Commercial Enrollment Form



PATIENT INFORMATION Please provide the following information about your patient. All fields are required unless otherwise noted. Patient first name Patient middle initial Patient last name Date of birth (MM/DD/YYYY) Male Female Nonbinary Last four digits of social security number Age Mailing address (for medication delivery) Street address ZIP City State Mobile phone number (preferred) Home phone number (optional) **Email address** Name of caregiver (optional) Relation of caregiver Primary mobile phone number of caregiver Email address of caregiver PATIENT DIAGNOSIS Please select the diagnosis code you have given your patient with PK deficiency. D55.21: Anemia due to pyruvate kinase deficiency D55.29: Anemia due to other disorders of glycolytic enzymes D55.8: Other anemias due to enzyme deficiency/disorders D55.9: Anemia due to enzyme disorder, unspecified D58.9: Hereditary hemolytic anemia, unspecified If your patient does not have a diagnosis code, please contact your Patient Support Manager (PSM). Your PSM will contact you to convey additional information. INSURANCE INFORMATION Please check and provide information about your patient's insurance(s). All fields are required unless otherwise noted. Be sure to include a copy of both sides of your patient's insurance card(s). Patient insurance (check all that apply): No insurance Medicare Medicaid Commercial/private Other Primary health insurance Prescription drug insurance Secondary insurance Plan name Plan name Plan name

Phone #_ Phone #_ Phone #_ Policy ID # Policy ID # Policy ID # Group # Policy holder (if other than patient) Policy holder (if other than patient) RX BIN # Name Name PCN# Date of birth Date of birth (MM/DD/YYYY) (MM/DD/YYYY)



4 CHECK BELOW FOR PYRUKYND® (MITAPIVAT) TABLETS PRESCRIPTION

Please be sure to fill in all the information below, including the number of refills. All fields are required unless otherwise noted.

Patient name		Date of birth (MM/DD/YYYY)			
Rx start date (MM/DD/YYYY)					
PYRUKYND Dosage Instructions: SIG:				Supplied: ment Pack (28-Day)	
Quantity: Refil	ls:		• 20 mg BID Trea	tment Pack (28-Day) tment Pack (28-Day)	
Please list any specific instructions you wou	ıld like to provide to your p	patient:	• 30 mg bib nea	itment rack (20-Day)	
Dispense as written (DAW)					
Physician Signature Dispense as written).			Date		
No stamps. NY providers must submit a vali	d NY prescription or eScril	be to dispensing pharmacy	:		
PRESCRIBER & PRACTIC	E CONTACT IN	NFORMATION A	AND DECLAR	ATION	
Please provide the following infor All fields are required unless other	mation about you ar	nd your practice. Prescriber specialty Hematologist/Onco			
Prescriber name		Hematologist	Other		
Practice name		Practice staff contact	name		
Street address		City	State	ZIP	
Practice staff contact phone number	Fax number	F	Practice staff contact	email address	
NPI number Medicare/	Medicaid provider num	ber Prov	ider Transaction Acce	ss Number (PTAN)	
Tax ID	State license num	ber			
Your patient has previously taken this	medication (eg, in the cli	nical trials ACTIVATE or AC	CTIVATE-T)		
Trial ID					
Trial ID I certify that the patient and physician inform have prescribed PYRUKYND based on my ju patient's treatment. I authorize, if appropriat my patient. I understand that neither I nor n free product received under the program. I other information as may be required by Ag initiating or continuing PYRUKYND therapy,	dgment of medical neces te, the forwarding of this p ny patient may seek reimb certify that I have obtained ios or its agents to assist r	sity, and in accordance wit prescription to an authorize ursement from any govern d my patient's authorization my patient in obtaining cov	h its labeled indication led specialty pharmacy or ment program or third- n to release the above in	will be supervising my n behalf of myself and party insurer for any formation and such	
Physician Signature	,	- 7 (D	ate	



6 PATIENT AUTHORIZATION TO USE/DISCLOSE HEALTH INFORMATION (REQUIRED)

Please have your patient agree to the terms below. All fields are required unless otherwise noted.

