

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:	
	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:	

DRUG INFORMATION (CONTINUED)						
	Medications Cove	ered as a Pharmacy Clai	im (select request	ed medication(s))		
Abiraterone	☐ Erivedge	☐ Jaypirca	☐ Ojemda	☐ Stivarga	☐ Vanflyta	
Afinitor	☐ Erleada	☐ Kisqali	☐ Ojjaara	Sunitinib	☐ Venclexta	
☐ Akeega	☐ Erlotinib	☐ Kisqali/Femara	Onureg	Sutent	☐ Verzenio	
Alecensa	☐ Everolimus	☐ Koselugo*	☐ Orgovyx	☐ Tabrecta	☐ Vitrakvi	
Alunbrig	☐ Exkivity	☐ Krazati	☐ Orserdu	☐ Tafinlar	☐ Vizimpro	
☐ Anastrazole*	☐ Femara*	☐ Lapatinib	☐ Pazopanib	☐ Tagrisso	☐ Vonjo	
☐ Arimidex*	☐ Fotivda	Lenalidomide	☐ Pemazyre	☐ Talzenna	☐ Votrient	
☐ Augtyro	☐ Fruzaqla	Lenvima	☐ Piqray	☐ Tarceva	☐ Welireg	
☐ Ayvakit	☐ Gavreto	☐ Letrozole*	☐ Pomalyst	☐ Targretin gel	☐ Xalkori	
Balversa	Gefitinib	Lonsurf	☐ Purixan	☐ Tasigna	☐ Xpovio	
BESREMi	Gilotrif	Lorbrena	Qinlock	☐ Tazverik	Xtandi	
Bosulif	☐ Ibrance	Lumakras	Retevmo	☐ Temodar	Yonsa	
☐ Braftovi	☐ Iclusig	Lynparza	Revlimid	☐ Temozolomide	Zelboraf	
Brukinsa	☐ Idhifa	Lytgobi	Rezlidhia	☐ Tepmetko	Zolinza	
☐ Cabometyx	☐ Imbruvica	☐ Mekinist	☐ Rezurock*	☐ Tibsovo	Zydelig	
☐ Calquence	☐ Inlyta	☐ Mektovi	Rozlytrek	☐ Torpenz	☐ Zykadia	
☐ Caprelsa	☐ Inqovi	☐ Nerlynx	Rubraca	☐ Truqap	Zytiga	
☐ Cometriq	☐ Inrebic	□ Ninlaro	Rydapt	☐ Tukysa		
☐ Copiktra	☐ Iressa	☐ Nubeqa	☐ Scemblix	☐ Turalio*		
☐ Cotellic	☐ Iwilfin	Odomzo	Soltamox	☐ Tykerb		
Daurismo	☐ Jakafi	☐ Ogsiveo	Sprycel	□ Valchlor		

Medications excluded from the above table may fall into one of the following categories:

• Available without prior authorization requirements

Beneficiary Name (Last, First):

- New to market medication
- Covered as a medical claim

Verification of PA status can be found on the pharmacy vendor website: https://ar.magellanrx.com/drug-lookup

Beneficiary Name (Last, First):
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.magellanrx.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)
☐ 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
☐ Dose requested.
Pregnancy test results if recommended in the package insert.
☐ ECOG performance status score and medical necessity of treatment with ECOG score of 4

Beneficiary Name (Last, First):
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:

Fax this form to 800-424-5851