SERYNTH HEMORRHOID- lidocaine 4%, phenylephrine hcl 0.25%, hemorrhoid cream Jiangxi Hemei Pharmaceutical Co., Ltd

84010-121

Active Ingredient

Lidocaine 4%

Phenylephrine HCL 0.25%

Purpose

Analgesic (pain relief) Vasoconstrictor

Use

Temporarily shrink hemorrhoidal tissue, relieving pain, soreness, burning, itching, and discomfort associated with hemorrhoids.

Warnings

For external use only, including the skin aroundthe anal canal

Do not use

Do not put this product into the rectum.

When Using

When using this product do not exceed recommended daily dosage unless directed by a doctor, when using finger cots or an applicator, apply externally or the skin of the anal canal only.

Stop Use

Ask a doctor before use if you have allergies to any of the components of this product, difficulty in urination due to enlargement of the prostate gland, diabetes, heart disease, high blood pressure, thyroid disease, or presently taking a prescription for high blood pressure or depression.

Ask Doctor

Stop use and ask a doctor if condition worsens or does not improve within seven days, bleeding occurs, an allergic reaction develops, the symptom being treated does not

subside or if redness, irritation, swelling, pain or other symptoms develop or increase.

Keep Out Of Reach Of Children

If swallowed, seek medical help or contact a Poison Control Center right away.

Directions

Children under 12 years of age consult your pediatrician. Adults: apply to affected area up to 3 times daily. Clean affected area with mild soap and warm water, rinse thoroughly, and then gently dry (patting or blotting) with tissue or soft cloth before use. To use dispensing cap, attach it to tube, lubricate well then gently insert part way into anus and squeeze tube to deliver medication. Thoroughly cleanse dispensing cap after use with mild soap and warm water and rinse thoroughly.

Other information

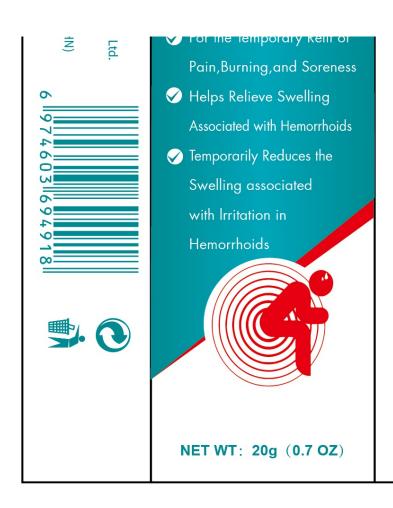
If pregnant or breast-feeding, ask a health care professional before use.

Inactive ingredients

Borneol, Carbomer Homopolymer, Chlorhexidine Acetate Glycerin, Thymol, Aloe

PRINCIPAL DISPLAY PANEL





JRRHOID CREAN

prescription for high blood pressure or depression. **Stop use and ask a doctor if** condition worsens or does not improve within seven days, bleeding occurs, an allergic reaction develops, the symptom being treated does not subside or if redness, irritation, swelling, pain or other symptoms develop or increase.

· KEEP OUT OF REACH OF CHILDREN

If swallowed, seek medical help or contact a Poison Control Center right away.

· DOSAGE & ADMINISTRATION

Children under 12 years of age consult your pediatrician. Adults: apply to affected area up to 3 times daily.Clean affected area with mild soap and warm water, rinse thoroughly, and then gently dry (patting or blotting) with tissue or soft cloth before use.To use dispensing cap, attach it to tube, lubricate well then gently insert part way into anus and squeeze tube to deliver medication. Thoroughly cleanse dispensing cap after use with mild soap and warm water and rinse thoroughly.

· INACTIVE INGREDIENT

Borneol, Carbomer Homopolymer, Chlorhexidine Acetate, Glycerin, Thymol, Aloe

· PREGNANCY OR BREAST FEEDING

If pregnant or breast-feeding, ask a health care professional before use.

SERYNTH HEMORRHOID

lidocaine 4%, phenylephrine hcl 0.25%, hemorrhoid cream

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:84010-121

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	4 g in 100 g
,	PHENYLEPHRINE HYDROCHLORIDE	0.25 g in 100 g

Inactive	Ingredients
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Ingredient Name

Strength

ALOE (UNII: V5VD430YW9)	
GLYCERIN (UNII: PDC6A3C0OX)	
THYMOL (UNII: 3J50XA376E)	
CHLORHEXIDINE ACETATE (UNII: 5908ZUF22Y)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
BORNEOL (UNII: M89NIB437X)	

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:84010-121- 01	20 g in 1 BOTTLE; Type 0: Not a Combination Product	05/23/2025	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	05/23/2025	

Labeler - Jiangxi Hemei Pharmaceutical Co., Ltd (724892056)

Establishment				
Name	Address	ID/FEI	Business Operations	
Jiangxi Hemei Pharmaceutical Co., Ltd		724892056	manufacture(84010-121)	

Revised: 5/2025 Jiangxi Hemei Pharmaceutical Co., Ltd