I understand that myAgios Patient Support Services is a service offered by Agios Pharmaceuticals, Inc. to help eligible patients who have been prescribed PYRUKYND® (mitapivat) tablets obtain insurance coverage and financial assistance for PYRUKYND, including through its Coverage Interruption and Patient Assistance Programs (the "Programs"). I give permission for my physician and their staff to disclose my health and other personal information, including, but not limited to the information on this form, to Agios Pharmaceuticals, Inc. and its agents and representatives (collectively "Agios") so that Agios may use and further disclose my information to healthcare providers, pharmacies, insurance companies, prescription drug plans, and other third-party payers and patient assistance groups (collectively, "Third Parties") in order to: (1) enroll me in the Programs; (2) facilitate the filling of my prescription for and the delivery and administration of PYRUKYND; (3) assist me in obtaining insurance coverage for PYRUKYND; (4) contact me about PYRUKYND and the Programs (this may include supplemental educational materials, information, offers, and services related to my therapy or my medical condition, or opportunities to participate in focus groups, surveys, or interviews); and (5) manage the Programs. I further authorize the Third Parties to disclose health and other personal information about me in their possession to Agios in order to assist Agios in accomplishing the purposes described above. I understand that once my information is disclosed pursuant to this authorization, it may no longer be protected by federal privacy laws (the Health Insurance Portability and Accountability Act) or state privacy laws and may be further disclosed to others. However, I understand that Agios will not release my information to any party, except as provided in this authorization or as permitted by applicable law, without first obtaining my (or my authorized representative's) separate written consent. I understand that I may refuse to sign this authorization and such refusal will not affect my ability to receive PYRUKYND that is paid for by my insurer, my treatment, payment for treatment, eligibility for or enrollment in health benefits, but it will limit my ability to receive support services for PYRUKYND, including participation in free medication programs. I understand that this authorization will remain in effect for 3 years, or a shorter period as may be required by state law, from the date of my signature, unless I revoke it earlier by contacting Agios in writing at AllCare c/o Agios Patient Support Services, 50 Bearfoot Rd, Northborough, MA, 01532. If I revoke this authorization, Agios and any Third Parties who are notified of my revocation will stop using and disclosing my information as soon as possible, but the revocation will not affect prior use or disclosure of my information in reliance on this authorization. I understand that the services described in this authorization may be reduced at any time, without prior notification. However, if any services are added, Agios will obtain my authorization to receive any such additional services. I understand that certain Third Parties may receive compensation in exchange for their disclosure of my information to Agios. I also understand that I have the right to receive a copy of this authorization. I verify the information provided is true and correct. If I am the caregiver/representative for the patient, I confirm I am authorized to sign on behalf of the patient.

Patient name	Caregiver/Guardian name	
	-	
Patient Signature		Date
Caregiver/Guardian Signature		Date



7 CONSENT TO RECEIVE MATERIAL FROM AGIOS (OPTIONAL)*

By enrolling in myAgios Patient Support Services, you may receive support and educational materials on pyruvate kinase deficiency and PYRUKYND® (mitapivat) tablets.

By clicking "I Agree," I consent that the information I am providing may be used by Agios and its agents to keep me informed about products, patient support services, or other opportunities that may be of interest to me via mail or email. Agios may also combine the information I provide with information about me from third parties to better match information with my interests. You may receive mail or emails that contain information to support and educate on pyruvate kinase deficiency as well as those that market or advertise Agios products or services. I understand from time to time, Agios' Online Privacy Policy may change and for the most recent version of the Online Privacy Policy, I should visit myAgios.com/privacypolicy.

Agios understands protecting your personally identifiable information is very important. We do not share any personally identifiable information you give us with third parties for their own marketing use.

Please see Important Safety Information below and accompanying Full Prescribing Information.

O I agree

Please have your patient agree to each of the following statements:

- I understand that I will receive educational information and updates about PK deficiency, research opportunities, and other information that may be of interest to me from Agios.
- O I consent to receive phone calls and text messages from and on behalf of Agios [and its affiliates] that contain information to support and educate on pyruvate kinase deficiency as well as those that market or advertise Agios products or services at the phone number(s) I provide. I understand that my consent is not required or a condition of purchasing goods or services from Agios.

8 SUBMIT FORM

Just one more step to go. To complete the enrollment process, please be sure to download your completed form, print it, and then fax it to myAgios Patient Support Services at 1-800-951-7814.

INDICATION

PYRUKYND is a pyruvate kinase activator indicated for the treatment of hemolytic anemia in adults with pyruvate kinase (PK) deficiency.

IMPORTANT SAFETY INFORMATION

Acute Hemolysis: Acute hemolysis with subsequent anemia has been observed following abrupt interruption or discontinuation of PYRUKYND in a dose-ranging study. Avoid abruptly discontinuing PYRUKYND. Gradually taper the dose of PYRUKYND to discontinue treatment if possible. When discontinuing treatment, monitor patients for signs of acute hemolysis and anemia including jaundice, scleral icterus, dark urine, dizziness, confusion, fatigue, or shortness of breath.

Adverse Reactions: Serious adverse reactions occurred in 10% of patients receiving PYRUKYND in the ACTIVATE trial, including atrial fibrillation, gastroenteritis, rib fracture, and musculoskeletal pain, each of which occurred in 1 patient. In the ACTIVATE trial, the most common adverse reactions including laboratory abnormalities (\geq 10%) in patients with PK deficiency were estrone decreased (males), increased urate, back pain, estradiol decreased (males), and arthralgia.

Drug Interactions:

- Strong CYP3A Inhibitors and Inducers: Avoid concomitant use
- Moderate CYP3A Inhibitors: Do not titrate PYRUKYND beyond 20 mg twice daily
- · Moderate CYP3A Inducers: Consider alternatives that are not moderate inducers. If there are no alternatives, adjust PYRUKYND dosage
- Sensitive CYP3A, CYP2B6, CYP2C Substrates Including Hormonal Contraceptives: Avoid concomitant use with substrates that have narrow therapeutic index
- UGT1A1 Substrates: Avoid concomitant use with substrates that have narrow therapeutic index
- P-gp Substrates: Avoid concomitant use with substrates that have narrow therapeutic index

Hepatic Impairment: Avoid use of PYRUKYND in patients with moderate and severe hepatic impairment.

 $\hbox{Please see $\underline{$full$ Prescribing Information}$ for PYRUKYND.}$

^{*}Patient is under no obligation to complete Section 7 to receive their prescription or to enroll in the Patient Support Program.