CHLORASEPTIC SORE THROAT MAX- phenol and glycerin spray
Prestige Brands Holdings, Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Chloraseptic Sore Throat Spray Max

Drug Facts

Active Ingredients

Phenol 1.5%

Purpose

Oral Anesthetic/Analgesic

Active Ingredients

Glycerin 33%

Purpose

Demulcent

Uses

for the temporary relief of ■ occasional minor irritation, pain, sore mouth and sore throat ■ minor discomfort and protection of irritated areas in sore mouth and sore throat

Warnings

Sore Throat Warning: Severe or persistent sore throat or sore throat accompanied by high fever, headache, nausea, and vomiting may be serious. Consult a doctor promptly. Do not use more than 2 days or administer to children under 3 years of age unless directed by a doctor.

When using this product

do not exceed recommended dosage.

If pregnant or breast-feeding,

ask a doctor before use.

Keep out of reach of children.

In case of overdose, or accidental poisoning, get medical help or contact a Poison Control Center right away.

Directions

Adults and children 3 years of age and older:

- apply to the affected area (one spray)
- allow to remain in place for at least 15 seconds, then spit out
- use every 2 hours or as directed by a doctor or dentist

Children under 12 years of age should be supervised by an adult in the use of this product.

Children under 3 years of age: consult a doctor or dentist.

Other Information

- store at room temperature
- check expiration date before using

Inactive ingredients

FD&C blue no. 1, FD&C red no. 40, flavor, purified water, sodium saccharin, sucralose

Questions?

1-800-552-7932 Chloraseptic.com

PRINCIPAL DISPLAY PANEL

Chloraseptic® Max Sore Throat

Phenol Oral Anesthetic /Demulcent Spray 1 fl oz (30 mL) | Wild Berries

CHLORASEPTIC SORE THROAT MAX

phenol and glycerin spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67172-937
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
PHENOL (UNII: 339NCG44TV) (PHENOL - UNII:339NCG44TV)	PHENOL	15 mg in 1 mL
GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	GLYCERIN	330 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	

WATER (UNII: 059QF0KO0R)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	

Product Characteristics			
Color	PURPLE	**Score**	
Shape		**Size**	
Flavor	BERRY	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67172-937-52	1 in 1 BOX	02/01/2011	
1		30 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product		
2	NDC:67172-937-04	1 in 1 BOX	06/01/2021	
2		118 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part356	02/01/2011	

Labeler - Prestige Brands Holdings, Inc. (159655021)

Establishment

Name	Address	ID/FEI	Business Operations
Denison Pharmaceuticals, LLC		001207208	MANUFACTURE(67172-937)

Revised: 1/2021 Prestige Brands Holdings, Inc.