

ONCOLOGY STEP REGULATION PRIOR AUTHORIZATION GUIDELINES

Administered by

Mediimpact



**ONCOLOGY STEP REGULATION
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ABIRATERONE SUBMICRONIZED

Generic	Brand				
ABIRATERONE ACET, SUBMICRONIZED	YONSA				

GUIDELINES FOR USE

Our guideline named **ABIRATERONE SUBMICRONIZED (Yonsa)** requires the following rule(s) be met for approval:

- A. You have metastatic castration-resistant prostate cancer (mCRPC: prostate cancer that has spread to other parts of the body and longer responds to testosterone lowering treatment)
- B. The requested medication will be used in combination with an oral corticosteroid (such as prednisone, prednisolone, methylprednisolone)
- C. You meet ONE of the following:
 - 1. You received a bilateral orchiectomy (both testicles have been surgically removed)
 - 2. You have a castrate level of testosterone (your blood testosterone levels are less than 50 ng/dL)
 - 3. The requested medication will be used together with a gonadotropin-releasing hormone (GnRH) analog (such as Lupron Depot [leuprolide], Zoladex [goserelin], Firmagon [degarelix])

Effective: 08/01/23



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DENOSUMAB-XGEVA

Generic	Brand				
DENOSUMAB	XGEVA				

GUIDELINES FOR USE

Our guideline named **DENOSUMAB-XGEVA** requires the following rule(s) be met for approval:

- A. You have ONE of the following:
 - 1. Multiple myeloma (a type of blood cancer)
 - 2. Bone metastases from a solid tumor (tumor has spread to the bone)
 - 3. Giant cell tumor of bone (noncancerous tumor)
 - 4. Hypercalcemia (high level of calcium in the blood) of malignancy (cancer)
- B. **If you have multiple myeloma or bone metastases from a solid tumor, approval also requires:**
 - 1. Xgeva will be used to prevent skeletal-related events (such as bone fractures, bone pain requiring radiation [a type of treatment])
- C. **If you have giant cell tumor of bone, approval also requires:**
 - 1. Your tumor is unresectable (unable to remove by surgery) or surgical resection (removal) is likely to result in severe morbidity (illness)
- D. **If you have hypercalcemia of malignancy, approval also requires:**
 - 1. You are refractory (disease did not respond to treatment) to a prior bisphosphonate therapy (such as Fosamax [alendronate], Actonel [risedronate], Boniva [ibandronate])

Effective: 04/10/26



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LORLATINIB

Generic	Brand				
LORLATINIB	LORBRENA				

GUIDELINES FOR USE

Our guideline named **LORLATINIB (Lorbrena)** requires the following rule(s) be met for approval:

- A. You have metastatic non-small cell lung cancer (NSCLC: type of lung cancer that has spread to other parts of the body)
- B. You are 18 years of age or older
- C. Your tumors are anaplastic lymphoma kinase (ALK: a type of enzyme)-positive, as detected by a Food and Drug Administration (FDA)-approved test

Effective: 04/01/26



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PALBOCICLIB

Generic	Brand				
PALBOCICLIB	IBRANCE				

GUIDELINES FOR USE

Our guideline named **PALBOCICLIB (Ibrance)** requires the following rule(s) be met for approval:

- A. You have breast cancer
- B. Your cancer is hormone receptor (HR: a type of protein)-positive, human epidermal growth factor receptor 2 (HER2: a type of protein)-negative
- C. **If you are using Ibrance in combination with an aromatase inhibitor (such as anastrozole, letrozole, exemestane) as initial endocrine (hormone)-based therapy, approval also requires:**
 - 1. Your cancer is advanced or metastatic (cancer that has progressed or has spread to other parts of the body)
- D. **If you are using Ibrance in combination with fulvestrant (Faslodex), approval also requires:**
 - 1. Your cancer is advanced or metastatic (cancer that has progressed or has spread to other parts of the body)
 - 2. Your disease has worsened after endocrine (hormone) therapy (such as anastrozole, letrozole, tamoxifen)
- E. **If you are using Ibrance in combination with Itovebi (inavolisib) and fulvestrant (Faslodex), approval also requires:**
 - 1. Your cancer is locally advanced or metastatic (cancer that has spread from where it started to nearby tissue or lymph nodes or to other parts of the body)
 - 2. Your tumor has a PIK3CA mutation (abnormal change in a type of gene), as detected by a Food and Drug Administration (FDA)-approved test
 - 3. You have experienced disease recurrence (disease has returned) on or after completing adjuvant (add-on) endocrine (hormone) therapy (such as anastrozole, letrozole, tamoxifen)

Effective: 04/10/26



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