

MENTHOL COUGH 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.

Menthol Cough Drops

Active ingredient 5.8 mg

Cough Suppressant

Oral anesthetic

Do not use if bag
is torn or open



◀ Tear here

Resealable Bag

Tear here ▶

NDC 82442-001-01

Compare to active ingredient in Halls®*

menthol cough drops

cough suppressant
oral anesthetic

helps soothe sore throat pain
temporarily relieves cough



80 DROPS

**Actual size depicted on back.



Uses temporarily relieves:

- cough as may occur with a cold
- occasional minor irritation and sore throat

Uses temporarily relieves:

cough as may occur with a cold
occasional minor irritation and sore throat

Warnings

Sore throat warning: severe or persistence 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 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 symptoms 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

blackcurrant and black carrot extract (coloring), citric acid, flavors, glucose syrup, sucrose

Leave blank for
LOT / EXP. DATE



Drug Facts	
Active ingredient (in each drop) Purposes Menthol 5.8 mg cough suppressant, oral anesthetic	
Uses temporarily relieves: cough as may occur with a cold. occasional minor irritation and sore throat.	
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 and consult 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 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: do not use.	
Other information 10 calories per drop. Store in a cool and dry place.	
Inactive ingredients Blackcurrant and blackcarrot extract (coloring), citric acid, flavors, glucose syrup, sucrose.	
Questions? Or to report and adverse event call 1-800-910-6874.	

100% satisfaction guaranteed or your money back.

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Mad in Spain
This product is not manufactured or distributed by Mondelez Global LLC, owner of the registered trademark Halls®



menthol cough drops
cough suppressant
oral anesthetic

no artificial flavors
colors from natural sources
no artificial preservatives



Resealable Bag

▶ Tear here

▶ Tear here



Questions? or to report and adverse event call: 1-800-423-0139

Drug Facts	
Active ingredient (in each drop) Purposes Menthol 5.8 mg cough suppressant, oral anesthetic	
Uses temporarily relieves: ■ cough as may occur with a cold. ■ occasional minor irritation and sore throat.	
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 and consult 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 the reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.	
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**Actual Size

menthol cough drops
cough suppressant
oral anesthetic
no artificial flavors
colors from natural sources
no artificial preservatives

Resealable Bag

▶ Tear here

▶ Tear here



MENTHOL COUGH DROPS

menthol pastille

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82442-001
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	5.8 mg

Inactive Ingredients

Ingredient Name	Strength
EUCALYPTOL (UNII: RV6J6604TK)	
BLACK CURRANT (UNII: 9755T40D11)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
BLACK CARROT (UNII: 27793ZFD2L)	
CHERRY (UNII: BUC5I9595W)	
SUCROSE (UNII: C151H8M554)	
CORN SYRUP (UNII: 9G5L16BK6N)	

Product Characteristics

Color	red	Score	score with uneven pieces
Shape	OVAL	Size	25mm
Flavor	CHERRY	Imprint Code	None
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82442-001-02	80 in 1 BAG; Type 0: Not a Combination Product	10/24/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	10/18/2022	

Labeler - Target Corporation Inc. (006961700)**Registrant** - Boston Nutraceutical Science SL (466061824)**Establishment**

Name	Address	ID/FEI	Business Operations
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Boston Nutraceutical Production SL		468121064	manufacture(82442-001) , label(82442-001) , pack(82442-001)
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Revised: 10/2022

Target Corporation Inc.