SENTARA COMMUNITY PLAN (MEDICAID)

MEDICAL PRIOR AUTHORIZATION/STEP-EDIT REQUEST*

Directions: The prescribing physician must sign and clearly print name (preprinted stamps not valid) on this request. All other information may be filled in by office staff; fax to 1-844-305-2331. No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct. If information provided is not complete, correct, or legible, authorization can be delayed.

Testosterone Replacement Therapy (TRT) Injectables

(MUST be purchased by Physician's Office)

erectile dysfunction drugs.

	NON-PREFERRED	
□ Aveed [®] (testosterone undecanoate) IM injection (J3145)	□ Azmiro (testosterone cypionate) IM injection (J1072) □ Testopel® (testosterone pellets) (11980/S0189)	
MEMBER & PRESCRIE	BER INFORMATION: Authorization may be delayed if incomplete.	
Member Name:		
	Date of Birth:	
Prescriber Name:		
	Date:	
Office Contact Name:		
Phone Number:	: Fax Number:	
NPI #:		
DRUG INFORMATION	: Authorization may be delayed if incomplete.	
Drug Form/Strength:		
	Length of Therapy:	
Diagnosis:	ICD Code, if applicable:	
Weight (if applicable):	Date weight obtained:	
	ng this box, the timeframe does not jeopardize the life or health of the mem in maximum function and would not subject the member to severe pain.	
NOTE: For the hypogonadist	m indication, testosterone drugs CANNOT be used in conjunction with	

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Maximum Dosage:

- Aveed 3 mL (750 mg) injected intramuscularly, followed by 3 mL (750 mg) injected after 4 weeks, then 3 mL (750 mg) injected every 10 weeks
- Azmiro 50 to 400 mg administered every two to four weeks as an intramuscular injection
- Testopel 75 mg implantable pellet; 6 pellets per 90-day supply (6 billable units every 90 days)
 - o 11980 = CPT1 code for subcutaneous hormone pellet implantation
 - o S0189 = HCPCS code for testosterone pellet, 75 mg

CLINICAL CRITERIA: Check below all that apply. All criteria must be met for approval. To support each line checked, all documentation, including lab results, diagnostics, and/or chart notes, must be provided or request may be denied.

Length of Authorization: 12 months

- ☐ Member must meet **ONE** of the following:
 - ☐ Member has Partial Androgen Insensitivity Syndrome with male gender identity/gender dysphoria or delayed male puberty

OR

- ☐ Member has hypogonadism confirmed by **BOTH** of the following:
 - □ TWO (2) MORNING (6AM to 11AM) testosterone levels obtained on different dates that are below 300 ng/dL or below the lower limit of normal for the reference range (attach lab results with reference ranges from the laboratory for both)

First testosterone level:	

- □ Repeat testosterone or free testosterone level:
- ☐ Member has at least <u>one</u> specific symptom and at least <u>two</u> non-specific symptoms [<u>NOTE</u>: If '<u>decreased spontaneous erections</u>' is the <u>ONLY</u> symptom documented in chart notes, the request will be denied as testosterone replacement is <u>EXCLUDED</u> from coverage for sexual dysfunction]:

Specific Symptoms: $(\geq 1 \text{ of the following})$	Non-Specific Symptoms: $(\geq 2 \text{ of the following})$
☐ Incomplete or delayed sexual development	☐ Decreased energy, motivation, initiative and
☐ Reduced sexual desire (libido) and activity	self-confidence
☐ Decreased spontaneous erections*	□ Depressed mood
☐ Breast discomfort, gynecomastia	□ Poor concentration and memory
☐ Loss of body hair (axillary, facial, and/or	☐ Sleep disturbances, increased sleepiness
pubic)	☐ Mild anemia (Hgb 10-11 gm/dL)
☐ Small testes (<5 mL) or shrinking testes	☐ Reduced muscle bulk and strength due to
☐ Low or zero sperm count	cachexia
☐ Height loss, low trauma fracture or low	☐ Increased body fat, BMI
bone mineral density	☐ Diminished physical or work performance
☐ Hot flushes, sweats	

	AND
	Member has had an inadequate response, contraindication or intolerance to at least a three-month trial with a topical agent e.g., testosterone gel, testosterone patch, testosterone topical solution, testosterone nasal gel (verified by pharmacy paid claims)
	(Please document date/name of drug)
	<u>AND</u>
	Member has had an inadequate response, contraindication or intolerance to at least a three-month trial of a preferred intramuscular testosterone medication e.g., testosterone cypionate, testosterone enanthate or testosterone undecanoate (verified by pharmacy paid claims)
	(Please document date/name of drug)
Med	lication being provided by (check box below that applies):
	Physician's office OR Specialty Pharmacy
review treatm	gent reviews: Practitioner should call Sentara Pre-Authorization Department if they believe a standard would subject the member to adverse health consequences. Sentara's definition of urgent is a lack of ent that could seriously jeopardize the life or health of the member or the member's ability to regain num function.
**	*Use of samples to initiate therapy does not meet step edit/preauthorization criteria. **
*Pre	evious therapies will be verified through pharmacy paid claims or submitted chart notes.