Follicular lymphoma (FL) is the most common indolent (slow-growing) form of B-cell non-Hodgkin lymphoma (NHL), accounting for 1 out of 5 lymphomas in the United States.

Common symptoms of FL include:

- Enlargement of the lymph nodes (bean-shaped structures that help the body fight infection, Figure 1) in the neck, underarms, abdomen, or groin
- Fatigue (extreme tiredness)

**DIAGNOSIS AND STAGING**

Typically, patients with FL have no obvious symptoms of the disease at diagnosis. Patients often only have an enlarged lymph node found upon examination by their doctor or found by chance on an imaging scan. Most patients with FL are aged 55 years or older when they are diagnosed.

To make a definite diagnosis of FL, doctors need to collect a sample of the affected lymph node. This procedure is called a biopsy. The biopsy is typically studied by a pathologist (doctor who specializes in the diagnosis of diseases by studying the cells from a patient’s body fluids and tissue samples) and preferably a hematopathologist (pathologist who has undergone additional training in the diagnosis of blood cancers, including lymphoma) who is experienced in diagnosing lymphoma. Determining the grade (level of large lymphocytes present in the affected lymph nodes) and if and how far the lymphoma has spread (staging) is important to define the best treatment for each patient.

FL is graded as 1 (low grade), 2 (mixed grade), or 3A or 3B (high grade) depending on the number of abnormal lymphocytes found on the lymph node tissue examined under the microscope. Grades 1 to 3A FL are slow growing and treated in the same way. However, grade 3B FL is usually fast growing, looks like a high-grade diffuse large B-cell lymphoma (DLBCL), and is treated the same way as DLBCL. In some patients (about 2%-3% per year), FL may transform (when a slow-growing lymphoma becomes a fast-growing lymphoma) into a more aggressive type of lymphoma, most commonly DLBCL. This transformed lymphoma usually requires more-intensive types of treatment. This change is characterized by an increase in the number of DLBCL cancer cells in the affected lymph node, which changes the follicular appearance of the cancer. For more information on transformed lymphomas, view the Transforming Lymphomas fact sheet on Lymphoma Research Foundation’s (LRF’s) website (lymphoma.org/publications).

*Figure 1. The lymphatic system (tissues and organs that produce, store, and carry white blood cells) and the lymph nodes.*
For staging, the results of the different tests [such as biopsies and scans] are used to determine the severity of the disease and the appropriate treatment. The Lugano staging system is used for FL and is depicted in Figure 1 below. This system categorizes FL from Stage I (limited disease) to IV (advanced disease), based on whether the disease is restricted to a single group of lymph nodes, has spread to other lymph nodes, or has reached the bone marrow (the spongy tissue inside the bones) and/or other organs (like the liver or lungs). Because FL is an indolent disease and might not cause any symptoms initially, it is often advanced (Stage III or IV) when it is diagnosed.

To predict the prognosis (how well the patient will do) of a patient with FL, physicians commonly use a score called the Follicular Lymphoma International Prognostic Index (FLIPI). The FLIPI score determines the risk level of each patient and predicts the chance of survival based on factors such as age and number of lymph nodes affected. Keep in mind that no two patients are alike and that statistics can only predict how a large group of patients will do, not what will happen to an individual patient. The doctor most familiar with the patient’s situation is in the best position to interpret these statistics, understand how well they apply to a patient’s particular situation, and respond to any questions you might have.

![Figure 1. Staging of FL according to the Lugano staging system. The Lugano system categorizes NHL from Stage I (limited disease) to IV (advanced disease), based on whether the cancer is restricted to a single group of lymph nodes, has spread to other lymph nodes, or has reached the bone marrow (the spongy tissue inside the bones) and/or other organs (like the liver or lungs).](image)

**TREATMENT OPTIONS**

If patients show no or very few symptoms, physicians may recommend not treating the disease right away. This approach is referred to as active surveillance (also known as “watchful waiting” or “observation”). Patients managed with active surveillance have survival outcomes similar to those treated early. In this case, patients’ overall health and disease are monitored through regular physical exams [to check for any swollen lymph nodes] or periodic imaging tests [like computed tomography (CT) scans]. If patients begin to have symptoms or signs of disease progression, treatment is initiated. There are various therapeutic options for FL based on how severe the symptoms are and how fast the cancer is growing.

