

Build your everyday routine

With IMBRUVICA® By Your Side* and stay on track with your treatment

*IMBRUVICA® By Your Side patient support program is not intended to provide medical advice, replace prescribed treatment plans, or provide treatment or case management services. Patients are advised to talk to their healthcare provider and treatment team about any medical decisions and concerns they may have.

Please review the Important Side Effect Information on pages 5-7. Please see the accompanying Important Product Information or go to <u>www.imbruvica.com/prescribing-information</u>.





Use this guide to help you stay on track

It's important to make taking your medicine a natural part of your day—every day. By building a routine for taking IMBRUVICA®, you can help yourself stay on track with your treatment plan.

My personal goals

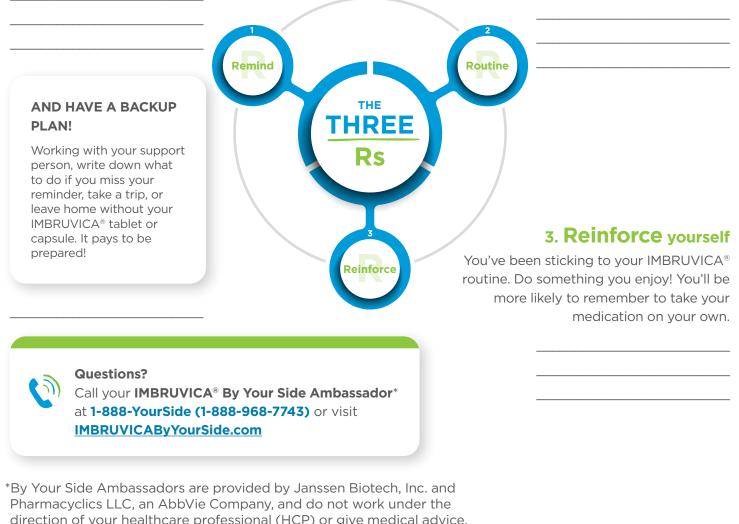
Just add "The Three Rs" to your day

1. Remind yourself

Leave yourself a reminder to take your tablet or capsule. Stick a note on your refrigerator door or ask a family member to remind you.

2. Make it Routine

IMBRUVICA® needs to be taken once daily, at about the same time each day, as directed by your doctor. Use daily activities (like lunchtime) as your reminder, until taking it becomes routine.



Pharmacyclics LLC, an AbbVie Company, and do not work under the direction of your healthcare professional (HCP) or give medical advice. They are trained to direct patients to their HCP for treatment-related advice, including further referrals.





Get your IMBRUVICA® Monthly Planner



Keep up with your IMBRUVICA® treatment by using the planner to remember key dates, such as appointments and prescription refills, and to capture your treatment experience.

Call your Ambassador at **1-888-YourSide (1-888-968-7743)** to receive your Monthly Planner.

Need more tips? Try these:



Get encouragement

Support from family, friends—even your Ambassador at **1-888-YourSide** (1-888-968-7743)—helps. My support person is _____



Take your IMBRUVICA® on the go

Living life on-the-go? You can still stick with your treatment when you leave home. Take your IMBRUVICA® tablet or capsule with you.



Questions?

Call your Ambassador at **1-888-YourSide (1-888-968-7743)** or visit **IMBRUVICAByYourSide.com**





Important information on how to take IMBRUVICA®

- Take IMBRUVICA® exactly as your healthcare provider tells you to take it.
- Take IMBRUVICA® 1 time a day at about the same time each day.

IMBRUVICA® comes as capsules, tablets, and oral suspension.

- If your healthcare provider prescribes IMBRUVICA® capsules or tablets:
- \circ Swallow IMBRUVICA $^{\scriptscriptstyle (\! R\!)}$ capsules or tablets whole with a glass of water.
- Do not open, break, or chew IMBRUVICA® capsules.
- Do not cut, crush, or chew IMBRUVICA® tablets.
- If your healthcare provider prescribes IMBRUVICA® oral suspension:
- See the detailed Instructions for Use that comes with IMBRUVICA[®] oral suspension for information about the correct way to give a dose to your child. If you have questions about how to give IMBRUVICA[®] oral suspension, talk to your healthcare provider.
- Do not use if the carton seal is broken or missing.
- If you miss a dose of IMBRUVICA[®] take it as soon as you remember on the same day. Take your next dose of IMBRUVICA[®] at your regular time on the next day. Do not take extra doses of IMBRUVICA[®] to make up for a missed dose.
- If you take too much IMBRUVICA® call your healthcare provider or go to the nearest hospital emergency room right away.

Call your doctor or pharmacist if you have any questions.

Questions? Call your Ambassador at 1-888-YourSide (1-888-968-7743) or visit IMBRUVICAByYourSide.com

Talk with your doctor and healthcare team about how you are feeling. You can report side effects to the FDA by calling **1-800-FDA-1088**.

Please review the Important Side Effect Information on pages 5-7. Please see the accompanying Important Product Information or go to <u>www.imbruvica.com/prescribing-information</u>.



IMPORTANT SIDE EFFECT INFORMATION

Before taking IMBRUVICA[®], tell your healthcare provider about all of your medical conditions, including if you:

- have had recent surgery or plan to have surgery. Your healthcare provider may stop IMBRUVICA® for any planned medical, surgical, or dental procedure.
- have bleeding problems
- have or had heart rhythm problems, smoke, or have a medical condition that increases your risk of heart disease, such as high blood pressure, high cholesterol, or diabetes
- have an infection
- have liver problems
- are pregnant or plan to become pregnant. IMBRUVICA® can harm your unborn baby. If you are able to become pregnant, your healthcare provider will do a pregnancy test before starting treatment with IMBRUVICA®. Tell your healthcare provider if you are pregnant or think you may be pregnant during treatment with IMBRUVICA®.
 - **Females** who are able to become pregnant should use effective birth control (contraception) during treatment with IMBRUVICA® and for 1 month after the last dose.
 - **Males** with female partners who are able to become pregnant should use effective birth control, such as condoms, during treatment with IMBRUVICA® and for 1 month after the last dose.
- are breastfeeding or plan to breastfeed. Do not breastfeed during treatment with IMBRUVICA® and for 1 week after the last dose.

