

Section 3: Terms of Participation and Program Disclosures

PATIENT ENROLLMENT SECTION Jaypirca[®] (pirtobrutinib)

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SUBMIT COMPLETED PAGES 1-5 VIA FAX AT 1-855-511-2515.

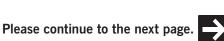
For assistance, call 1-800-LillyRx (1-800-545-5979), Monday-Friday, 8am-10pm ET.

THIS PAGE MUST BE SUBMITTED

	Patient Name (First, MI, Last)		Date of Birth (MM/DD/YYYY)		
ion	Address	City	State	Zip	
Patient Information	US or Puerto Rico Resident Yes No	Gender M F P	referred Language English Spa	anish Other	
ient In	Phone* (000-000-0000)	Email			
Pat	*By checking the box, I agree to receive automated marketing calls and texts from and on behalf of Eli Lilly and Company. I understand that I am not required to provide my number as a condition of receiving goods and services. Message and data rates may apply. By checking the box, I agree to be contacted to: provide feedback on my experience with the related products, services, and programs; to share my story; and, to participate in market and medical research studies about products and services.				
ccuon 2. nce Information	Must select one of the following: No Insurance Coverage Copy of Policyholder's Insurance Card (Front and Back) Is Attached Provide Information Below				
	Primary Prescription Insurance Company				
	Insurance Company Phone (000-000-0000) _		Cardholder Name		
Insurance	Policy/ID		Group #		
	RX BIN		PCN		

TERMS OF PARTICIPATION AND PROGRAM DISCLOSURES:

Your healthcare provider has talked with you about using Jaypirca®, an Eli Lilly and Company medicine. Lilly Support Services[™] for Jaypirca[®] offers personalized support to Patients at no charge and was created to help you have a positive experience as you get started with and use this medicine. By signing and submitting this form, you consent to your enrollment into Lilly Support Services[™] for Jaypirca[®]. As part of your participation in Lilly Support Services[™] for Jaypirca®, you understand and authorize Lilly USA, LLC to retain and use your personal information for the purposes described in this form. Eli Lilly and Company, Lilly USA, LLC and its affiliates, agents, representatives, and service providers (together "Lilly") may use, disclose, and/or transfer the personal information you supply to provide services related to your condition and treatment to administer the program. The Lilly Support Services™ for Jaypirca® Support team can contact you by email, mail or telephone to provide personalized services and information and materials directly related to your condition and therapy; responding to customer service requests and/or questions about your treatment; disclosing your enrollments and use of these services to your doctors and insurers; analyzing and/or measuring program performance and program effectiveness for future enhancements; and other activities related to your condition and therapy that are part of Lilly Support Services[™] for Jaypirca[®]. Your personal information, including information that may be related to your health, is needed to fulfill your request. To cancel your participation in the program, please contact us at 1-800-LillyRx (1-800-545-5979) Monday-Friday, 8am-10pm ET. For information about Lilly's privacy practices, please see our Privacy Statement at https://privacynotice.lilly.com and the Consumer Health Privacy Notice at https://www.lillyhub.com/legal/lillyusa/CHPN.html.







PRESCRIBER ENROLLMENT SECTION Jaypirca® (pirtobrutinib)

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	Prescription
	9
Section of	ICP Service Selection & Prescription

