## Prior Authorization Request Form for **Zytiga 250mg (abiraterone)**



## JOHNS HOPKINS HEALTHCARE

7231 Parkway Drive, Suite 100, Hanover, MD 21076

FAX Completed Form and Applicable Progress Notes to: (410) 424-4037

## **USFHP Pharmacy Prior Authorization Form**

To be completed by Requesting provider				
Drug Name:	Strength:			
Dosage/Frequency (SIG):	Duration of Therapy:			

Questions? Contact the Pharmacy Dept at: (888) 819-1043, option 4

Clinical Documentation must accompany form in order for a determination to be made.

Step	Please complete patient and physician information (please print):				
1	Patient Name: Ph	ysician Name:			
	Address:	Address:			
	Sponsor ID #	Phone #:			
	Date of Birth:	Secure Fax #:			
Step	Please complete the clinical assessment:				
2	1. Yonsa is the Department of Defense's preferred CYP-17 Inhibitor agent. Has the patient tried Yonsa?	☐ Yes Proceed to question 3	□ No Proceed to question 2		
	2. Does the patient have or have they had a contraindication/inadequate response/adverse reaction to Yonsa that is not expected to occur with the requested agent?	☐ Yes Proceed to question 3	□ No STOP Coverage not approved		
	3. Is the patient greater than or equal to 18 years of age?	☐ Yes Proceed to question 4	□ No STOP Coverage not approved		
	4. Is the requested medication being prescribed by or in consultation with an oncologist or urologist?	☐ Yes Proceed to question 5	□ No STOP Coverage not approved		
	5. For which indication is the requested medication being prescribed?	☐ Metastatic castration-resistant prostate cancer (mCRPC) Proceed to question 8			
		☐ Metastatic castration-sensitive prostate cancer (mCSPC) Proceed to question 8			
		☐ Regional disease (TxN1M0) - Proceed to question 8			
		☐ Other indication - Pro	oceed to question 6		

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	o. Flease provide the diagnosis.			
			Proceed to	question <b>7</b>
7. Is the diagnosis cited in the National Comprehensive Cancer Network (NCCN) guidelines as a category 1, 2A, or 2B recommendation?			☐ Yes Proceed to question 11	□ No STOP Coverage not approved
	8. Is the patient receiving concomitant therapy with prednisone?	ith	☐ Yes	□ No
			Proceed to question 9	STOP Coverage not approved
	9. Is the patient receiving concomitant therapy w		□ Yes	□ No
	gonadotropin-releasing hormone (GnRH) analog (for example: Eligard, Lupron, Trelstar, or Zoladex)?		Proceed to question 11	Proceed to question 10
	10. Has the patient had bilateral orchiectomy?		□ Yes	□ No
			Proceed to question 11	STOP Coverage not approved
	11. Zytiga 250 mg is the DoD's preferred strength		□ Yes	□ No
	Is the prescription for Zytiga 250mg OR will the prescription be changed to the 250 mg? Note: the prescription is being changed to the 250 mg strength, please submit a new prescription.		Sign and date below	Proceed to question 12
	12. Please state why the patient cannot take multiple 250 mg tablets to achieve the patient's daily dose.			
			Sign and date	below
Step 3	I certify the above is true to the best of my kr Please sign and date:	nowled	lge.	
	Prescriber Signature		Date	
				[31 July 2019
r Interr	nal Use Only			
Approved:		Duration of Approval:month(s)		
Denied:		Authorized By:		
Incomplete/Other:			PA#:	
te Faxe	e Faxed to MD:		Date Decision Rendered:	