

# SmartPA Criteria Proposal

<b>Drug/Drug Class:</b>	Oxandrin Clinical Edit
<b>First Implementation Date:</b>	November 25, 2002
<b>Revised Date:</b>	February 18, 2021
<b>Prepared for:</b>	MO HealthNet
<b>Prepared by:</b>	MO HealthNet/Conduent
<b>Criteria Status:</b>	<input checked="" type="checkbox"/> Existing Criteria <input type="checkbox"/> Revision of Existing Criteria <input type="checkbox"/> New Criteria

## Executive Summary

**Purpose:** Ensure appropriate utilization and control of Oxandrin® (oxandrolone)

**Why Issue Selected:** Oxandrin® is an oral tablet formulation of the anabolic steroid oxandrolone. It was FDA approved in 1964 and became a Schedule III controlled substance in 1991. Oxandrin is indicated as adjunctive therapy to promote weight gain after weight loss following extensive surgery, chronic infections, or severe trauma or in patients who fail to gain or maintain normal weight without definite pathophysiologic reasons. It is also indicated to offset the protein catabolism associated with prolonged administration of corticosteroids and for the relief of the bone pain frequently accompanying osteoporosis.

Program-Specific Information:	Date Range FFS 7-1-2019 to 6-30-2020				
	Drug	Claims	Spend	Cost per tab	Avg spend per claim
	OXANDRIN 2.5 MG TABLET	109	\$27,977.14	\$2.79 MAC	\$256.67
	OXANDRIN 10 MG TABLET	8	\$6,199.07	\$13.39 MAC	\$774.88

**Type of Criteria:**  Increased risk of ADE  Preferred Drug List  
 Appropriate Indications  Clinical Edit

**Data Sources:**  Only Administrative Databases  Databases + Prescriber-Supplied

## Setting & Population

- Drug class for review: Oxandrin® (oxandrolone)
- Age range: All appropriate MO HealthNet participants

## Approval Criteria

- Diagnosis of bone pain associated with osteoporosis **OR**
- Diagnosis of protein catabolism associated with chronic corticosteroids **OR**
- To promote weight gain:

*SmartPA Clinical Proposal Form*

© 2021 Conduent Business Services, LLC. All rights reserved. Conduent™ and Conduent Design™ are trademarks of Conduent Business Services, LLC in the United States and/or other countries.

Other company trademarks are also acknowledged.

- Documented history of extensive surgery, chronic infection, or severe trauma **OR**
- Documented failure to gain or maintain at least 90% of ideal body weight due to underlying disease state (ex. COPD or AIDS) **OR**
- Approval based on clinical consultant review

## Denial Criteria

- Therapy will be denied if all approval criteria are not met
- Participant is currently pregnant

## Required Documentation

Laboratory Results:   
 MedWatch Form:

Progress Notes:   
 Other:

## Disposition of Edit

Denial: Exception code “0682” (Clinical Edit)  
 Rule Type: CE

## Default Approval Period

30 days

## References

- Oxandrin® (oxandrolone tablets) [package insert]. East Brunswick, NJ: Savient Pharmaceuticals, Inc.; January 2006.
- Facts & Comparisons. Oxandrolone Oral. Accessed August 5, 2020.
- Clinical Pharmacology. Oxandrolone. Accessed August 5, 2020.