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Prescriber I	Prescriber Name (First, Last)		NPI #		
Practice Name Office Phone (000-000-0000)					
Office Fax (000-000-0000)	Office Address			
Practice Na Office Fax (Office State 0	ffice Zip G	Group Tax ID	
Office Conta	act Name	Office	Contact Phone (000-000-000	00)	
Office Conta	Office Contact Email Secondary Office Contact				
		Patient Addre			
Patient City Diagnosis:		Patient State	Patient Zip .		
Adult na	tients with chronic lymphocytic leuk	emia or small lymphocytic lymphoma	(CLL/SLL) who have received a	at least two prior lin	es of therapy,
I — `	g a BTK inhibitor and a BCL-2 inhibitor	or /R) mantle cell lymphoma (MCL) after a	at least two lines of systemic th	perany including a	RTK inhibitor
		ompendia and not unsupported in any	-	icrapy, including a i	DTR IIIIIDIOI
Jaypirc Jaypirc Program Patients in insura determin the Patie 15-day si is only av or other-	Program™ does not guarantee coverage e program at any time. Additional Terms KED, MUST READ ELIGIBILITY REQUI	ON SECTION BELOW. Fort—The Jaypirca Interim Access at no charge for eligible, insured ng a minimum 5-business-day delay hose insurers have made a final nere is a persistent coverage delay, by be eligible for up to three additional Jaypirca Interim Access Program™ laity Pharmacy. No purchase contingency action. The Jaypirca Interim ge. Lilly reserves the right to change and Conditions apply - see page 5.	Specialty Pharmacy Phone N on below and sign at the	·	
	Dosing		Quantity to be Dispensed	<u> </u>	Refills
You must select the appropriate Dosing	Recommended Dosing: 200 mg (2 x 100mg tablets) orally of 300mg (3 x 100mg tablets) orally or 150mg (3 x 50mg tablets) orally or 100mg (2 x 50mg tablets) orally or 50mg cyclly once doily	once daily nce daily	60 Tablets (30 day supply) 90 Tablets (30 day supply) 90 Tablets (30 day supply) 60 Tablets (30 day supply) 30 Tablets (30 day supply)		Refills (1-11)
USA, LLC, their has directed my made through ti identified in this only to the dispe	affiliates, agents, representatives, business predisclosure of their information to Lilly so that he duration of the Patient's therapy; 4) I will not form, the prescription complies with my state ensing pharmacy. I understand that by signing	ary and that this information is accurate to the be artners, and service providers (together "Lilly") to Lilly may contact the Patient to further enable se to seek reimbursement from any third party for the specific prescribing requirements and I appoint I; this form, I am requesting support from Eli Lilly a pendia and the use is not listed as unsupported, r	Ist of my knowledge; 2) I am disclosing it help enable treatment for this Patient; rvices for those purposes and that such e support Lilly provides; and 5) I am lice Lilly as my agent for the limited purpose and Company for Patients receiving Jay	3) The Patient is aware of consent and direction a ensed to prescribe the press of conveying this pres- pirca® pursuant to an FD	of, has consented to, and pplies to disclosures rescription medication cription by facsimile A approved indication
D	ispense as written	May substitute/brand	l exchange permitted		(MM/DD/YYYY)







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You have selected Eli Lilly and Company ("Lilly") to coordinate certain services related to your health and to provide information related to your health (Lilly's "Programs and Services"). In order for Lilly to offer the Programs and Services, Lilly may need to obtain or exchange your protected health information ("PHI") as defined under the Health Insurance Portability and Accountability Act of 1996, as amended ("HIPAA") from your Health Care Entities (as defined below). PHI can be inclusive of "sensitive data" as defined by applicable U.S. privacy laws. After your PHI has been released to Lilly, it is no longer covered by HIPAA. By signing this form, you understand and authorize your Health Care Entities to share your PHI with Lilly and use as explained below.

PHI includes the following individually identifiable information:

- Information about your health insurance or benefits, including how much coverage you have
- All relevant records about your treatment, including medication histories and prescriptions
- Information about your payment for treatment, including any insurance coverage
- Whether you're staying on your medicine or treatment

If you agree, your PHI may be collected from and shared by these entities (together "Health Care Entities"):

- Your doctors and other healthcare providers
- Your healthcare plan or health insurance company
- Clearinghouses or other agents
- Your pharmacy
- Others who might have your PHI on behalf of your healthcare providers, pharmacies and healthcare plans

How Your PHI Will Be Used

Your PHI will be used to enroll you in, provide you with, and operate and administer the Programs and Services, consistent with Lilly's Privacy Statement and Consumer Health Privacy Notice, including to:

- understand how much of your Lilly treatment is covered by your insurance
- help you find ways to afford such treatment
- track the shipment, receipt, and use of your Lilly treatment and Programs and Services
- share information with your Health Care Entities and communicate with them regarding Lilly Programs and Services
- contact you about Lilly Programs and Services related to your health needs
- measure Lilly Programs and Services' performance in order to make improvements and drive business decisions and metrics
- de-identify your data for analytics including reports about Health Care Entities' use of Lilly Programs and Services.



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Other things you should know about how we may use and share your PHI:

We do not ask for any PHI that we do not need, but we may receive some in the health records sent to us. Your PHI will be released to Lilly and its wholly owned subsidiaries ("Lilly" or "we") and/or entities or persons that work on behalf of, or in partnership with, Lilly but are not Lilly employees ("Third Parties").

