

**INDIANA HEALTH COVERAGE PROGRAMS (IHCP) PHARMACY BENEFIT
TESTOSTERONES PRIOR AUTHORIZATION REQUEST FORM**



CareSource Pharmacy Prior Authorization Form

**P.O. Box 8738
Dayton, OH 45401-8738
Fax: (866) 930-0019**

Today's Date

/ /

Non-Urgent ☐

Urgent ☐

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

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

| | |
|--|---|
| Patient's CareSource # <input type="text"/> | Date of Birth <input type="text"/> / <input type="text"/> / <input type="text"/> |
| Patient's Name | Prescriber's Name |
| Prescriber's IN License # <input type="text"/> | Specialty |
| Prescriber's NPI # <input type="text"/> | Office Contact |
| Prescriber's Fax: <input type="text"/> - <input type="text"/> - <input type="text"/> | Prescriber's Phone <input type="text"/> - <input type="text"/> - <input type="text"/> |
| Prescriber's Address | Date(s) of Service: _____ Start Date: _____ |
| Diagnosis | Diagnosis Code |

| Requested Medication | Strength | Quantity | Directions for Use |
|----------------------|----------|----------|--------------------|
| | | | |

DEPO-TESTOSTERONE, TESTOSTERONE CYPIONATE

Initial Authorization:

1. Please select one of the following:

- ☐ Member has a diagnosis of delayed puberty
- ☐ Member has a total testosterone level \leq 350 ng/dL within the past 3 months (Documentation is required)

2. For **ALL** indications:

Provider attests that member has none of the following contraindications to therapy: ☐ Yes ☐ No

- Breast cancer in a member assigned male at birth
- Pregnancy
- Prostate cancer

If **no**, please specify contraindication and medical rationale for use:

Prescriber Signature: _____ **Date:** _____

Reauthorization:

1. Total testosterone level is ≤ 1000 ng/dL within the past 6 months (Documentation is required) ☐ Yes ☐ No
2. Provider attests that member remains a candidate for treatment, indicating that they have not developed any of the contraindication(s) listed under initial authorization above ☐ Yes ☐ No

If **no**, please specify contraindication and medical rationale for use:

Prescriber Signature: _____ **Date:** _____

TESTOSTERONE ENANTHATE**Initial Authorization:**

1. Please select one of the following:

- ☐ Member has a diagnosis of delayed puberty
- Has the member had a previous trial and failure of ALL preferred injectable testosterone agents (reference PA criteria)? ☐ Yes ☐ No

If **no**, please provide medical justification for use of requested agent over ALL preferred injectable testosterone agents:

- ☐ Member has a total testosterone level ≤ 350 ng/dL within the past 3 months (Documentation is required)
- Has the member had a previous trial and failure of ALL preferred injectable testosterone agents (reference PA criteria)? ☐ Yes ☐ No

If **no**, please provide medical justification for use of requested agent over ALL preferred injectable testosterone agents:

- ☐ Member needs medication for palliative treatment of metastatic breast cancer

2. For **ALL** indications:

Provider attests that member has none of the following contraindications to therapy: ☐ Yes ☐ No

- Breast cancer in a member assigned male at birth
- Pregnancy
- Prostate cancer

If **no**, please specify contraindication and medical rationale for use:

Prescriber Signature: _____ **Date:** _____

Reauthorization:

1. Total testosterone level is ≤ 1000 ng/dL within the past 6 months (Documentation is required) ☐ Yes ☐ No
2. Has the member had a previous trial and failure of at least ONE preferred injectable testosterone agent (not required for palliative treatment of breast cancer) [reference PA criteria]? ☐ Yes ☐ No

If **no**, please provide medical justification for use of requested agent over ALL preferred injectable testosterone agents:

3. Provider attests that member remains a candidate for treatment, indicating that they have not developed any of the contraindication(s) listed under initial authorization above ☐ Yes ☐ No

If **no**, please specify contraindication and medical rationale for use:

Prescriber Signature: _____ **Date:** _____

AVEED, TESTOPEL PELLET, XYSOTED

Initial Authorization:

1. Please select one of the following:

- ☐ Member has a diagnosis of delayed puberty
- Has the member had a previous trial and failure of ALL preferred injectable testosterone agents (reference PA criteria)? ☐ Yes ☐ No

If **no**, please provide medical justification for use of requested agent over ALL preferred injectable testosterone agents:

- ☐ Member has a total testosterone level ≤ 350 ng/dL within the past 3 months (Documentation is required)
- Has the member had a previous trial and failure of ALL preferred injectable testosterone agents (reference PA criteria)? ☐ Yes ☐ No