FL is generally very responsive to radiation and chemotherapy. Radiation alone can provide long-lasting remission (disappearance of signs and symptoms) in some patients with early-stage disease.

Chemotherapy drugs used to treat FL include cyclophosphamide, chlorambucil, or bendamustine hydrochloride [Treanda]. Chemotherapy can be used alone or in combination with monoclonal antibodies [proteins made in the laboratory that bind to cancer cells and help the body fight cancer] that target CD20 (a protein at the surface of cancer cells). These include obinutuzumab [Gazyva], rituximab [Rituxan], and rituximab/hyaluronidase human [Rituxan Hycela, a rituximab product that is administered under the skin]. This combination is called chemoimmunotherapy and can increase the treatment response when compared to chemotherapy alone. Common chemoimmunotherapy regimens used to treat FL include:

- Bendamustine hydrochloride [Treanda] and obinutuzumab [Gazyva]
- R-Bendamustine [rituximab [Rituxan] and bendamustine]
- R-CHOP [rituximab, cyclophosphamide, Doxorubicin Hydrochloride [Hydroxydaunomycin], vincristine [Oncovin], and prednisone]
- R-CVP [rituximab, cyclophosphamide, vincristine, and prednisone]
- R-Lenalidomide [rituximab and lenalidomide [Revlimid]], often referred to as R² [R-squared]

Patients seeking information about monoclonal antibodies should view the Immunotherapy and Other Targeted Therapies fact sheet on LRF’s website [lymphoma.org/publications].

Some monoclonal antibodies, such as obinutuzumab [Gazyva] or rituximab [Rituxan], can also be used as maintenance therapy to prolong remission in patients with no signs of lymphoma after initial treatment. Patients seeking information about maintenance therapy should view the Understanding Lymphoma and Maintenance Therapy fact sheet on LRF’s website [lymphoma.org/publications].

After treatment, many patients can go into durable remission (disappearance of signs of cancer for a long period) that lasts for years; however, FL should be considered a chronic or lifelong condition. In some cases, the disease can relapse (return after treatment) or become refractory (no longer responds to treatment). For more information on relapsed and refractory FL, view the Follicular Lymphoma Relapsed/Refractory fact sheet on LRF’s website [lymphoma.org/publications].
For patients with relapsed or refractory FL, the same treatments listed on the previous page may be used, depending on the number and type of past treatments, duration of previous remission, age, health status, and patient preference. Below are other common treatments for relapsed or refractory FL:

- Targeted therapies (drugs that target specific molecules that cancer cells use to survive and spread) like copanlisib (Aliqopa) and tazemetostat (Tazverik)
- Immunotherapies (drugs that help the body’s immune system fight cancer) like Mosunetuzumab-axgb (Lunsumio) and lenalidomide (Revlimid)
- Chimeric antigen receptor (CAR) T-cell therapies (a special type of immunotherapy that uses the patient’s immune cells to fight cancer) like axicabtagene ciloleucel (Yescarta)
- Stem cell transplantation (the patient is treated with high-dose chemotherapy or radiation to remove their blood-forming cells or stem cells, then receives healthy stem cells to restore the immune system and the bone marrow’s ability to make new blood cells); for some patients with multiple relapsed FL, high-dose chemotherapy followed by stem cell transplantation may be an option

Patients seeking more information about stem cell transplantation and/or CAR T cell therapy should view the Understanding Cellular Therapy guide on LRF’s website [lymphoma.org/publications].

**TREATMENTS UNDER INVESTIGATION**

Many treatments (also referred to as investigational drugs) are currently being tested in clinical trials in patients who are previously untreated or newly diagnosed with FL. Results from these clinical trials may improve or change the current standard of care (the proper treatment that is widely used by healthcare professionals and accepted by medical experts). Table 1 below lists some of these investigational drugs that can be accessed through a clinical trial.

