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



Uses temporarily relieves:

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

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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 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

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





Questions? Or to report and adverse event call 1-888-423-0139.

extract (coloring), citric acid, flavors, glucose syrup, sucrose Inactive ingredients Blackcurrant and blackcarrot

> Store in a cool and dry place. Other information = 10 calones per drop.

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St nenblida and children 12 Directions

Center right away. overdose, get medical help or contact a Poison Control before use. Keep out of the reach of children. In case of If pregnant or breast-feeding, ask a health professional

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> do not exceed recommended dosage. When using this product

> > swelling, nausea or vomiting.

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Warnings

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

USes temporarily relieves:

Active ingredient (in each drop) Menthol 5.8 mg cough suppressant, oral anesthetic Purposes

Drug Facts

owner of the registered trademark Halls®. distributed by Mondelez International, *This product is not manufactured or

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100% satisfaction guaranteed or your money back.



no artificial preservatives colors from natural sources no artificial flavors

oral anesthetic condy suppressant menthol cough drops

Resealable Bag



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





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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			

ı	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:82442-001- 01	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

Boston Nutraceutical Production	468121064	manufacture(82442-001) , label(82442-001) , pack(82442-001)
JL		001)

Revised: 10/2022 Target Corporation Inc.