If **no**, please provide medical justification for use of requested agent over ALL preferred injectable testosterone agents:

2. For **ALL** indications:

Provider attests that member has none of the following contraindications to therapy: ☐ Yes ☐ No

- Breast cancer in a member assigned male at birth
- Hypogonadal conditions not associated with structural or genetic etiologies (Xyosted ONLY)
- Pregnancy
- Prostate cancer

If **no**, please specify contraindication and medical rationale for use:

Prescriber Signature: _____ **Date:** _____

Reauthorization:

1. Total testosterone level is ≤ 1000 ng/dL within the past 6 months (Documentation is required) ☐ Yes ☐ No

2. Has the member had a previous trial and failure of at least ONE preferred injectable testosterone agent (reference PA criteria)? ☐ Yes ☐ No

If **no**, please provide medical justification for use of requested agent over ALL preferred injectable testosterone agents:

3. Provider attests that member remains a candidate for treatment, indicating that they have not developed any of the contraindication(s) listed under initial authorization above ☐ Yes ☐ No

If **no**, please specify contraindication and medical rationale for use:

Prescriber Signature: _____ Date: _____

TESTOSTERONE 1% (25 MG)/ 2.5 GM GEL PACKETS, TESTOSTERONE 1% (12.5 MG)/ACT GEL PUMP, TESTOSTERONE 1.62% (20.25 MG)/ACT METERED PUMP GEL, TESTIM 1% (50 MG)/5 GM GEL PACKETS

Initial Authorization:

1. Please select one of the following:

- ☐ Member is 16 years of age or older, has a total testosterone level ≤ 350 ng/dL within the past 3 months (Documentation is required), and is requesting to use topical testosterone **within the established quantity limits**

Requested dose: _____

- ☐ Member is 16 years of age or older, has a total testosterone level ≤ 400 ng/dL **while on topical testosterone therapy** (Documentation is required) and is requesting to **exceed established quantity limits**

Requested dose: _____

Member has utilized ≥ 14 days of topical testosterone therapies ☐ Yes ☐ No

Name of medication: _____

Dose: _____

Start and End date: _____

If **no**, please provide medical justification as to why member is requesting a dose beyond established quantity limits:

2. For **ALL** indications:

Provider attests that member has none of the following contraindications to therapy: ☐ Yes ☐ No

- Breast cancer in a member assigned male at birth
- Pregnancy
- Prostate cancer

If **no**, please specify contraindication and medical rationale for use:

Prescriber Signature: _____ Date: _____

Reauthorization:

1. Total testosterone level is ≤ 1000 ng/dL within the past 6 months (Documentation is required) ☐ Yes ☐ No
2. Provider attests that member remains a candidate for treatment, indicating that they have not developed any of the contraindication(s) listed under initial authorization above ☐ Yes ☐ No

If **no**, please specify contraindication and medical rationale for use:

Prescriber Signature: _____ **Date:** _____

NATESTO, TESTOSTERONE 1% (12.5 MG)/ACT GEL PUMP, TESTOSTERONE 1.62% (40.5 MG)/2.5 GM GEL PACKETS, TESTOSTERONE 1.62% (20.25 MG)/1.25 GM GEL PACKETS, TESTOSTERONE 2% (10 MG)/ACT METERED PUMP, TESTOSTERONE 30 MG/ACT SOLUTION, VOGELXO 1% (50 MG)/5 GM GEL PACKETS, VOGELXO 1% (12.5 MG)/ACT GEL PUMP

Initial Authorization:

1. Please select one of the following:

- ☐ Member is 16 years of age or older, has a total testosterone level ≤ 350 ng/dL within the past 3 months (Documentation is required), and is requesting to use topical testosterone **within the established quantity limits**

Requested dose: _____

- ☐ Member is 16 years of age or older, has a total testosterone level ≤ 400 ng/dL **while on topical testosterone therapy** (Documentation is required) and is requesting to **exceed established quantity limits**

Requested dose: _____

Member has utilized ≥ 14 days of topical testosterone therapies: ☐ Yes ☐ No

Name of medication: _____

Dose: _____

Start and End date: _____

If **no**, please provide medical justification as to why member is requesting a dose beyond established quantity limits:

2. Previous trial and failure of ALL preferred topical testosterone agents (reference PA criteria) ☐ Yes ☐ No

If **no**, please provide medical justification for use of requested agent over ALL preferred topical testosterone agents:

3. For **ALL** indications:

Provider attests that member has none of the following contraindications to therapy: ☐ Yes ☐ No

- Breast cancer in a member assigned male at birth
- Pregnancy

- Prostate cancer

If **no**, please specify contraindication and medical rationale for use:

Prescriber Signature: _____ **Date:** _____

Reauthorization:

1. Total testosterone level is ≤ 1000 ng/dL within the past 6 months (Documentation is required) ☐ Yes ☐ No

2. Previous trial and failure of at least ONE preferred topical testosterone agent ☐ Yes ☐ No

If **no**, please provide medical justification for use of requested agent over ALL preferred topical testosterone agents:

3. Provider attests that member remains a candidate for treatment, indicating that they have not developed any of the contraindication(s) listed under initial authorization above ☐ Yes ☐ No

If **no**, please specify contraindication and medical rationale for use:

Prescriber Signature: _____ **Date:** _____

Initial Authorization (approval up to 6 months):

1. Member diagnosis(es): _____

Note: Approvable diagnoses include angioedema prophylaxis for heredity angioedema, autoimmune hemolytic anemia, discoid lupus erythematosus, endometriosis, fibrocystic breast disease, myelosclerosis with myeloid metaplasia

2. For **ALL** indications:

Provider attests that member has none of the following contraindications to therapy: ☐ Yes ☐ No

- Active or history of thrombosis or thromboembolic disease
- Androgen-dependent tumor
- Cardiac disease
- Porphyria
- Pregnancy or breast-feeding
- Severe hepatic disease
- Severe renal disease
- Undiagnosed genital bleeding

If **no**, please specify contraindication and medical rationale for use:

Prescriber Signature: _____ **Date:** _____

Reauthorization (approval up to 6 months):

1. Documentation from prescriber indicating continued benefit from the medication without significant adverse events ☐ Yes ☐ No
2. Provider attests that member remains a candidate for treatment, indicating that they have not developed any of the contraindication(s) listed under initial authorization above ☐ Yes ☐ No

If **no**, please specify contraindication and medical rationale for use:

Prescriber Signature: _____ **Date:** _____

JATENZO (TESTOSTERONE UNDECANOATE):**Initial Authorization:**

1. Member is 18 years of age or older ☐ Yes ☐ No
2. Member has a diagnosis of hypogonadism with a total testosterone level ≤ 350 ng/dL within the past 3 months (Documentation is required) ☐ Yes ☐ No
3. Previous trial and failure of at least ONE preferred injectable testosterone agent (reference PA criteria) ☐ Yes ☐ No

If **no**, please provide medical justification for use of requested agent over ALL preferred injectable testosterone agents:

4. For **ALL** indications:

Provider attests that member has none of the following contraindications to therapy: ☐ Yes ☐ No

- Breast cancer in a member assigned male at birth
- Hypogonadal conditions not associated with structural or genetic etiologies
- Pregnancy
- Prostate cancer

If **no**, please specify contraindication and medical rationale for use:

Prescriber Signature: _____ **Date:** _____

Reauthorization:

1. Total testosterone level is ≤ 1000 ng/dL within the past 6 months (Documentation is required) ☐ Yes ☐ No
2. Provider attests that member remains a candidate for treatment, indicating that they have not developed any of the contraindication(s) listed under initial authorization above ☐ Yes ☐ No

If **no**, please specify contraindication and medical rationale for use:

Prescriber Signature: _____ **Date:** _____

3. Previous trial and failure of at least ONE preferred injectable testosterone agent (reference PA criteria)

☐ Yes ☐ No

If **no**, please provide medical justification for use of requested agent over ALL preferred injectable testosterone agents:

KYZATREX (TESTOSTERONE UNDECANOATE):

Initial Authorization:

1. Member is 18 years of age or older ☐ Yes ☐ No

2. Member has a diagnosis of hypogonadism and a total testosterone level ≤ 350 ng/dL within the past 3 months (Documentation is required) ☐ Yes ☐ No

3. Previous trial and failure of at least ONE preferred injectable testosterone agent (reference PA criteria)

☐ Yes ☐ No

If **no**, please provide medical justification for use of requested agent over ALL preferred injectable testosterone agents:

4. For **ALL** indications:

Provider attests that member has none of the following contraindications to therapy: ☐ Yes ☐ No

- Breast cancer
- Hypogonadal conditions not associated with structural or genetic etiologies
- Pregnancy
- Prostate cancer

If **no**, please specify contraindication and medical rationale for use:

Prescriber Signature: _____ **Date:** _____

Reauthorization:

1. Total testosterone level is ≤ 1000 ng/dL within the past 6 months (Documentation is required) ☐ Yes ☐ No

2. Provider attests that member remains a candidate for treatment, indicating that they have not developed any of the contraindication(s) listed under initial authorization above ☐ Yes ☐ No

If **no**, please specify contraindication and medical rationale for use:

