

MEDICAL PHARMACY PRIOR AUTHORIZATION REQUEST FORM

TESTOPEL®

For authorization, please answer each question and fax this form PLUS chart notes back to the U of U Health Plans Prior Authorization Department at 801-213-1547. Failure to submit clinical documentation to support this request will result in delay and/or denial of the request.

If you have prior authorization questions, please call for assistance: Healthy U: 855-856-5694, University of Utah Health Employees: 855-856-5690, Individual & Family Plans: 855-869-4769, Commercial Groups: 855-859-4892, MHC: 855-885-7695, Advantage U Part B: 888-605-0858

Date:	Member Name:	ID#:
DOB:	Gender:	Physician:
Office Phone:	Office Fax:	Office Contact:
Height/Weight:	HCPCS Code:	

Member must try formulary preferred drugs before a request for a non-preferred drug may be considered. If treatment with preferred products has not been successful, you must submit which preferred products have been tried, dates of treatment, and reason for failure. Reasons for failure must meet the Health Plan medical necessity criteria.

Product being requested: Testopel® (testosterone pellets)

Please note that testosterone injectable and topical testosterone are the plans preferred products.

Dosing/Frequency: _____

If the request is for reauthorization, proceed to reauthorization section.

Questions	Yes	No	Comments/Notes
1. Is the member 18 years of age or older?	<input type="checkbox"/>	<input type="checkbox"/>	
2. Is the member male?	<input type="checkbox"/>	<input type="checkbox"/>	
3. Does the member have a confirmed diagnosis of one of the following? <ul style="list-style-type: none"> • Primary hypogonadism • Hypogonadotropic hypogonadism 	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
4. Does the member have 2 confirmed early morning low serum testosterone levels at least 24 hours apart, defined as one of the following: <ul style="list-style-type: none"> • Total testosterone(TT) <464ng/dL (9.2nmol/L) for CDC certified TT assays • Free testosterone (FT) level less than the laboratory's normal reference range 	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
5. Has the member had at least a 6 month trial and failure of injectable testosterone?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
6. Has the member had at least a 6 month trial and failure of topical testosterone?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation

REAUTHORIZATION

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1. Is the requesting for reauthorization of therapy?	<input type="checkbox"/>	<input type="checkbox"/>	
2. Does clinical documentation show continued medical necessity and that the treatment is effective?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
What medications and/or treatment modalities have been tried in the past for this condition? Please document name of treatment, reason for failure, treatment dates, etc.			
Additional information:			
Physician Signature:			

****Failure to submit clinical documentation to support this request will result in delay and/or denial of the request****

Policy PHARM- M008
 Origination Date: 08/15/2019
 Reviewed/Revised Date: 08/19/2020
 Next Review Date: 08/19/2021
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