meeting one of the following GELF criteria:

- Any nodal or extranodal tumor mass > 7 cm in diameter
- Involvement of at least 3 nodal sites, each with a diameter > 3 cm
- Presence of any systemic or B symptoms
- Splenic enlargement with inferior margin below the umbilical line
- Compression syndrome (ureteral, orbital, gastrointestinal)
- Pleural or peritoneal serous effusion (irrespective of cell content)
- Leukemic phase (> 5.0 x 10⁹/L circulating malignant cells)
- Cytopenia (granulocyte count < 1.0 x 10⁹/L and/or platelets < 100 x 10⁹/L)
- Serum lactate dehydrogenase or beta2-microglobulin level above normal values

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Advance Care Planning (ACP) Conversation Workflow (ATT1925)

Continued on next page

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DEVELOPMENT CREDITS

This survivorship algorithm is based on majority expert opinion of the Lymphoma Survivorship workgroup at the University of Texas MD Anderson Cancer Center. It was developed using a multidisciplinary approach that included input from the following:

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