Prescriber Signature: _____ **Date:** _____

3. Previous trial and failure of at least ONE preferred injectable testosterone agent (reference PA criteria)

☐ Yes ☐ No

If **no**, please provide medical justification for use of requested agent over ALL preferred injectable testosterone agents:

METHITEST (METHYLTESTOSTERONE)

Initial Authorization (approval up to 6 months):

1. Please select one of the following:

- ☐ Member has a diagnosis of cryptorchidism
- ☐ Member has a diagnosis of delayed puberty
- ☐ Member has a diagnosis of hypogonadism (primary or hypogonadotropic) with a total testosterone ≤ 350 ng/dL within the past 3 months (Documentation is required)
- ☐ Member needs medication for palliative treatment of metastatic breast cancer

2. Previous trial and failure of at least ONE preferred injectable testosterone agent (reference PA criteria)

☐ Yes ☐ No

If **no**, please provide medical justification for use of requested agent over ALL preferred injectable testosterone agents:

3. For **ALL** indications:

Provider attests that member has none of the following contraindications to therapy: ☐ Yes ☐ No

- Breast cancer in a member assigned male at birth
- Pregnancy
- Prostate cancer

If **no**, please specify contraindication and medical rationale for use:

Prescriber Signature: _____ **Date:** _____

Reauthorization (approval up to 6 months):

1. Please select one of the following:

- ☐ Member has a diagnosis of hypogonadism and a total testosterone level ≤ 1000 ng/dL within the past 6 months (Documentation is required)
- ☐ Member has a diagnosis of delayed puberty, palliative treatment of metastatic breast cancer, or cryptorchidism AND prescriber has submitted documentation indicating continued benefit from the medication without significant adverse events:

2. For **ALL** indications:

Provider attests that member remains a candidate for treatment, indicating that they have not developed any of the contraindication(s) listed under initial authorization above ☐ Yes ☐ No

If **no**, please specify contraindication and medical rationale for use:

Prescriber Signature: _____ **Date:** _____

3. Previous trial and failure of at least ONE preferred injectable testosterone agent (reference PA criteria)

☐ Yes ☐ No

If **no**, please provide medical justification for use of requested agent over ALL preferred injectable testosterone agents:

OXANDRIN (OXANDROLONE):

Initial Authorization (approval up to 6 months):

1. Member diagnosis(es): _____

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

2. For ALL indications:

Provider attests that member has none of the following contraindications to therapy: ☐ Yes ☐ No

- Breast cancer
- Hypercalcemia
- Pregnancy
- Prostate cancer
- Severe renal disease

If **no**, please specify contraindication and medical rationale for use:

Prescriber Signature: _____ **Date:** _____

Reauthorization (approval up to 6 months):

1. Documentation from prescriber indicating continued benefit from the medication without significant adverse events ☐ Yes ☐ No

2. Provider attests that member remains a candidate for treatment, indicating that they have not developed any of the contraindication(s) listed under initial authorization above ☐ Yes ☐ No

If **no**, please specify contraindication and medical rationale for use:

Prescriber Signature: _____ **Date:** _____

TLANDO (TESTOSTERONE UNDECANOATE)

Initial Authorization:

1. Member is 18 years of age or older ☐ Yes ☐ No

2. Member has a diagnosis of hypogonadism and a total testosterone level ≤ 350 ng/dL within the past 3 months (Documentation is required) ☐ Yes ☐ No

3. Previous trial and failure of at least ONE preferred injectable testosterone agent (reference PA criteria)
☐ Yes ☐ No

If **no**, please provide medical justification for use of requested agent over ALL preferred injectable testosterone agents:

4. For **ALL** indications:

Provider attests that member has none of the following contraindications to therapy: ☐ Yes ☐ No

- Breast cancer

- Hypogonadal conditions not associated with structural or genetic etiologies
- Pregnancy
- Prostate cancer

If **no**, please specify contraindication and medical rationale for use:

Prescriber Signature: _____ **Date:** _____

Reauthorization:

1. Total testosterone level is ≤ 1000 ng/dL within the past 6 months (Documentation is required) ☐ Yes ☐ No
2. Prescriber attests that member remains a candidate for treatment, indicating that they have not developed any of the contraindication(s) listed under initial authorization above ☐ Yes ☐ No

If **no**, please specify contraindication and medical rationale for use:

Prescriber Signature: _____ **Date:** _____

3. Previous trial and failure of at least ONE preferred injectable testosterone agent (reference PA criteria)
☐ Yes ☐ No

If **no**, please provide medical justification for use of requested agent over ALL preferred injectable testosterone agents:

I attest that the provided information above is accurate:

Physician Signature: _____ **Date:** _____

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