

lt's Wholecare.

I. Requirements for Prior Authorization of Oncology Agents, Breast Cancer

A. Prescriptions That Require Prior Authorization

Prescriptions for Oncology Agents, Breast Cancer that meet any of the following conditions must be prior authorized:

- 1. A non-preferred Oncology Agent, Breast Cancer. See the Preferred Drug List (PDL) for the list of preferred Oncology Agents, Breast Cancer at: <u>https://papdl.com/preferred-drug-list</u>.
- 2. A prescription for letrozole.

B. Review of Documentation for Medical Necessity

In evaluating a request for prior authorization of a prescription for an Oncology Agent, Breast Cancer, the determination of whether the requested prescription is medically necessary will take into account the whether the beneficiary:

- 1. For a non-preferred agent, has a history of therapeutic failure, contraindication, or intolerance to the preferred Oncology Agents, Breast Cancer; **AND**
- 2. For letrozole, is being treated for a diagnosis that is indicated in the U.S. Food and Drug Administration (FDA)-approved package labeling OR a medically accepted indication, excluding use to promote fertility. The requesting prescriber must provide documentation from the medical record of the diagnosis

NOTE: If the beneficiary does not meet the clinical review guidelines listed above but, in the professional judgment of the physician reviewer, the services are medically necessary to meet the medical needs of the beneficiary, the request for prior authorization will be approved.

C. Clinical Review Process

Prior authorization personnel will review the request for prior authorization and apply the clinical guidelines in Section B. above to assess the medical necessity of a prescription for an Oncology Agent, Breast Cancer. If the guidelines in Section B. are met, the reviewer will prior authorize the prescription. If the guidelines are not met, the prior authorization request will be referred to a physician reviewer for a medical necessity determination. Such a request for prior authorization will be approved when, in the professional judgment of the physician reviewer, the services are medically necessary to meet the medical needs of the beneficiary.



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ONCOLOGY ANALYTICS

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Chemotherapy and Supportive Care Prior Authorization Request Form

REC	QUEST DAT	'E:	_ TREATMENT START DATE:				Standard 🛛 Expedited				
I.	MEMBER I	NFORMATION									
First:			Last:			DOB:		🗆 Male 🗆	🗆 Male 🛛 Female		
Height:			Weight:			BSA (m ²):					
Diagnosis:			ICD-10:			Stage (0-4):					
Insurance:			Line of Business (e.g., Medicare):			Member ID:					
II. ANTI-CANCER TREATMENT REQUEST New Retrospective Re-authorization											
#	Billing Code	Drug Name	2	Route	Dose			Cycles or Refills	(B = Buy	Billing Method (B = Buy & Bill or P = Pharmacy)	
1									В	ΠP	
2									□B	□ P	

III. SUPPORTING CARE DRUGS REQUESTED (see attached drug list for reference)

#	Billing Code	Drug Name	Route	Dose	Frequency & Schedule	Condition (e.g. nausea)	Billing N (B = Buy P = Pha	& Bill or
1							□В	□ P
2							□в	□ P
3							□В	□ P
4							□В	□ P
5							□В	□ P
If bone agents requested, select indication: 🗆 osteo 🗆 bone metastases 🗆 hypercalcemia 🗆 adjuvant breast cancer								
If ESAs requested, select indication: 🗆 CKD 🔅 CIA 🔅 MDS								

IV. PROVIDER AND PLACE OF TREATMENT INFORMATION

Ordering Provider:	NPI #:	TIN #:			
	Phone:	Fax:			
Treating Provider: (if different)	NPI #:	TIN #:			
Place of Treatment: (if different)	NPI #:	TIN #:			
Has the member been receiving cancer treatments from the requesting treating provider? Yes No Unknown					
Is treating provider in-network? 🗆 Yes 🖾 No 🗆 Unknown					
Requestor's Name:	Phone:	Fax:			

SUBMIT PROGRESS NOTES, CHEMO ORDERS, LABS, PATHOLOGY AND IMAGING RESULTS WITH REQUEST.

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SUPPORTIVE DRUGS REFERENCE PAGE

Note: This page is for reference and does not need to be faxed to Oncology Analytics.

Anti-emetics: nausea					
J1626 granisetron hydrochloride (Kytril) - IV					
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Q0166 granisetron hydrochloride (Kytril) - PO					
J1627 granisetron ER (Sustol) - SubQ					
J2405 ondansetron (Zofran) - IV					
Q0162	ondansetron (Zofran) - PO				
J2469 palonosetron (Aloxi) - IV					
J8655	netupitant/palonosetron HCl (Akynzeo) - PO				
J1454	netupitant/palonosetron HCl (Akynzeo) - IV				
J8670	rolapitant HCl (Varubi) - PO				
J1453	fosaprepitant dimeglumine (Emend) - IV				
J8501 aprepitant (Emend) – PO					
J0185 aprepitant (Cinvanti) - IV					
Request Notes: Include latest MD progress notes					
Bone Ag	ents				
J0897	denosumab (Xgeva) – SQ				
J0897	denosumab (Prolia) – SQ				
J3489	zoledronic acid (Zometa) - IV				
J3489	zoledronic acid (Reclast) - IV				
J2430	pamidronate (Aredia) – IV				
Request Notes: Include bone scan and bone density					
test results and latest MD progress notes.					

Erythropoiesis-stimulating agents (ESA): anemia						
J0885	J0885 epoetin alfa (Procrit) – SQ					
Q5106	Q5106 epoetin alfa-epbx (Retacrit) – SQ					
J0881 darbepoetin alfa (Aranesp) - SQ						
Request Notes: Include recent CBC, Iron Sat % and						
Ferritin. EPO level for initiation with MDS. Check						
indication for use on the request form: chronic kidney						
disease (CKD), chemotherapy induced anemia (CIA) or						
myelody	myelodysplastic syndrome (MDS)					
Granulo	cyte Colony Stimulating Growth Factors (G-					
CSF): neu	utropenia					
Q5101	filgrastim-sndz (Zarxio) – SQ					
J2505 pegfilgrastim (Neulasta) – SQ						
J1442 filgrastim (Neupogen) – SQ						
Q5110 filgrastim-aafi (Nivestym) – SQ						
J1447 tbo-filgrastim (Granix) – SQ						
Q5111	pegfilgrastim-cbqv (Udenyca) – SQ					
Q5108	peg filg rastim-jmdb (Fulphila) – SQ					
J9999	pegfilgras tim-bmez (Ziextenzo) – SQ					
J2820 sargramostim (Leukine) – SQ						
Request Notes: Include most recent CBC with diff,						
lowest ANC, any history of fever, febrile neutropenia,						
neutropenia on chemotherapy, current chemotherapy						
regimen, and a latest MD progress note.						