<table>
<thead>
<tr>
<th>Agent (Drug)</th>
<th>Class (Type of Treatment)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Atezolizumab (Tencetriq)</td>
<td>Immunotherapy, immune checkpoint inhibitor; anti-PD-1</td>
</tr>
<tr>
<td>Copanlisib (Aliqopa)</td>
<td>Targeted therapy; PI3K Inhibitor</td>
</tr>
<tr>
<td>Mosunetuzumab (Lunsumio)</td>
<td>Bispecific antibody; anti-CD20</td>
</tr>
<tr>
<td>Epcoritamab-bysp (Epkinly)</td>
<td>Bispecific antibody; anti-CD20</td>
</tr>
<tr>
<td>Venetoclax (Venclexa)</td>
<td>Targeted therapy; BCL-2 inhibitor</td>
</tr>
<tr>
<td>Pirtorubinib (Jaypirca)</td>
<td>Targeted therapy; BTK inhibitor</td>
</tr>
<tr>
<td>Toripalimab</td>
<td>Immunotherapy, immune checkpoint inhibitor; anti-PD-1</td>
</tr>
</tbody>
</table>

Abbreviations: BCL-2, B-cell lymphoma-2 protein; BTK, Bruton tyrosine kinase; CD20, cluster of differentiate 20; PD-1, programmed cell death protein 1; Phosphatidylinositol 3-kinase (PI3K)

It is important to remember that scientific research is always evolving. Treatment options may change as new treatments are discovered and current treatments are improved. Therefore, it is important that patients check with their physician or with LRF for any treatment updates that may have recently appeared. It is also very important that patients consult with a specialist to clear up any questions.

**CLINICAL TRIALS**

Clinical trials are crucial in identifying effective drugs and the best treatment doses for patients with lymphoma. Patients interested in participating in a clinical trial should view the Understanding Clinical Trials fact sheet on LRF’s website [lymphoma.org/publications] and the Clinical Trials Search Request Form at lymphoma.org, talk to their physician, or contact the LRF Helpline for an individualized clinical trial search by calling (800) 500-9976 or emailing helpline@lymphoma.org.

**FOLLOW-UP**

Since FL is generally characterized by multiple disease relapses after responses to a variety of treatments, patients should have regular visits with their physician. During these visits, medical tests (such as blood tests, CT scans, positron emission tomography [PET] scans, and biopsies of suspicious masses or bone marrow) may be required to evaluate the need for additional treatment. Some treatments can cause long-term side effects (occur during treatment and continue for months or years) or late side effects (appear only months, years, or decades after treatment has ended). These side effects can vary depending on the following factors:

- Duration of treatment (how long the treatment was given)
- Frequency of treatment (how often long the treatment was administered)
- Type of treatment given
- Patient’s age and gender
- Patient’s overall health at the time of treatment

A physician will check for these side effects during follow-up care. Visits may become less frequent the longer the patient stays in remission.

Patients and their caregivers are encouraged to keep copies of all medical records. These include test results as well as information on the types, amounts, and duration of all treatments received. Medical records are important for keeping track of any side effects resulting from treatment or potential disease recurrence. LRF’s award-winning Focus on Lymphoma mobile app [lymphoma.org/mobileapp] and Lymphoma Care Plan [lymphoma.org/publications] can help patients manage this documentation.
LYMPHOMA CARE PLAN AND PATIENT EDUCATION PROGRAMS

Keeping your information in one location can help you feel more organized and in control. This also makes it easier to find information pertaining to your care and saves valuable time. LRF’s Lymphoma Care Plan document organizes information on your health care team, treatment regimen, and follow-up care. You can also keep track of health screenings and any symptoms you experience to discuss with your health care provider during future appointments. The Lymphoma Care Plan document can be accessed by visiting lymphoma.org/publications. LRF also offers a variety of educational activities, including live meetings and webinars, for individuals looking to learn directly from lymphoma experts. To view our schedule of upcoming programs, please visit lymphoma.org/programs.

LRF FOCUS ON LYMPHOMA MOBILE APP

Focus on Lymphoma is the first app to provide patients and their caregivers with tailored content based on lymphoma subtype and with actionable tools to better manage diagnosis and treatment. The app helps with comprehensive lymphoma management, conveniently in one secure and easy-to-navigate app, no matter where you are on the care continuum. Get the right information, with resources from the entire LRF content library, use unique tracking and reminder tools, and connect with a community of specialists and patients. To learn more about this resource, visit our website at lymphoma.org/mobileapp, or contact the LRF Helpline at 800-500-9976 or helpline@lymphoma.org.

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