VICKS VAPOCOOL SEVERE SUGAR-FREE- menthol lozenge
The Procter & Gamble Manufacturing Company

Vicks ®VapoCOOL ™SEVERE MENTHOL • ORAL ANESTHETIC
SUGAR FREE WINTERFROST MEDICATED DROPS

Drug Facts

Active ingredient (per drop)

Menthol 20 mg

Purpose

Oral anesthetic

Uses

temporarily relieves occasional minor irritation and pain due to
- sore throat
- sore mouth

Warnings

Sore Throat Warning-

Severe or persistent sore throat or sore throat accompanied by high fever, headache, nausea, and vomiting may be serious. Consult doctor promptly. Do not use more than 2 days or administer to children under 12 years of age.

Ask a doctor before use if you have
- a severe sore throat accompanied by difficulty in breathing or that lasts more than 2 days
- a sore throat accompanied by fever, headache, rash, swelling, nausea or vomiting

Stop use and ask a doctor if
- sore mouth symptoms do not improve in 7 days, or if irritation, pain, or redness persists or worsens.

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

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

Directions

- adults and children 12 years and over: dissolve 1 drop slowly in the mouth. Repeat every 2 hours as needed or as directed by a doctor.
- children 12 years and under: do not use

Other information

- Store at no greater than 25°C (77°F)
- contains Soy.

Inactive ingredients

acacia, acesulfame potassium, ascorbyl palmitate, beta-carotene, FD&C Blue No. 1, flavor, glycerin, hydrogenated starch hydrolysate, isomalt, maltodextrin, potassium sorbate, sucralose, sucrose, sunflower oil, vitamin E DL-alpha, water

Questions?

1-800-707-1709

DIST. BY:

PROCTER & GAMBLE

CINCINNATI, OH 45202

SUGAR FREE

VICKS ®

VapoCOOL™

SEVERE

MENTHOL - ORAL ANESTHETIC

SUGAR FREE

MAX STRENGTH - FAST RELIEF

WINTERFROST

30 MEDICATED DROPS

VICKS VAPOCOOL SEVERE SUGAR-FREE

menthol lozenge

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:84126-313
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	20 mg

Inactive Ingredients	
Ingredient Name	**Strength**
SUCROSE (UNII: C151H8M554)	
ACESULFAME POTASSIUM (UNII: 23OV73Q5G9)	
HYDROGENATED STARCH HYDROLYSATE (UNII: 27F77DSJ5V)	
ISOMALT (UNII: S870P55O2W)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
ASCORBYL PALMITATE (UNII: QN83US2B0N)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	

SUNFLOWER OIL (UNII: 3W1JG795YI)	
BETA CAROTENE (UNII: 01YAE03M7J)	
ACACIA (UNII: 5C5403N26O)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
GLYCERIN (UNII: PDC6A3C0OX)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
WATER (UNII: 059QF0KO0R)	
.ALPHA.-TOCOPHEROL, DL- (UNII: 7QWA1RIO01)	

Product Characteristics

Color	white (with blue specks)	**Score**	no score
Shape	OVAL	**Size**	23mm
Flavor	MENTHOL	**Imprint Code**	vapor
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84126-313-30	30 in 1 BAG	06/28/2024	
1		1 in 1 POUCH; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M022	06/28/2024	

Labeler - The Procter & Gamble Manufacturing Company (004238200)

Revised: 9/2024 The Procter & Gamble Manufacturing Company