

**Clinical Policy: Colonoscopy Preparation Products** 

Reference Number: HIM.PA.04

Effective Date: 01.01.20 Last Review Date: 11.20 Line of Business: HIM

**Revision Log** 

See <u>Important Reminder</u> at the end of this policy for important regulatory and legal information.

#### **Description**

Colonoscopy preparation products contain a combination of osmotic laxatives, stimulant laxatives, and electrolytes used for cleansing of the colon to allow for imaging during a colonoscopy.

## FDA Approved Indication(s)

MoviPrep<sup>®</sup>, OsmoPrep<sup>®</sup>, and Sutab<sup>®</sup> are indicated for cleansing of the colon as a preparation for colonoscopy in adults.

Clenpiq® and Prepopik® are indicated for cleansing of the colon as a preparation for colonoscopy in adults and pediatric patients ages 9 years and older.

#### Policy/Criteria

Provider must submit documentation (such as office chart notes, lab results or other clinical information) supporting that member has met all approval criteria.

It is the policy of health plans affiliated with Centene Corporation<sup>®</sup> that Clenpiq, MoviPrep, OsmoPrep, Prepopik, and Sutab are **medically necessary** when the following criteria are met:

#### I. Initial Approval Criteria

### A. Colonoscopy Preparation (must meet all):

1. For members 12 years and older, failure of Suprep<sup>®</sup>, unless contraindicated or clinically significant adverse effects are experienced.

**Approval duration: 4 weeks (one colonoscopy preparation)** 

#### B. Other diagnoses/indications

1. Refer to the off-label use policy for the relevant line of business if diagnosis is NOT specifically listed under section III (Diagnoses/Indications for which coverage is NOT authorized): HIM.PHAR.21 for health insurance marketplace.

#### **II. Continued Therapy**

### A. Colonoscopy Preparation (must meet all):

1. Currently receiving medication via Centene benefit or member has previously met initial approval criteria.

**Approval duration: 4 weeks (one colonoscopy preparation)** 



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

- 1. Currently receiving medication via Centene benefit and documentation supports positive response to therapy.
  - Approval duration: Duration of request or 12 months (whichever is less); or
- 2. Refer to the off-label use policy for the relevant line of business if diagnosis is NOT specifically listed under section III (Diagnoses/Indications for which coverage is NOT authorized): HIM.PHAR.21 for health insurance marketplace.

#### 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 for health insurance marketplace or evidence of coverage documents.

#### IV. Appendices/General Information

Appendix A: Abbreviation/Acronym Key FDA: Food and Drug Administration

Appendix B: Therapeutic Alternatives

This table provides a listing of preferred alternative therapy recommended in the approval criteria. The drugs listed here may not be a formulary agent for all relevant lines of business

and may require prior authorization.

Drug Name	Dosing Regimen	Dose Limit/
		<b>Maximum Dose</b>
Suprep®	Split-dose regimen:	Not applicable
(magnesium	Total volume of liquid consumed over the course of	
sulfate,	treatment: 2880 mL (96 oz)	
potassium		
sulfate,	Evening before colonoscopy: Drink the entire contents	
sodium	of 1 bottle, diluted to a final volume of 480 mL (16 oz).	
sulfate)	Then drink 2 additional containers of water each (filled	
	to the 16-ounce line) over the next hour, for an	
	additional volume of 960 mL (32 oz).	
	Morning of the colonoscopy (10-12 hours after the	
	evening dose): Repeat entire process with the second	
	bottle: Drink entire contents of second bottle diluted to	
	a final volume of 480 mL (16 oz); then drink 2	
	additional containers of water (each filled to the 16-	
	ounce line) over the next hour, for an additional	
	volume of 960 mL (32 oz). Complete at least 2 hours	
	before the procedure.	

Therapeutic alternatives are listed as Brand name<sup>®</sup> (generic) when the drug is available by brand name only and generic (Brand name<sup>®</sup>) when the drug is available by both brand and generic.

Appendix C: Contraindications/Boxed Warnings

• Contraindication(s):



- All colonoscopy prep products: gastrointestinal obstruction, ileus (except OsmoPrep), or gastric retention (except OsmoPrep); bowel perforation; toxic colitis or toxic megacolon; hypersensitivity
- o Prepopik: severely reduced renal function (creatinine clearance less than 30 mL/min)
- OsmoPrep: biopsy-proven acute phosphate nephropathy, gastric bypass or stapling surgery
- Boxed warning(s): OsmoPrep acute phosphate nephropathy

