

Arkansas Medicaid Prescription Drug Program Oncology Medication Prior Authorization Fax Form

Fax completed form and required documentation to Arkansas Medicaid Pharmacy Program

Fax: 800-424-5851 For questions, call: 501-683-4120.

This prior authorization request form pertains to pharmacy processed oncology medications. Oncology medications obtained through medical billing should not be requested with this form.

Prime Therapeutics partners with CoverMyMeds to allow for the submission of electronic PA requests. **For faster coverage determinations**, go to www.coverMyMeds.com

If the following information is not complete, correct, or legible, the prior authorization (PA) process can be delayed. Please use one form per beneficiary.

Requestor Name:	Title:
BENEFICIARY INFORMATION	
Medicaid ID:	Date of Birth:
Beneficiary Last Name:	
Beneficiary First Name:	
PRESCRIBER INFORMATION	
Prescriber Last Name:	
Prescriber First Name:	
Prescriber NPI:	DEA Number:
Prescriber Phone:	Prescriber Fax:
DIAGNOSIS AND TREATMENT HISTORY	
Diagnosis:	
ICD-10 Code:	
☐ New Therapy ☐ Renewal	
If renewal, duration of therapy (specific dates):	to
DRUG INFORMATION	
Drug Name:	
	Dosage Form:
Directions:	

Beneficiary Name (Last, First):					
DRUG INFORMA	ATION (CONTINUE	D)			
Me	edications Covered	as a Pharmacy Claim (s	select requested medicat	tion(s))	
Abiraterone	Afinitor tablets	Afinitor Disp (PPB)	Akeega	Alecensa	
Alunbrig	Anastrazole*	Arimidex*	Augtyro	Avmapki/ Fakzynja	
Ayvakit	Balversa	BESREMI	Bosulif	☐ Braftovi	
Brukinsa	☐ Cabometyx	Calquence	Caprelsa	Cometriq	
Copiktra	Cotellic	Danziten	Dasatinib	☐ Daurismo	
Erivedge	Erleada	Erlotinib	☐ Everolimus	Exkivity	
Femara*	Fotivda	Fruzaqla	Gavreto	Gefitinib	
Gilotrif	Gomekli*	Hernexeos	☐ Ibrance	☐ Ibtrozi	
Iclusig	☐ Idhifa	Imbruvica	☐ Imkeldi	☐ Inlyta	
Inqovi	☐ Inrebic	☐ Iressa	☐ Itovebi	☐ Iwilfin	
Jakafi	Jaypirca	Kisqali	Kisqali/Femara	☐ Koselugo*	
☐ Krazati	Lapatinib	Lazcluze	Lenalidomide	Lenvima	
Letrozole*	Lonsurf	Lorbrena	Lumakras	Lynparza	
Lytgobi	Mekinist	Mektovi	Mercaptopurine Susp	Modeyso	
Nerlynx	Nexavar (PPB)	Ninlaro	☐ Nubeqa	Odomzo	
Ogsiveo	Ojemda	Ojjaara	Onureg	Orgovyx	
Orserdu	Pazopanib	Pemazyre	Phyrago	Piqray	
Pomalyst	Purixan	Qinlock	Retevmo	Revlimid (PPB)	
Revuforj	Rezlidhia	Rezurock*	Romvimza	Rozlytrek	
Rubraca	Rydapt	Scemblix	Soltamox	Sprycel (PPB)	
Stivarga	Sunitinib	Sutent (PPB)	☐ Tabrecta	☐ Tafinlar	
Tagrisso	☐ Talzenna	☐ Tarceva	Targretin gel	Tasigna (PPB)	
Tazverik	Temodar	Temozolomide	Tepmetko	Tibsovo	
Truqap	Tukysa	☐ Turalio*	Tykerb	☐ Valchlor	
☐ Vanflyta	☐ Venclexta	Verzenio	☐ Vitrakvi	Vizimpro	
Vonjo	Voranigo	☐ Votrient (PPB)	Welireg	Xalkori	
Xospata	Xpovio	Xtandi	Yonsa	Zejula	
Zelboraf	Zolinza	Zydelig	Zykadia	Zytiga	
PPB=Plan prefers	brand name				

Ben	eficiary Name (Last, First):
• • Veri	dications excluded from the above table may fall into one of the following categories: Available without prior authorization requirements New to market medication Covered as a medical claim fication of PA status can be found on the pharmacy vendor website: s://ar.primetherapeutics.com/drug-lookup
CRI	TERIA
Poli	cy guidelines:
•	Prior authorization criteria for oncology medications covered under this policy will be based on the FDA-approved label and support found in the NCCN treatment guidelines with NCCN level of evidence 1 or 2a unless otherwise noted with an asterisk*.
•	Medications noted with an asterisk follow DUR Board approved criteria found on the pharmacy vendor website: https://ar.primetherapeutics.com . Arimidex® (anastrazole) and Femara® (letrozole) will process at point-of-sale without a prior authorization if the beneficiary's medical history includes a female with breast cancer billed in the last 3 years.
•	Requests for an indication, dosage, age, or duration of treatment outside of the FDA-approved label and NCCN treatment recommendations are considered off-label.
•	Off-label requests will be reviewed for medical necessity on a case-by-case basis while referencing official compendia, peer-reviewed literature, and tumor board (case conference) review along with documentation submitted with the request.
•	All prior authorization requests must be submitted by or in consultation with an oncologist or hematologist.
•	Documentation supporting the prior authorization request must be submitted at the time of the request.
•	Quantity limits apply to all medications based on FDA-approved dosing.
pert	en submitting an initial prior authorization request for an oncology product, providing all inent information with the initial request will expedite reviews. At a minimum, the prescriber st submit:
	Current chart notes
□ T	Type of cancer with documentation of any mutations
	All previous therapies tried with timelines and response (i.e., medications and surgeries)
	Current labs specific to the type of cancer and treatment requesting (e.g., complete blood count, renal function labs, liver function panel)
	Specific imaging requirements per the package insert (e.g., MRI or CT imaging)
	Letter of medical necessity outlining the rationale for the treatment requested especially if the equest is off-label.
	Current weight or body surface area

Beneficiary Name (Last, First):
CRITERIA (CONTINUED)
☐ Dose requested.
$\ \square$ Pregnancy test results if recommended in the package insert.
☐ ECOG performance status score and medical necessity of treatment with ECOG score of 4
For prior authorization renewal requests, the prescriber must submit the following:
☐ Current chart notes
Current lab work
Current weight or body surface area
☐ Dose requested
☐ Documentation of current response to treatment
☐ Attestation that the patient exhibits a positive response from treatment without intolerable side effects.
Initial requests may be approved for 3 months, unless otherwise noted, with renewal pending a positive response to treatment without intolerable side effects. Prior authorization renewals may be approved for 3–6 months depending on the level of monitoring required for the treatment.
☐ Attachments
Prescriber Signature: Date:
(Required) Prescriber's original signature required; copied, stamped, or e-signature is not allowed. This certifies that the information provided in the Statement of Medical Necessity is accurate and substantiated by the patient's medical record.

Fax this form to 800-424-5851