Tell your healthcare provider about all the medicines you take, including prescription and over-the-counter medicines, vitamins, and herbal supplements. Taking IMBRUVICA[®] with certain other medicines may affect how IMBRUVICA[®] works and can cause side effects.

How should I take IMBRUVICA®?

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IMPORTANT SIDE EFFECT INFORMATION (CONT'D)

What should I avoid while taking IMBRUVICA®?

• You should not drink grapefruit juice, eat grapefruit, or eat Seville oranges (often used in marmalades) during treatment with IMBRUVICA[®]. These products may increase the amount of IMBRUVICA[®] in your blood.

What are the possible side effects of IMBRUVICA®?

IMBRUVICA® may cause serious side effects, including:

- Bleeding problems (hemorrhage) are common during treatment with IMBRUVICA[®], and can also be serious and may lead to death. Your risk of bleeding may increase if you are also taking a blood thinner medicine. Tell your healthcare provider if you have any signs of bleeding, including: blood in your stools or black stools (looks like tar), pink or brown urine, unexpected bleeding, or bleeding that is severe or that you cannot control, vomit blood or vomit looks like coffee grounds, cough up blood or blood clots, increased bruising, dizziness, weakness, confusion, change in your speech, or a headache that lasts a long time or severe headache.
- **Infections** can happen during treatment with IMBRUVICA[®]. These infections can be serious and may lead to death. Tell your healthcare provider right away if you have fever, chills, weakness, confusion, or other signs or symptoms of an infection during treatment with IMBRUVICA[®].
- **Heart problems.** Serious heart rhythm problems (ventricular arrhythmias, atrial fibrillation and atrial flutter), heart failure and death have happened in people treated with IMBRUVICA[®], especially in people who have an infection, an increased risk for heart disease, or have had heart rhythm problems in the past. Your heart function will be checked before and during treatment with IMBRUVICA[®]. Tell your healthcare provider if you get any symptoms of heart problems, such as feeling as if your heart is beating fast and irregular, lightheadedness, dizziness, shortness of breath, swelling of the feet, ankles or legs, chest discomfort, or you faint. If you develop any of these symptoms, your healthcare provider may do tests to check your heart and may change your IMBRUVICA[®] dose.
- **High blood pressure (hypertension).** New or worsening high blood pressure has happened in people treated with IMBRUVICA[®]. Your healthcare provider may start you on blood pressure medicine or change current medicines to treat your blood pressure.
- Decrease in blood cell counts. Decreased blood counts (white blood cells, platelets, and red blood cells) are common with IMBRUVICA[®], but can also be severe. Your healthcare provider should do monthly blood tests to check your blood counts.
- Second primary cancers. New cancers have happened during treatment with IMBRUVICA®, including cancers of the skin or other organs.
- **Tumor lysis syndrome (TLS).** TLS is caused by the fast breakdown of cancer cells. TLS can cause kidney failure and the need for dialysis treatment, abnormal heart rhythm, seizure, and sometimes death. Your healthcare provider may do blood tests to check you for TLS.

The most common side effects of IMBRUVICA[®] in adults with B-cell malignancies (CLL/SLL and WM) include:

- diarrhea
- tiredness

- rash
- bruising
- muscle and bone pain
- nausea



IMPORTANT SIDE EFFECT INFORMATION (CONT'D)

The most common side effects of IMBRUVICA® in adults or children 1 year of age and older with cGVHD include:

• tiredness

diarrhea

- low platelet count
- muscle and joint pain
- low red blood cell count (anemia)
 bruising
- muscle spasms

fever

- mouth sores (stomatitis)
- bleeding
- nausea
- stomach pain
- pneumonia
- headache

Diarrhea is a common side effect in people who take IMBRUVICA®. Drink plenty of fluids during treatment with IMBRUVICA® to help reduce your risk of losing too much fluid (dehydration) due to diarrhea. Tell your healthcare provider if you have diarrhea that does not go away.

These are not all the possible side effects of IMBRUVICA[®]. Call your doctor for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088.

General information about the safe and effective use of IMBRUVICA®

Medicines are sometimes prescribed for purposes other than those listed in a Patient Information leaflet. Do not use IMBRUVICA® for a condition for which it was not prescribed. Do not give IMBRUVICA® to other people, even if they have the same symptoms that you have. It may harm them. You can ask your pharmacist or healthcare provider for information about IMBRUVICA® that is written for health professionals.

USES

What is IMBRUVICA® (ibrutinib)?

IMBRUVICA® (ibrutinib) is a prescription medicine used to treat:

- Adults with chronic lymphocytic leukemia (CLL)/Small lymphocytic lymphoma (SLL).
- Adults with chronic lymphocytic leukemia (CLL)/Small lymphocytic lymphoma (SLL) with 17p deletion.
- Adults with Waldenström's macroglobulinemia (WM).
- Adults and children 1 year of age and older with chronic graft versus host disease (cGVHD) after failure of 1 or more lines of systemic therapy.

It is not known if IMBRUVICA® is safe and effective in children under 1 year of age.

Please see the accompanying Important Product Information or go to <u>www.imbruvica.com/prescribing-information</u>.





imbruviča® By Your Side Patient Support

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For more information about PCYC's privacy practices or how to opt-out, visit https://www.pharmacyclics.com/privacy-policy.



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