- You don't have to give permission to share your PHI with Lilly to receive treatment from your Health Care Entities, your prescription from your pharmacy, or benefits from your healthcare plan, but Lilly Programs and Services may not be able to help you without your Authorization.
- Your Health Care Entities may receive compensation from us in exchange for sharing your PHI. They may also be paid by us to use your PHI to provide services, such as contacting you about Lilly products.
- Your signed authorization to share and use your PHI lasts for the duration of your participation in Lilly Programs and Services from the date of your signature or earlier as required by state law. In any case, you may revoke this Authorization for Lilly Programs and Services and you may request to obtain PHI from your Health Care Entities at any time by writing to PO Box 501847, San Diego, CA 92150. Your revocation of this Authorization will not have any effect on any uses or disclosures of your PHI that occurred prior to Lilly's receipt of your revocation.
- Your revocation of this Authorization will be effective when your Health Care Entities receive notice of your cancellation or revocation and will not apply to any information shared with Lilly prior to receipt of the notice.

AUTHORIZATION TO USE AND DISCLOSE PROTECTED HEALTH INFORMATION: I authorize my Health Care Entities to disclose my PHI and sensitive data for the purposes as described in this HIPAA Authorization. This HIPAA Authorization replaces any prior HIPAA Authorizations that I may have provided at a specific program level.

By signing this form, I attest that I have read and agree to the Patient HIPAA Authorization. I understand I am entitled to a copy of this signed Authorization.



Signature of Patient				
Printed Name of Patient				
Signature Date (MM/DD/YYYY)				
DOB (MM/DD/YYYY)				







PRESCRIBER ENROLLMENT SECTION Jaypirca® (pirtobrutinib)

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Jaypirca Interim Access Program™ Requirements

To be eligible for the Jaypirca Interim Access Program™, a patient must be: 1) prescribed Jaypirca® for the first time and have insurance (commercial or governmental); 2) experiencing a minimum 5-business day delay in insurance coverage determination; 3) prescribed Jaypirca® for an FDA-approved indication or an indication medically supported by CMS-recognized compendia; 4) enrolled in Lilly Support Services; and 5) a resident of the United States or Puerto Rico.

Uninsured Patients are not eligible for the Interim Access Program™.

Please note: An insurer may require that the Healthcare provider submit a Prior Authorization in order to obtain coverage. This Program is provided by Sonexus Non-Commercial Specialty Pharmacy (NCSP) rather than your in-office dispensary or any other Specialty Pharmacy your patient may later use. We will contact your office when the patient has received his/her dose. Sonexus NCSP supplies and executes a 15-day supply of Jaypirca® at no charge for eligible, insured patients.

Terms and Conditions:

The Jaypirca Interim Access Program (or "Program") provides a 15-day supply of Jaypirca at no charge for eligible, insured patients who are:

- 1) prescribed Jaypirca for the first time,
- 2) experiencing a minimum 5-business-day delay in insurance coverage determination,
- 3) prescribed Jaypirca for an FDA-approved indication or an indication medically supported by CMS-recognized compendia,
- 4) enrolled in Lilly Support Services, and
- residents of the United States or Puerto Rico.

May not be combined with any other offer. Not available to patients whose insurers have made a final determination to deny the patient coverage for Jaypirca. If a denial is received after the initial 5 business days have passed and appeal rights are being pursued, or if there is a persistent coverage delay, the patient, under appropriate circumstances, may be eligible for up to 3 additional 15-day supplies of Jaypirca. Product provided through the Program is only available through the Program non-commercial specialty pharmacy. Product is provided free of charge and may not be sold, bartered, or returned for credit. Reimbursement cannot be sought from any third party for product provided under the program. Patients enrolled in Medicare Part D are prohibited from counting any portion of the cost of the product provided under the Program towards true out-of-pocket ("TrOOP") costs for prescription drug calculations. No purchase contingency or other obligation accompanies program participation. This Program is not health insurance and does not guarantee coverage. Lilly reserves the right to change or end the program at any time. Benefits under the program are not transferable.

