

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.

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:	
	DEA Number:
Prescriber Phone:	Prescriber Fax:
DIAGNOSIS AND TREATMENT HISTORY	
Diagnosis:	
☐ New Therapy ☐ Renewal	
If renewal, duration of therapy (specific dates):	to
DRUG INFORMATION	
Drug Name:	
	Dosage Form:
Directions:	

Beneficiary Name (Last, First):						
DRUG INFORMATION (CONTINUED)						
Med	dications Covered as	s a Pharmacy Claim (se	lect requested medicati	on(s))		
Abiraterone	Fotivda	Lorbrena	Rozlytrek	Vitrakvi		
Afinitor	Fruzaqla	Lumakras	Rubraca	Vizimpro		
Akeega	Gavreto	Lynparza	Rydapt	Vonjo		
Alecensa	Gefitinib	Lytgobi	Scemblix	Voranigo		
Alunbrig	Gilotrif	Mekinist	Soltamox	Votrient		
Anastrazole*	Gomekli*	Mektovi	Sprycel	Welireg		
Arimidex*	☐ Ibrance	Mercaptopurine Suspension	Stivarga	☐ Xalkori		
Augtyro	Clusig	Nerlynx	Sunitinib	Xospata		
Ayvakit	☐ Idhifa	Ninlaro	Sutent	Xpovio		
Balversa	Imbruvica	Nubeqa	Tabrecta	Xtandi		
BESREMi	Imkeldi	Odomzo	Tafinlar	Yonsa		
Bosulif	Inlyta	Ogsiveo	Tagrisso	Zejula		
☐ Braftovi	Inqovi	Ojemda	Talzenna	Zelboraf		
Brukinsa	Inrebic	Ojjaara	Tarceva	Zolinza		
Cabometyx	Iressa	Onureg	☐ Targretin gel	Zydelig		
Calquence	Itovebi	Orgovyx	Tasigna	Zykadia		
Caprelsa	☐ Iwilfin	Orserdu	Tazverik	Zytiga		
Cometriq	Jakafi	Pazopanib	Temodar			
Copiktra	Jaypirca	Pemazyre	Temozolomide			
Cotellic	Kisqali	Piqray	Tepmetko			
Danziten	Kisqali/Femara	Pomalyst	Tibsovo			
☐ Dasatinib	☐ Koselugo*	Purixan	Truqap			
Daurismo	☐ Krazati	Qinlock	Tukysa			
Erivedge	Lapatinib	Retevmo	☐ Turalio*			
Erleada	Lazcluze	Revlimid	Tykerb			
Erlotinib	Lenalidomide	Revuforj	☐ Valchlor			
Everolimus	Lenvima	Rezlidhia	☐ Vanflyta			
Exkivity	Letrozole*	Rezurock*	☐ Venclexta			
☐ Femara*	Lonsurf	Romyimza	Verzenio			

Beneficiary Name (Last, First):
 Medications 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 Verification of PA status can be found on the pharmacy vendor website:
https://ar.primetherapeutics.com/drug-lookup
CRITERIA
Policy 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.
When submitting an initial prior authorization request for an oncology product, providing all pertinent information with the initial request will expedite reviews. At a minimum, the prescriber must submit:
Current chart notes
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)
\square 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 request is off-label.
Current weight or body surface area

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Beneficiary Name (Last, First):
☐ Dose requested.
☐ Pregnancy test results if recommended in the package insert.
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CRITERIA (CONTINUED)
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