SEVERE THROAT SORE DROPS- menthol pastille
Target Corporation 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.

Severe Throat Drops Oral Anesthetic

Active Ingredient in each drop Purposes

Menthol 20mg Cough Suppressant

Oral anesthetic

Temporarily relieves ocassional minor irritation and pain due to:
- sore throat
- sore mouth

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Warnings

Sore throat warning: severe or persistence sore throat accompanied or followed 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 12 years of age.

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

Stop use and consult a doctor if
- sore mouth symptons do not improve in 7 days or if irritation, pains or redness persists or worsens.

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

Keep this and all drugs out of the reach of children.

In case of accidental 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 mouth. Repeated every 2 hours as needed or as directed by a doctor.
- Children under 12 years and under do not use

- Store in a cool and dry place

eucalyptus oil, FD&C Blue No.1, glucose syrup, gum arabic, sucrose

Drug Facts

Active ingredient (in each drop)	Purposes
Menthol 20 mg	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 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 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.

When using this product
- do not exceed recommended dosage.

Stop use and consult 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 the 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 under 12 years and under:	do not use.

Other information
- Store in a cool and dry place.

Inactive ingredients: Eucalytus oil, FD&C blue no. 1, glucose syrup, gum arabic, sucrose.

Questions? Or to report and adverse event call **1-617-848-4560**. Monday - Friday, 9AM to 4PM EST.

100% satisfaction guaranteed or your money back.

Distributed by Target Corporation
Minneapolis, MN 55403
Made in Spain

*This product is not manufactured or distributed by Procter & Gamble, owner of the registered trademark Vicks Vapocool™ severe.

PAPER BOX
how2recycle.info

094 04 TBD 1 R00
C-001227-01-006

80% UPC
FOR POSITION
ONLY

0 00000 00000 0

Leave blank for
LOT / EXP. DATE

Questions? or to report and adverse event call: 1-617-848-4560

Monday to Friday 9AM to 4 PM east

Do not use if bag is torn or open

reclosable bag

NDC 00000-000-00

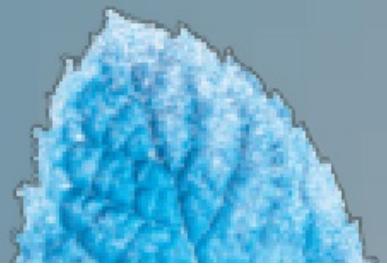
MAX strength

Compare to active ingredient in
Vicks® Severe VapoCool®*

severe
throat drops
oral anesthetic

helps soothe severe sore throat pain

maximum strength menthol for greater cooling effect



SEVERE THROAT SORE DROPS

menthol pastille

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-143
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	0.69 mg

Inactive Ingredients

Ingredient Name	Strength
EUCALYPTUS OIL (UNII: 2R04ONI662)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
WATER (UNII: 059QF0KO0R)	
ACACIA (UNII: 5C5403N26O)	
SUCROSE (UNII: C151H8M554)	
CORN SYRUP (UNII: 9G5L16BK6N)	

Product Characteristics

Color	white (with menthol drops)	**Score**	score with uneven pieces
Shape	RECTANGLE	**Size**	17mm
Flavor	MENTHOL	**Imprint Code**	None
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-143-01	45 in 1 BAG; Type 0: Not a Combination Product	10/12/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	10/12/2019	

Labeler - Target Corporation Inc. (006961700)

Registrant - Boston Nutraceutical Science SL (466061824)

Establishment

Name	Address	ID/FEI	Business Operations
Boston Nutraceutical Production SL		468121064	manufacture(11673-143) , label(11673-143) , pack(11673-143)

Revised: 1/2023 Target Corporation Inc.