

STANDARD COMMERCIAL DRUG FORMULARY PRIOR AUTHORIZATION GUIDELINES

ABIRATERONE SUBMICRONIZED

Generic	Brand	HICL	GCN	Medi-Span	Exception/Other
ABIRATERONE	YONSA	44946		GPI-10	
ACET,				(2140601025)	
SUBMICRONIZED					

GUIDELINES FOR USE

- 1. Does the patient have a diagnosis of metastatic castration-resistant prostate cancer (mCRPC) and meet **ALL** of the following criteria?
 - The requested medication will be used in combination with an oral corticosteroid (e.g., prednisone, prednisolone, methylprednisolone)
 - The patient had a trial of or contraindication to Zytiga (abiraterone acetate)

If yes, continue to #2. If no, do not approve.

DENIAL TEXT: See the denial text at the end of the guideline.

- 2. Does the patient meet **ONE** of the following criteria?
 - The patient had a bilateral orchiectomy
 - The patient has a castrate level of testosterone (i.e., < 50 ng/dL)
 - The requested medication will be used concurrently with a gonadotropin-releasing hormone (GnRH) analog (e.g., Lupron Depot [leuprolide], Zoladex [goserelin], Firmagon [degarelix])

If yes, continue to #3. If no, do not approve.

DENIAL TEXT: See the denial text at the end of the guideline.

3. Is the patient concomitantly using a strong CYP3A4 inducer (e.g., phenytoin, carbamazepine, rifampin, rifabutin, rifapentine, phenobarbital)?

If yes, approve for 12 months by HICL or GPI-10 with a quantity limit of #8 per day. If no, approve for 12 months by HICL or GPI-10 with a quantity limit of #4 per day.

CONTINUED ON NEXT PAGE

Copyright © 2023 MedImpact Healthcare Systems, Inc. All rights reserved. This document is proprietary to MedImpact. MedImpact maintains the sole and exclusive ownership, right, title, and interest in and to this document.

Revised: 6/16/2023 Page 1 of 2



STANDARD COMMERCIAL DRUG FORMULARY PRIOR AUTHORIZATION GUIDELINES

ABIRATERONE SUBMICRONIZED

GUIDELINES FOR USE (CONTINUED)

DENIAL TEXT: *Some terms are already pre-defined in parenthesis. Please use these definitions if the particular text you need to use does not already have definition(s) in it.

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 no 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 have tried or have a contraindication to (harmful for) Zytiga (abiraterone acetate)
- D. You meet ONE of the following:
 - 1. You had 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])

Your doctor told us [INSERT PT SPECIFIC INFO PROVIDED]. We do not have information showing you [INSERT UNMET CRITERIA]. This is why your request is denied. Please work with your doctor to use a different medication or get us more information if it will allow us to approve this request.

RATIONALE

For further information, please refer to the Prescribing Information and/or Drug Monograph for Yonsa.

REFERENCES

Yonsa [Prescribing Information]. Cranbury, NJ: Sun Pharmaceuticals Industries, Inc.; March 2022.

Library	Commercial	NSA
Yes	Yes	No

Part D Effective: N/A Created: 03/23

Commercial Effective: 08/01/23 Client Approval: 06/23 P&T Approval: 07/23

Copyright © 2023 MedImpact Healthcare Systems, Inc. All rights reserved. This document is proprietary to MedImpact. MedImpact maintains the sole and exclusive ownership, right, title, and interest in and to this document.

Revised: 6/16/2023 Page 2 of 2