MHS PHARMACY BENEFIT TESTOSTERONES PRIOR AUTHORIZATION REQUEST FORM

MHS 550 N. Meridian St. Suite 101 Indianapolis, IN, 46204-1208 Phone: (877) 647-4848 Fax: (866) 399-0929



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Note: This form must be completed by the prescribing provider.

******All sections must be completed or the request will be returned******

Patient's Medicaid #	Date of Birth
Patient's Name	Prescriber's Name
Prescriber's IN License #	Specialty
Prescriber's NPI #	Prescriber's Signature
Return Fax # - -	Return Phone # - -
Check box if requesting retroactive PA	Date(s) of service requested for retroactive eligibility (if applicable):

Note: Submit PA requests for retroactive claims (dates of service prior to eligibility determination, but within established eligibility timelines) with dates of service prior to 30 calendar days of submission separately from current PA requests (dates of service 30 calendar days or less and going forward).

Requested Medication	Strength	Quantity	Dosage Regimen

INJECTABLE TESTOSTERONE AGENTS:

Please select one of the following:

- □ Member has a diagnosis of palliative treatment of breast cancer □ Yes □ No (Excluding Aveed, Testopel, and Xyosted)
- $\hfill\square$ Member has a diagnosis of delayed puberty $\hfill\square$ Yes \hfill No
- □ Total testosterone level is <= 350 ng/dL at therapy initiation (within the past 3 months) and <=1000 ng/dL

at renewal (within the past 6 months) (Documentation is required)

TOPICAL TESTOSTERONE AGENTS:

- 1. Member is 16 years of age or older \Box Yes \Box No
- 2. Total testosterone level is <= 350 ng/dL at therapy initiation (within the past 3 months) or <=1000 ng/dL at renewal (within the past 6 months) (Documentation is required) □ Yes □ No

The following criteria must be met to <u>exceed</u> the established topical testosterone quantity limit:

- 1. Member has utilized topical testosterone therapy for at least 14 days \Box Yes \Box No
- 2. Total testosterone level after at least 14 days of therapy is <= 400ng/dL (within the last 3 months)

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that apply.)
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Dng/dL at initiation
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2. Does the member have any of the following contraindications to therapy? (Check all that apply.)

□ Prostate cancer

Note: Approvable diagnoses include hypogonadism (documentation required of total testosterone <=350ng/dL at initiation [within past 3 months] or <=1000ng/dL at renewal [within past 6 months]).

Agents previously trialed:

DANOCRINE (DANAZOL):

1. Member diagnosis(es):_

Rationale for use of Kyzatrex over injectable testosterone formulations:

METHITEST (METHYLTESTOSTERONE):

- 1. Member diagnosis(es):
- 2. Does the member have any of the following contraindications to therapy? (Check all that apply.)
 - □ Breast cancer
 - □ Pregnancy
 - □ Prostate cancer

Note: Approvable diagnoses include hypogonadism (documentation required of total testosterone <=350ng/dL at initiation [within past 3 months] or <=1000ng/dL at renewal [within past 6 months]), delayed puberty, breast cancer, and cryptorchidism.

OXANDRIN (OXANDROLONE):

- 1. Member diagnosis(es):
- 2. Does the member have any of the following contraindications to therapy? (Check all that apply.)
 - □ Breast cancer
 - □ Hypercalcemia
 - □ Pregnancy
 - □ Prostate cancer
 - □ Severe renal disease

Note: Approvable diagnoses include adjunct treatment of severe burns during the catabolic and rehabilitative phases, AIDS-associated wasting syndrome, alcoholic hepatitis, cachexia.

TLANDO (TESTOSTERONE UNDECANOATE)

- 1. Member diagnosis(es):_
- 3. Does the member have any of the following contraindications to therapy? (Check all that apply.)
 - Breast cancer
 - □ Hypogonadal conditions not associated with structural or genetic etiologies
 - □ Pregnancy
 - □ Prostate cancer

Note: Approvable diagnoses include hypogonadism (documentation required of total testosterone <=350ng/dL at initiation [within past 3 months] or <=1000ng/dL at renewal [within past 6 months]).

Agents previously trialed:

Rationale for use of Tlando over injectable testosterone formulations AND Kyzatrex:

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