

2 A DOCTOR DISCUSSION GUIDE FOR PREPARING FOR NEXT STEPS

Thinking about the future when facing a cancer diagnosis can be overwhelming, and we want you to know that we are here for you every step of the way. Use this guide to help navigate discussions with your care team. Write down any additional questions or concerns you may have regarding your diagnosis or treatment plan.

WHAT IS CHRONIC LYMPHOCYTIC LEUKEMIA (CLL)?

CLL is a type of blood cancer that starts in B cells—a type of white blood cell that forms in the bone marrow.

BEFORE YOUR APPOINTMENT...

How do you feel? Nervous, hopeful—maybe both? Sharing how you feel can help you and your care team work better together.



QUESTIONS TO ASK YOUR CARE TEAM

1 | Do I need treatment? If so, what are my treatment options (including clinical trials), and what are the differences between them?

2 | Why did you select this treatment?

3 | How does this treatment work, and what's the goal of my treatment?

4 | How should I prepare for treatment before it begins?

5 | How long will I need to be on treatment?

6 | How will we know if the treatment is working?

7 | How will this treatment affect my daily life? Will I be able to go to work/do my normal activities?

8 | What types of side effects could I expect and how could I manage them? When should I see a doctor about side effects?

*Be sure to note the date and time, communicate any changes, and report all side effects to your healthcare provider right away. Call your doctor for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088.

9 | If needed, am I eligible for savings programs, and where can I find financial assistance?

Select Safety Information About CALQUENCE® (acalabrutinib) Capsules

CALQUENCE is a prescription oral treatment for adults with chronic lymphocytic leukemia or small lymphocytic lymphoma. May cause serious side effects including: serious infections, bleeding problems, decrease in blood cell count, new cancers, and heart rhythm problems. Some may lead to death. Tell your doctor if you experience infections such as flu-like symptoms; unexpected bleeding such as blood in your stool or urine; or heart rhythm problems such as fast or irregular heartbeat. Use sun protection when outside.

PLEASE SEE FULL PRESCRIBING INFORMATION, INCLUDING PATIENT INFORMATION.

ADDITIONAL NOTES

ADDITIONAL RESOURCES

To find more information and other support and resources, check out the organizations below.

American Cancer Society

cancer.org

The American Cancer Society offers programs, support, and information. Learn about treatment options, get advice on coping with side effects, and get answers to questions about health insurance, free services, and much more.

CLL Society

cllsociety.org

The CLL Society is a patient-centric, physician-curated resource for patients with CLL and their caregivers. In addition to providing helpful information, the CLL Society is a community designed to help meet the unmet needs of those with CLL.

Leukemia & Lymphoma Society (LLS)

lls.org

LLS is a source of free, highly specialized blood cancer information, education, and support for patients, survivors, families, and healthcare professionals. They have chapters across the US.

National Comprehensive Cancer Network (NCCN)

nccn.org/patients/guidelines/cll/index.html

NCCN Guidelines for Patients® are step-by-step guides to the treatment options likely to have the best results. They are based on treatment guidelines used by healthcare providers worldwide and are designed to help you discuss these options with your doctors.

Lymphoma Research Foundation (LRF)

lymphoma.org

LRF is the nation's largest non-profit organization devoted exclusively to funding innovative lymphoma research and providing support and services to people with lymphoma, their families, and caregivers.

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HAVE SPECIFIC QUESTIONS ABOUT CLL?

Check out our *Beginning Your Journey* guide.