→	I certify and agree to the above Jaypirca Interim Access Signature of Patient Not signing this form will result in an incomplete submission ar	Date of Signature (MM/DD/YYYY)					
Patient's Name (First, MI, Last)							
Patient's DOB (MM/DD/YYYY) Patient's Address							
Patie	ent's City	Patient's State Patient's	Zip				
\(\)	Jaypirca® Prescription - Fill out corresponding prescription below and sign at the bottom of page						
	Primary Diagnosis:	Dosing:	Quantity to be Dispensed				
You must	Adult patients with chronic lymphocytic leukemia or small lymphocytic lymphoma (CLL/SLL) who have received at least two prior lines of therapy, including a BTK inhibitor and a BCL-2 inhibitor	Recommended Dosing: 200 mg (2 x 100mg tablets) orally once daily 300mg (3 x 100mg tablets) orally once daily	30 Tablets (15 day supply) 45 Tablets (15 day supply)				
select a Diagnosis and Dosing	Adult patients with relapsed or refractory (R/R) mantle cell lymphoma (MCL) after at least two lines of systemic therapy, including a BTK inhibitor	150mg (3 x 50mg tablets) orally once daily	45 Tablets (15 day supply) 30 Tablets (15 day supply)				
	Diagnosis supported by CMS recognized compendia and not unsupported in any CMS approved compendia	50mg orally once daily	15 Tablets (15 day supply)				
Prescriber: I certify that I understand and agree: 1) To the terms and conditions of the Jaypirca Interim Access Program™; 2) I am licensed to prescribe the medication identified in this form, and that the prescription complies with my state-specific prescribing requirements; 3) In my medical judgment, Jaypirca® is clinically appropriate for the Patient named above and its use is consistent with the FDA-approved indication or an indication medically supported by CMS recognized compendia and not unsupported in any CMS approved compendia; 4) This supply of Jaypirca® is specifically for the Patient named above; and 5) I will not seek reimbursement from any third party for the support Lilly provides. PRESCRIBER SIGNATURE: PRESCRIBER MUST MANUALLY SIGN AND DATE. Rubber stamps, signature by other office personnel for the Prescriber, and computer-generated signatures will not be accepted.							
→	Dispense as written Not signing this form will result in an incomplete submission and a	0 .	ned Date (MM/DD/YYYY)				





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Privacy Notice:

This Privacy Notice ("Notice") is intended to supplement the Eli Lilly and Company Privacy Statement (https://privacynotice.lilly.com) and the Consumer Health Privacy Notice (https://www.lillyhub.com/legal/lillyusa/CHPN.html) that can be accessed in the footers of Lilly's websites. This Notice is to provide you with information about the personal information, including health information, we may collect, use, disclose or otherwise process, and your rights and choices with respect to your information.

The categories of health information we collect will depend on how you interact with Lilly Services and the information you choose to provide. We may collect:

- Health conditions, treatments, diseases, or diagnosis
- Social, psychological, behavioral, and medical interventions
- Health-related surgeries or procedures
- Use or purchase of prescribed medication
- Bodily functions, vital signs, symptoms, or measurements of other types of consumer health data
- Diagnoses or diagnostic testing, treatment, or medication

- Reproductive or sexual health information
- Biometric data
- Genetic data
- Data that identifies a consumer seeking health care services
- Other information that may be used to infer or derive data related to the above or other health information.

With your consent, we may use the health information we collect for the following purposes, as further described in our privacy statements:

- Providing Services and support.
- Analytics and improvement.
- Customization and personalization.
- Marketing and advertising.

- Security and protection of rights.
- · Legal proceedings and obligations.
- General business and operational support.

Lilly does not sell or share your health information with third parties without your consent or authorization. We may disclose health information to our processors for our business purposes or at your direction to provide you with products and Services that you request.

We may use and save your personal information to meet legal or regulatory obligations that are in the legitimate interest of Lilly, to fulfill legitimate and lawful business purposes in accordance with Lilly's record retention policies and applicable laws and regulations, and to respond to lawful requests by public authorities, including to comply with national security or law enforcement requests.

Some of this personal information may be considered sensitive under applicable laws, such as information about your health or medical diagnosis and demographic information collected in some circumstances, such as race, ethnic origin, and sexual orientation. We may process your sensitive PI with your consent, or as otherwise permitted by law.

Upon verification, you have rights with respect to the collection, use and storage of your information. These rights may include access to your information and how it is being used or shared, the right to correct, delete or limit use of your information or to withdraw consent for us to collect and use your information. There may be certain exceptions and limitations that apply to your request including the right to have your information transmitted to another entity or person in a machine-readable format. To exercise your rights, you or your authorized representative may submit a request to datarights@lilly.com or 1-800-Lilly-Rx (1-800-545-5979). You will not be discriminated against for exercising any of your rights. You may be entitled, in accordance with applicable law, to appeal a refusal to take action on your request. To do so, please contact us by using one of the methods listed here or in How to Contact Us section of the online Privacy Statement.

If you wish to raise a complaint on how we have handled your personal information, you can contact the Global Privacy Office and Data Protection Officer at privacy@lilly.com, who will investigate the matter. If you are not satisfied with our response or have any concerns about how your data is being processed, you can register a complaint with a relevant regulatory authority (e.g., a Data Protection Authority (DPA) or Attorney General).

