

Clinical Policy: Topical Testosterone
Reference Number: HIM.PA.87
Effective Date: 12/14
Last Review Date: 08/17
Line of Business: Health Insurance Marketplace

[Coding Implications](#)
[Revision Log](#)

See [Important Reminder](#) at the end of this policy for important regulatory and legal information.

Description

The following are topical testosterone agents requiring prior authorization: testosterone transdermal system (Androderm®)

FDA approved indication

Topical testosterone agents are indicated:

- For the treatment of primary hypogonadism (congenital or acquired)
- For the treatment of hypogonadotropic hypogonadism (congenital or acquired)

Limitation of use:

- Safety and efficacy of Androderm in men with “age-related hypogonadism” have not been established.
- Safety and efficacy of Androderm in males less than 18 years old have not been established.
- Topical testosterone products may have different doses, strengths, or application instructions that may result in different systemic exposure.

Policy/Criteria

Provider must submit documentation (including office chart notes and lab results) supporting that member has met all approval criteria

I. Initial Approval Criteria

A. Hypogonadism (primary and hypogonadotropic types) (must meet all):

1. Diagnosis of hypogonadism;
2. Failure of testosterone cypionate or testosterone enanthate injection at up to maximally indicated dose unless contraindication or clinically significant adverse effects are experienced (*Failure is demonstrated by lower than normal total testosterone levels as compared to laboratory reference values*);
3. Dose does not exceed the FDA approved maximum recommended dose for the relevant drug.

Approval duration: 12 months

B. Other diagnoses/indications

1. Refer to HIM.PHAR.21 if diagnosis is NOT specifically listed under section III (Diagnoses/Indications for which coverage is NOT authorized).

II. Continued Therapy

A. Hypogonadism (primary and hypogonadotropic types) (must meet all):

CLINICAL POLICY

Topical Testosterone

1. Currently receiving medication via Centene benefit or member has previously met initial approval criteria;
2. Documentation of positive response to therapy;
3. Dose does not exceed the FDA approved maximum recommended dose for the relevant drug.

Approval duration: 12 months

B. Other diagnoses/indications (must meet 1 or 2):

1. Currently receiving medication via health plan benefit and documentation supports positive response to therapy.

Approval duration: Duration of request or 12 months (whichever is less); or

2. Refer to HIM.PA.21 if diagnosis is NOT specifically listed under section III (Diagnoses/Indications for which coverage is NOT authorized)

III. Diagnoses/Indications for which coverage is NOT authorized:

- A.** Non-FDA approved indications, which are not addressed in this policy, unless there is sufficient documentation of efficacy and safety according to the off label use policy – HIM.PHAR.21 or evidence of coverage documents

IV. Appendices/General Information

Appendix A: Abbreviation Key

FDA: Food and Drug Administration

V. References

1. Androderm Prescribing Information. Irvine, CA: Allergan USA, Inc.; October 2016. Available at <https://www.allergan.com>. Accessed April 7, 2017.
2. Testosterone Drug Monograph. Clinical Pharmacology. Accessed January 24, 2017. <http://www.clinicalpharmacology-ip.com/default.aspx>.
3. Androgel Prescribing Information. North Chicago, IL: AbbVie Inc.; October 2016. Available at <https://www.androgel.com>. Accessed April 7, 2017.
4. Axiron Prescribing Information. Indianapolis, IN: Lilly USA, LLC; February 2017. Available at <http://www.axiron.com>. Accessed April 7, 2017.
5. Vogelxo Prescribing Information. Maple Grove, MN: Upsher-Smith Laboratories, Inc.; September 2016. Available at <http://www.upsher-smith.com/wp-content/uploads/Vogelxo-MI.pdf>. Accessed April 7, 2017.
6. The World Professional Association for Transgender Health (WPATH), Standards of Care for the Health of Transsexual, Transgender, and Gender Nonconforming People, 7th Version.

Reviews, Revisions, and Approvals	Date	P&T Approval Date
Reformatted guideline to new format. Added Workflow reference document. Removed exclusion for use in gender reassignment transitioning	12/15	12/15
Removed gender restriction to male. Updated references.	11/16	11/16

CLINICAL POLICY

Topical Testosterone

Reviews, Revisions, and Approvals	Date	P&T Approval Date
Converted to new template. Added testosterone enanthate injection as a trial option.	4/17	08/17

Important Reminder

This clinical policy has been developed by appropriately experienced and licensed health care professionals based on a review and consideration of currently available generally accepted standards of medical practice; peer-reviewed medical literature; government agency/program approval status; evidence-based guidelines and positions of leading national health professional organizations; views of physicians practicing in relevant clinical areas affected by this clinical policy; and other available clinical information. The Health Plan makes no representations and accepts no liability with respect to the content of any external information used or relied upon in developing this clinical policy. This clinical policy is consistent with standards of medical practice current at the time that this clinical policy was approved. “Health Plan” means a health plan that has adopted this clinical policy and that is operated or administered, in whole or in part, by Centene Management Company, LLC, or any of such health plan’s affiliates, as applicable.

The purpose of this clinical policy is to provide a guide to medical necessity, which is a component of the guidelines used to assist in making coverage decisions and administering benefits. It does not constitute a contract or guarantee regarding payment or results. Coverage decisions and the administration of benefits are subject to all terms, conditions, exclusions and limitations of the coverage documents (e.g., evidence of coverage, certificate of coverage, policy, contract of insurance, etc.), as well as to state and federal requirements and applicable Health Plan-level administrative policies and procedures.

This clinical policy is effective as of the date determined by the Health Plan. The date of posting may not be the effective date of this clinical policy. This clinical policy may be subject to applicable legal and regulatory requirements relating to provider notification. If there is a discrepancy between the effective date of this clinical policy and any applicable legal or regulatory requirement, the requirements of law and regulation shall govern. The Health Plan retains the right to change, amend or withdraw this clinical policy, and additional clinical policies may be developed and adopted as needed, at any time.

This clinical policy does not constitute medical advice, medical treatment or medical care. It is not intended to dictate to providers how to practice medicine. Providers are expected to exercise professional medical judgment in providing the most appropriate care, and are solely responsible for the medical advice and treatment of members. This clinical policy is not intended to recommend treatment for members. Members should consult with their treating physician in connection with diagnosis and treatment decisions.

Providers referred to in this clinical policy are independent contractors who exercise independent judgment and over whom the Health Plan has no control or right of control. Providers are not agents or employees of the Health Plan.

CLINICAL POLICY

Topical Testosterone

This clinical policy is the property of the Health Plan. Unauthorized copying, use, and distribution of this clinical policy or any information contained herein are strictly prohibited. Providers, members and their representatives are bound to the terms and conditions expressed herein through the terms of their contracts. Where no such contract exists, providers, members and their representatives agree to be bound by such terms and conditions by providing services to members and/or submitting claims for payment for such services.

Note: For Medicaid members, when state Medicaid coverage provisions conflict with the coverage provisions in this clinical policy, state Medicaid coverage provisions take precedence. Please refer to the state Medicaid manual for any coverage provisions pertaining to this clinical policy.

©2016 Centene Corporation. All rights reserved. All materials are exclusively owned by Centene Corporation and are protected by United States copyright law and international copyright law. No part of this publication may be reproduced, copied, modified, distributed, displayed, stored in a retrieval system, transmitted in any form or by any means, or otherwise published without the prior written permission of Centene Corporation. You may not alter or remove any trademark, copyright or other notice contained herein. Centene® and Centene Corporation® are registered trademarks exclusively owned by Centene Corporation.