

3 A DOCTOR DISCUSSION GUIDE FOR STAYING ON TRACK

Living with cancer can be overwhelming. We want you to remember 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 CALQUENCE?

CALQUENCE is a targeted, oral therapy for adults with chronic lymphocytic leukemia (CLL) or small lymphocytic lymphoma (SLL).

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 | How should I expect to feel from treatment? How do we know if it's working? What will we do if it doesn't work?

2 | How often should I check in with you or my other doctors once I've started treatment?

3 | If I'm experiencing a side effect, when should I contact your office?

*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.

4 | Is there anything I need to know about other medications or foods interacting with my treatment? What about my other health conditions?

5 | Is there anything I should change about my lifestyle during treatment? How do diet and exercise play a role?

6 | What will my follow-up care be like? Who will be leading this?

7 | Who can I reach out to for emotional support and resources?

8 | What should I do if I feel worse while I am taking this drug?

9 | What will I do if my cancer progresses?

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](https://www.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](https://www.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](https://www.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/ctl/index.html](https://www.nccn.org/patients/guidelines/ctl/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](https://www.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.

SOCIAL

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