

SUGAR FREE MENTHOL COUGH DROPS- menthol lozenge
Topco Associates

Menthol 5.8mg

Active ingredient (in each drop)

Menthol 5.8 mg

Purpose

Cough suppressant / Oral anesthetic

Uses temporarily relieves:

cough as may occur with a cold

occasional minor irritation and sore throat

Warnings

Sore throat warning: if sore throat is severe, persists for more than 2 days, is accompanied or followed by fever, headache, rash, swelling, nausea, or vomiting, consult a doctor promptly. If sore mouth symptoms do not improve in 7 days, see your dentist or doctor promptly. These symptoms may be serious.

Ask a doctor before use if you have:

persistent or chronic cough such as occurs with smoking, asthma, or emphysema

cough accompanied by excessive phlegm (mucus)

Stop use and ask doctor if

cough persists for more than 7 days, tends to recur, or is accompanied by fever, rash, or persistent headache. These could be signs of a serious condition.

sore throat is severe, or irritation, pain or redness lasts or worsens

sore mouth does not improve in 7 days

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

Keep out of reach of children.

Directions

adults and children 5 years and over - dissolve 1 lozenge slowly in mouth. Repeat every 2 hours as needed or as directed by a doctor.

children under 5 years - ask a doctor

Other information

- 6 calories per drop
- Excess consumption may cause a laxative effect.

Inactive ingredients: Acesulfame potassium, corn starch, eucalyptus oil, FD&C blue no. 1, FD&C red no. 40, glycerin, gum arabic, isomalt, maltitol syrup, medium chain triglycerides, natural and artificial flavors, potassium sorbate, soybean oil, sucralose and

water.

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



RESEALABLE BAG

← Tear here — — — — — Tear here →

Do not use if bag is torn or open

SUGAR FREE

COMPARE TO HALLS®
ACTIVE INGREDIENT*

Cough Drops

MENTHOL COUGH SUPPRESSANT/
ORAL ANESTHETIC

- **Soothes Sore Throat Pain**
- **Temporarily Relieves Cough**



MENTHOL
FLAVOR

25 DROPS



This product is not manufactured or distributed by the owner of the registered trademark HALLS™.

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Directions	adults and children	dissolve 1 lozenge slowly in mouth. Repeat every 2 hours as needed or as directed by a doctor.
	children under 5 years	ask a doctor

If pregnant or breast-feeding, ask a health professional before use. Keep out of reach of children.

■ sore mouth does not improve in 7 days or worsens
■ sore throat is severe, or irritation, pain or redness lasts. These could be signs of a serious condition.
■ cough persists for more than 7 days, tends to recur, or is accompanied by fever, rash, or persistent headache.
Stop use and ask doctor if
■ cough persists for more than 7 days, tends to recur, or is accompanied by excessive phlegm (mucus)
■ persistent or chronic cough such as occurs with smoking, asthma, or emphysema

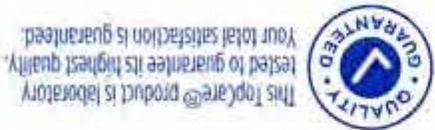
symptoms may be serious.

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QUESTIONS? 1-888-423-0139
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MADE IN USA OF U.S. AND
IMPORTED INGREDIENTS.

