US Family Health Plan Prior Authorization Request Form for

Androgeltestosterone1% and 1.62% MDP, geland gelpackets, 2% testosterone solution MDP, 1% testosterone gel in unit-dose tubes (Testim, Vogelxo, generics)

To be completed and signed by the prescriber. To be used only for prescriptions which are to be filled through the Department of Defense (DoD) US Family Health Plan Pharmacy Program. US Family Health Plan is a TRICARE contractor for DoD.

The completed form may be faxed to 855-273-5735

OR

The patient may attach the completed form to the prescription and **mail** it to:

Attn: Pharmacy, 77 Warren St, Brighton, MA 02135

QUESTIONS? Call 1-877-880-7007

Clinical d	ocun	nentation including labs may be required.					
Initial the required.		approves for 1 year, renewal approves indefinitely. For	renewal of the	rapy, an initial Tricare prior	authorization approval is		
Step	Ple	ase complete patient and physician information ((please print):				
1	Patient Name: Address:		Physician Name:				
			Address:				
	Sno	onsor ID #	_ Pho	one #:			
	Date of Birth:		Secure Fax #:				
Step	Ple	Please complete the clinical assessment:					
2	1.	Will the requested medication be used to enhance athletic		□ Yes	□ No		
		performance?		STOP	Proceed to question 2		
				Coverage not approved			
	2. Will the requested medication be used concomitation other testosterone products?		tantly with	□ Yes	□ No		
				STOP	Proceed to question 3		
				Coverage not approved			
	3.	3. Has the patient received this medication under the TRICARE benefit in the last 6 months? Please choose "No" if the patient did not previously have a TRICARE approved PA for the requested medication.		□ Yes	□ No		
				Proceed to question 4	Proceed to question 6		
	4.	Has the patient had a positive response to therapy?		☐ Yes	□ No		
				Proceed to question 5	STOP		
					Coverage not approved		
	5.	Do the benefits of continued therapy outweigh the risks?		□ Yes	□ No		
				Sign and date on page 3	STOP		
					Coverage not approved		

6. What is the indication or diagnosis?		☐ Hypogonadism - Proceed to question 7	
		☐ Female-to-male gender-affirming hormone therapy in a natal female patient (assigned female at birth) - Proceed to question 14	
		☐ Other - Proceed to que	estion 22
7.	Is the patient a male who is 18 years of age or older?	□ Yes	□ No
		Proceed to question 8	STOP
			Coverage not approved
8.	Does the patient have a confirmed diagnosis of hypogonadism as evidenced by morning total serum	□ Yes	□ No
	testosterone levels below 300 ng/dL taken on at least two separate occasions?	Proceed to question 10	Proceed to question 9
9.	Is testosterone being prescribed by an endocrinologist or urologist who has made the diagnosis of hypogonadism	□ Yes	□ No
	based on unequivocally and consistently low serum total	Proceed to question 10	STOP
	testosterone or free testosterone levels?		Coverage not approved
10.	Is the patient experiencing signs and symptoms associated with hypogonadism?	□ Yes	□ No
		Proceed to question 11	STOP
			Coverage not approved
11.	Has the provider investigated the etiology of the low testosterone levels?	☐ Yes	□ No
		Proceed to question 12	STOP
			Coverage not approved
12.	Has the provider assessed the risks versus benefits of initiating testosterone therapy in this patient?	□ Yes	□ No
	initiating testosterone therapy in this patient:	Proceed to question 13	STOP
			Coverage not approved
13.	Does the provider acknowledge that testosterone therapy is clinically appropriate and needed?	☐ Yes	□ No
	is clinically appropriate and needed:	Proceed to question 23	STOP
			Coverage not approved
14.	Is the patient greater than or equal to 14 years of age?	□ Yes	□ No
		Proceed to question 15	STOP
			Coverage not approved
15.	Does the patient have a diagnosis of gender dysphoria made by a TRICARE-authorized mental health provider according to the most current edition of the DSM?	□ Yes	□ No
		Proceed to question 16	STOP
			Coverage not approved

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16.	Is the requested medication being prescribed by an endocrinologist or a physician who specializes in the treatment of transgender patients?		☐ Yes Proceed to question 17	□ No STOP		
			·	Coverage not approved		
17.	Is the patient an adult, or an adolescent wi		□ Yes	□ No		
	irreversible treatment?	ior tills partially	Proceed to question 18	STOP		
				Coverage not approved		
18.	8. Has the patient experienced puberty to at least Tanner stage 2?		□ Yes	□ No		
	31age 2:		Proceed to question 19	STOP		
				Coverage not approved		
19.	Is the patient a biological female of childbe	earing potential?	☐ Yes	□ No		
		Proceed to question 20	Proceed to question 21			
20.	20. Is the patient pregnant or breastfeeding?		☐ Yes	□ No		
			STOP	Proceed to question 21		
			Coverage not approved			
21.	Does the patient have a psychiatric comor		☐ Yes	□ No		
	confound a diagnosis of gender dysphoria or interfere with treatment (for example: unresolved body dysmorphic disorder; schizophrenia or other psychotic disorders that have not been stabilized with treatment)?		STOP Coverage not approved	Proceed to question 23		
22.	. Document the requested indication and rationale for use.					
			Proceed to	question 23		
23.	What is the requested medication?			o gel (for example, generic Androgel, generic n and date below		
			62% gel (for example, generic Androgel, etc.) - low			
		☐ testosterone 2% Sign and date bel	solution (for example, generic Axiron, etc.) - ow			
		☐ Other (for example Androderm, Fortesta (2% testosterone gel multi-dose pump (MDP)), Natesto, brand Testosterone 1% gel packet, brand Vogelxo) - Proceed to question 24				
24.	24. Has the patient tried and failed a 3-month trial of one of the following medications: testosterone 1% gel (for example,		□ Yes	□ No		
generic Androgel, generic Testim, etc.), 1.62% gel (for example, generic Androgel, etc.), or 2% solution (for example, generic Axiron, etc.)?		Sign and date below	Proceed to question 25			

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	25. Has the patient experienced a clinically significant adverse reaction to one of the following medications: testosterone 1% gel (for example, generic Androgel, generic Testim, etc.), 1.62% gel (for example, generic Androgel, etc.), or 2% solution (for example, generic Axiron, etc.)?	☐ Yes Sign and date below	☐ No Proceed to question 26
	26. Has the patient had a contraindication or relative contraindication to one of the following medications: testosterone 1% gel (for example, generic Androgel, generic Testim, etc.), 1.62% gel (for example, generic Androgel, etc.), or 2% solution (for example, generic Axiron, etc.)?	☐ Yes Sign and date below	□ No STOP Coverage not approved
Step 3	I certify the above is true to the best of my knowledge. Please sign	and date:	
	Prescriber Signature	Date	
			[29 May 2024]