

## Iowa Department of Human Services

## **FAX Completed Form To** 1 (800) 574-2515

**Provider Help Desk** 1 (877) 776-1567

## Request for Prior Authorization TESTOSTERONE PRODUCTS

(PLEASE PRINT – ACCURACY IS IMPORTANT)

IA Medicaid Member ID #	Patient name	DOB			
Patient address					
Provider NPI	Prescriber name	Phone			
Prescriber address Fax					
Pharmacy name	Address	Phone			
Prescriber must complete all information above. It must be legible, correct, and complete or form will be returned.					
Pharmacy NPI	Pharmacy fax	NDC			

Prior authorization is required for testosterone products. Payment will be considered with documentation of a specific testicular or hypothalamic/pituitary disease (primary hypogonadism or hypogonadotropic hypogonadism) that results in classic hypogonadism. Requests for FDA approved indications other than hypogonadism will not be subject to prior authorization criteria with adequate documentation of diagnosis. Payment for non-preferred testosterone products will be authorized only for cases in which there is documentation of previous trials and therapy failures with two preferred agents. Requests for erectile dysfunction, infertility, and age-related hypogonadism will not be considered. Payment will be considered under the following conditions:

- 1) Patient is male and 18 years of age or older (or 12 years of age and older for testosterone cypionate); and
- 2) Patient has two (2) morning pre-treatment testosterone levels below the lower limit of the normal testosterone reference range of the individual laboratory used (attach results); and
- 3) Patient has primary hypogonadism or hypogonadotropic hypogonadism (further defined below)
  - Primary hypogonadism (congenital or acquired) caused by testicular failure due to one of the following: cryptorchidism, bilateral torsion, orchitis, vanishing testes syndrome, orchiectomy, Klinefelter's syndrome, chemotherapy, toxic damage from alcohol or heavy metals
  - Hypogonadotropic hypogonadism: idiopathic gonadotropin or luteinizing hormone-releasing (LHRH) deficiency, pituitary-hypothalamic injury from tumors, trauma, or radiation
- 4) Patient does not have:
  - Breast or prostate cancer
  - Palpable prostate nodule or prostate-specific antigen (PSA) > 4ng/mL
  - Hematocrit > 50%
  - Untreated severe obstructive sleep apnea
  - Severe lower urinary tract symptoms
  - Uncontrolled or poorly controlled heart failure

If criteria for coverage are met, initial authorizations will be given for 3 months. Requests for continuation of therapy will require the following:

- An updated testosterone level (attach result); and
- Documentation the patient has not experienced a hematocrit > 54% or an increase in PSA > 1.4ng/mL in the past 12 months.

The required trials may be overridden when documented evidence is provided that use of these agents would be medically contraindicated.

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Preferred Androderm Testosterone Cypionate Testosterone Enanthate Testosterone Gel 1% Packets	Non-Preferred Androgel Aveed Depo-Testostero Fortesta	☐ Jaten: ☐ Methione ☐ Methy ☐ Nates	test Itestostero	one  Testoste	☐ Tlando one Gel 1.62% ☐ Xyosted rone Gel Pump ☐ Vogelxo rone Topical Solution	
Strength Dosa	ige Instructions			Quantity	Days Supply	
Complete for diagnosis of hypogo	nadism:					
<ul> <li>□ Primary Hypogonadism (congeni</li> <li>□ Cryptorchidism □ Bilatera</li> <li>□ Klinefelter's syndrome □ 0</li> <li>□ Other:</li> <li>□ Hypogonadotropic Hypogonadism</li> </ul>	torsion	☐ Vanishir	ng testes sy m alcohol c	ndrome	-	
Idiopathic gonadotropin or luteinizing hormone-releasing (LHRH) deficiency						
☐ Pituitary-hypothalamic injury from tumors, trauma, or radiation						
Please indicate setting in which medication is to be administered:  List & attach results of two (2) morning pre-treatment testosterone levels below the lower limit of the normal testosterone reference range of the individual laboratory used:  Level 1: Date: Level 2: Date:						
Does patient have any of the follow Breast or prostate cancer: Palpable prostate nodule or prostate- Hematocrit > 50%: Untreated severe obstructive sleep a Severe lower urinary tract symptoms Uncontrolled or poorly controlled hea	specific antigen (PSA)	Yes	No Do No No No No	Yes	No	
Renewal Requests:						
List & attach updated testosterone level: Level:			[	Date:		
Hematocrit > 54%: Increase in PSA > 1.4ng/mL: Other medical conditions to consider						
Prescriber signature (Must match pre		<i>,</i>		Date of submiss	sion	

**IMPORTANT NOTE:** In evaluating requests for prior authorization the consultant will consider the treatment from the standpoint of medical necessity only. If approval of this request is granted, this does not indicate that the member continues to be eligible for Medicaid. It is the responsibility of the provider who initiates the request for prior authorization to establish by inspection of the member's Medicaid eligibility card and, if necessary by contact with the county Department of Human Services, that the member continues to be eligible for Medicaid.

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