MENTHOL COUGH SUPPRESSANT/
ORAL ANESTHETIC

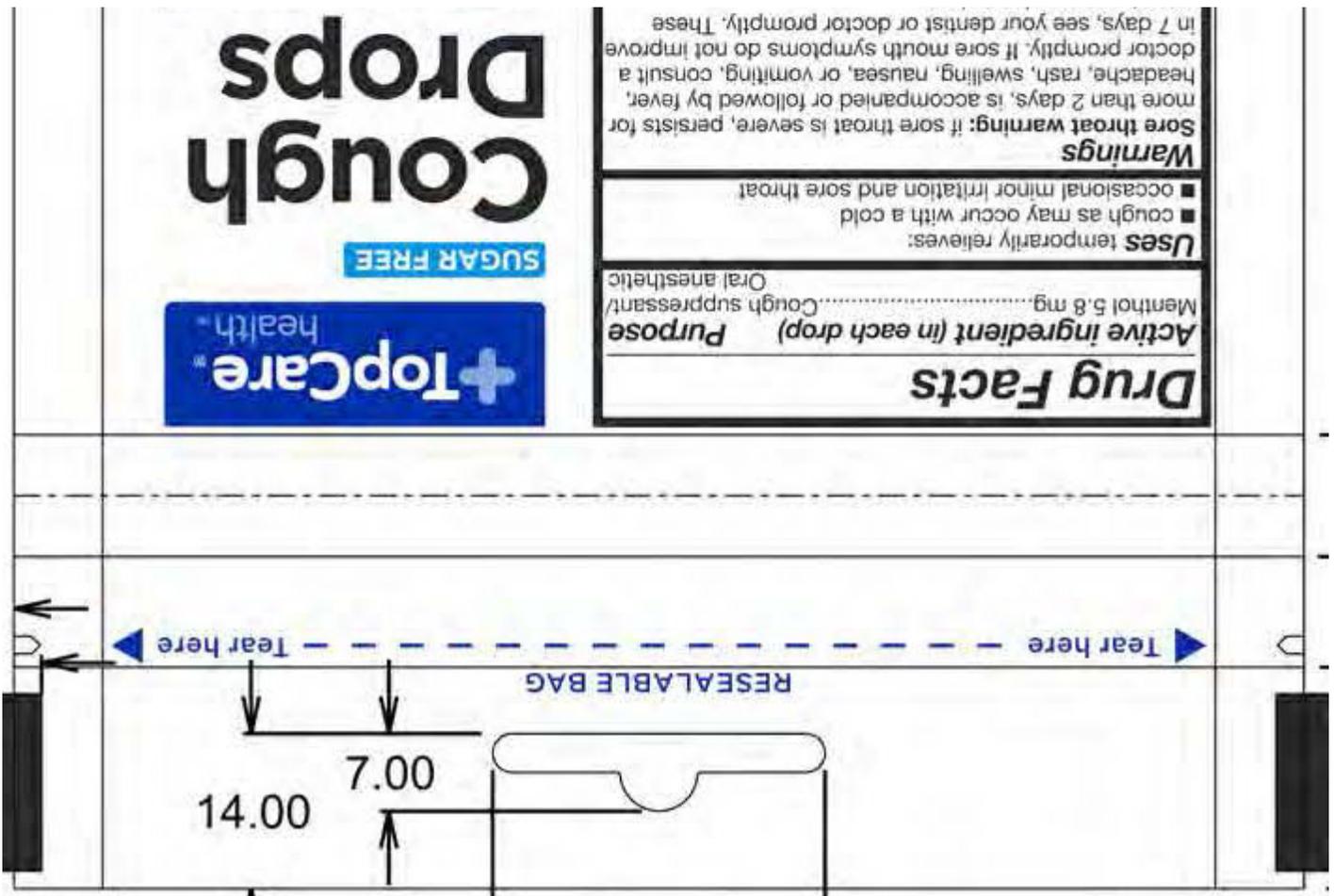
- Soothes Sore Throat Pain
- Temporarily Relieves Cough

MENTHOL FLAVOR



This Topcare® product is laboratory tested to guarantee its highest quality. Your total satisfaction is guaranteed.





SUGAR FREE MENTHOL COUGH DROPS			
menthol lozenge			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:36800-819
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	
WATER (UNII: 059QF0KO0R)			
Product Characteristics			
Color	blue (light blue with blue flakes)	Score	no score
Shape	OVAL	Size	16mm
Flavor	MENTHOL	Imprint Code	B

Contains**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:36800-819-25	25 in 1 BAG; Type 0: Not a Combination Product	03/22/2013	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	03/22/2013	

Labeler - Topco Associates (006935977)**Registrant** - Bestco Inc. (002149136)

Establishment

Name	Address	ID/FEI	Business Operations
Bestco Inc.		002149136	manufacture(36800-819)

Revised: 12/2025

Topco Associates