



Please complete ALL information below and fax your request to 1-888-671-5285

### Androgens Prior Authorization Request Form (Page 1 of 2)

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Member Information (required)			Provider Information (required)		
Member Name:			Provider Name:		
Insurance ID#:			NPI#:	Specialty:	
Date of Birth:			Office Phone:		
Street Address:			Office Fax:		
City:	State:	Zip:	Office Street Address:		
Phone:			City:	State:	Zip:

Medication Information (required)		
Medication Name:	Strength:	Dosage Form:
<input type="checkbox"/> Check if <b>generic substitution</b> is acceptable <input type="checkbox"/> Check if request is for <b>continuation of therapy</b>		Directions for Use:

Clinical Information (required)
<p><b>Select the diagnosis below:</b></p> <input type="checkbox"/> Delayed puberty <input type="checkbox"/> Gender dysphoria <input type="checkbox"/> Hypogonadism (e.g., testicular hypofunction, male hypogonadism, ICD-10 code E29.1) <input type="checkbox"/> Inoperable breast cancer, palliative treatment <input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s): _____
<p><b>Continuation of therapy:</b>  Is this a continuation of testosterone therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No</p>
<p><b>Which gender was the patient at birth?</b> (Select from one of the options below)  <input type="checkbox"/> Female  <input type="checkbox"/> Male</p>
<p><b>Medication History:</b>  <b>For nasal, oral, &amp; topical testosterone requests, select the medications the patient has a failure, contraindication, or intolerance to:</b>  <input type="checkbox"/> Androderm (testosterone patch)      <input type="checkbox"/> Androgel 1.62% gel (testosterone gel)      <input type="checkbox"/> Fortesta (testosterone gel)  <input type="checkbox"/> Androgel 1% (testosterone gel)      <input type="checkbox"/> Androgel 1.62% pump (testosterone pump)      <input type="checkbox"/> Testim (testosterone gel)</p> <p><b>For implant &amp; injectable testosterone requests, select the medications the patient has a failure, contraindication, or intolerance to:</b>  <input type="checkbox"/> Testosterone cypionate      <input type="checkbox"/> Testosterone enanthate</p>
<p><b>Hypogonadism in men (e.g., testicular hypofunction, male hypogonadism, ICD-10 code E29.1):</b>  Does the patient have a condition that may cause altered sex-hormone binding globulin (SHBG) (e.g., thyroid disorder, HIV disease, liver disorder, diabetes, obesity)? <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Does the patient have a history of one of the following: Bilateral orchiectomy, panhypopituitarism, or a genetic disorder known to cause hypogonadism (e.g., congenital anorchia, Klinefelter's syndrome)? <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p><b>Laboratory information:</b>  <b>Total testosterone level:</b>  Does the patient have two pre-treatment serum total testosterone levels less than 300ng/dL (&lt; 10.4 nmol/L) or less than the reference range for the lab? <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p><b>Calculated free or bioavailable testosterone level:</b>  Does the patient have one pre-treatment calculated free or bioavailable testosterone level less than 5 ng/dL (&lt; 0.17 nmol/L) or less than the reference range for the lab? <input type="checkbox"/> Yes <input type="checkbox"/> No</p>

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## Androgens Prior Authorization Request Form (Page 2 of 2)

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### Gender dysphoria:

Is the patient a female-to-male transsexual?  Yes  No

### Reauthorization:

**If this is a reauthorization request, answer the following questions:**

Is the patient's follow-up **total serum** testosterone level drawn within the past 6 months for patients new to testosterone therapy, or 12 months for patients continuing testosterone therapy, within or below the normal limits of the reporting lab?  Yes  No

Is the patient's follow-up **total serum** testosterone level drawn within the past 6 months for patients new to testosterone therapy, or 12 months for patients continuing testosterone therapy, outside of the upper limits of normal for the reporting lab and the dose has been adjusted?  Yes  No

Is the patient's follow-up **calculated free or bioavailable** testosterone level drawn within the past 6 months for patients new to testosterone therapy, or 12 months for patients continuing testosterone therapy, within or below the normal limits of the reporting lab?  Yes  No

Is the patient's follow-up **calculated free or bioavailable** testosterone level drawn within the past 6 months for patients new to testosterone therapy, or 12 months for patients continuing testosterone therapy, outside of the upper limits of normal for the reporting lab and the dose has been adjusted?  Yes  No

Does the patient have a condition that may cause altered sex-hormone binding globulin (SHBG) (e.g., thyroid disorder, HIV disease, liver disorder, diabetes, obesity)?  Yes  No

**Are there any other comments, diagnoses, symptoms, medications tried or failed, and/or any other information the physician feels is important to this review?**

Please note: This request may be denied unless all required information is received.