## V. Dosage and Administration

Drug Name	Dosing Regimen	<b>Maximum Dose</b>
Clenpiq	Split-dose regimen: 160 mL evening before colonoscopy. Second 160 mL the morning of the	Not applicable
	colonoscopy	
	Day-before regimen: 160 mL during afternoon or	
	early evening before colonoscopy. Second 160 mL 6 hours later during evening before colonoscopy	
MoviPrep	Split dose (2 day regimen) (preferred method):	Not applicable
	Dose 1: Evening before colonoscopy (10-12 hours before dose 2): 240 mL (8 oz) every 15 minutes until	
	1 L (entire contents of container) is consumed. Then	
	fill container with 480 mL (16 oz) of clear liquid and consume prior to going to bed.	
	Dose 2: On the morning of the colonoscopy	
	(beginning at least 3.5 hours prior to procedure): 240 mL (8 oz) every 15 minutes until 1 L (entire contents	
	of container) is consumed. Then fill container with	
	480 mL (16 oz) of clear liquid and consume at least 2 hours before the procedure.	
OsmoPrep	Evening before colonoscopy: Four tablets with 8	Not applicable
	ounces of clear liquids every 15 minutes for a total of 20 tablets	
	Next morning: Four tablets with 8 ounces of clear	
Duan an ila	liquids every 15 minutes for a total of 12 tablets	Not applicable
Prepopik	Adults and pediatrics: <u>Split-dose regimen (preferred):</u> 150 mL (5 oz) the	Not applicable
	evening before the colonoscopy (5 PM-9 PM),	
	followed by a second 150 mL (5 oz) dose ~5 hours before the colonoscopy	
	before the colonoscopy	
	<u>Day-before regimen (alternative):</u> 150 mL (5 oz) in	
	the early evening before the colonoscopy (4 PM-6 PM), followed by a second 150 mL (5 oz) dose 6	



Drug Name	Dosing Regimen	Maximum Dose
	hours later (10 PM-12 AM) the night before the	
	colonoscopy	
Sutab	Split dose (2 day regimen) (preferred method):	Not applicable
	Day 1 (day prior to colonoscopy): Take 12 tablets PO,	
	then drink 16 oz of water over 15-20 minutes. 1 hour	
	after the last tablet is ingested, drink another 16 oz of	
	water over 30 minutes. 30 minutes after the second	
	container of water is finished, drink another 16 oz of	
	water over 30 minutes	
	Day 2 (day of colonoscopy): The morning of	
	colonoscopy (5 to 8 hours prior to the colonoscopy	
	and no sooner than 4 hours from starting Dose 1), take	
	12 tablets PO, then drink 16 oz of water over 15-20	
	minutes. 1 hour after the last tablet is ingested, drink	
	another 16 oz of water over 30 minutes. Complete all	
	tablets and water at least two hours prior to	
	colonoscopy	

### VI. Product Availability

Drug Name	Availability		
Clenpiq	Oral solution: Each bottle contains 10 mg of sodium picosulfate, 3.5 g		
	of magnesium oxide, and 12 g of anhydrous citric acid in 160 mL of		
	solution		
MoviPrep	Oral solution: Pouch A – 100 grams PEG 3350, 7.5 grams sodium		
	sulfate, 2.691 grams sodium chloride, 1.015 grams potassium		
	chloride; Pouch B – 4.7 grams ascorbic acid, 5.9 grams sodium		
	ascorbate		
OsmoPrep	Tablet: 1.5 g of sodium phosphate		
Prepopik	Powder for oral solution: 2 packets each containing 10 mg sodium		
	picosulfate, 3.5 g magnesium oxide, and 12 g anhydrous citric acid		
Sutab	Tablet: 1.479 g sodium sulfate, 0.225 g magnesium sulfate, and 0.188		
	g potassium chloride		

#### VII. References

- 1. Prepopik Prescribing Information. Parsippany, NJ: Ferring Pharmaceuticals Inc.; August 2018. Available at: http://www.prepopik.com. Accessed November 19, 2020.
- 2. Osmoprep Prescribing Information. Bridgewater, NJ: Salix Pharmaceuticals, Inc.; March 2019. Available at: <a href="https://shared.salix.com/shared/pi/osmoprep-pi.pdf">https://shared.salix.com/shared/pi/osmoprep-pi.pdf</a>. Accessed November 19, 2020.
- 3. Micromedex® Healthcare Series [Internet database]. Greenwood Village, Colo: Thomson Healthcare. Updated periodically. Accessed July 7, 2020.
- 4. Clenpiq Prescribing Information. Parsippany, NJ: Ferring Pharmaceuticals Inc.; October 2019. Available at: www.clenpiq.com. Accessed November 19, 2020.



- 5. MoviPrep Prescribing Information. Bridgewater, NJ: Valeant Pharmaceuticals; May 2019. Available at: <a href="https://moviprep.salix.com">https://moviprep.salix.com</a>. Accessed November 19, 2020.
- 6. Sutab Prescribing Information. Holbrook, MA: Braintree Laboratories, Inc.; November 2020. Available at: <a href="https://www.accessdata.fda.gov/drugsatfda\_docs/label/2020/213135s000lbl.pdf">https://www.accessdata.fda.gov/drugsatfda\_docs/label/2020/213135s000lbl.pdf</a>. Accessed November 19, 2020.

Reviews, Revisions, and Approvals		P&T
		Approval Date
Policy created per SDC and prior clinical guidance; adapted from	12.04.19	02.20
CP.CPA.245 for HIM line of business requiring redirection to Suprep		
for members 18 or older.		
4Q 2020 annual review: modified Suprep redirection to require down	07.07.20	11.20
to age 12 per RT4 to address updated prescribing information for		
pediatric extension; modified approval duration to 4 weeks (one		
colonoscopy preparation); references reviewed and updated.		
RT4: added new dosage form, Sutab.	11.19.20	

## **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.



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