

**BERRY MEDLEY THROAT DROPS- menthol lozenge**  
**Ricola USA Inc.**

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**BERRY MEDLEY THROAT DROPS**

**Drug Facts**

**Purpose**

Oral anesthetic

**Active Ingredient (in each drop)**

Menthol, 1.7 mg

**Uses**

temporarily relieves occasional minor irritation and pain associated with:

- sore mouth
- sore throat

**Warnings**

**Do not use**

- in children under 6 years of age unless directed by a doctor.

**Stop use and ask a doctor if**

- sore throat is severe, persists for more than 2 days, or is accompanied by fever, headache, rash, nausea or vomiting.
- sore mouth symptoms do not improve in 7 days.

**Keep out of reach of children.**

**Directions**

- adults and children 6 years and older: dissolve 2 drops (one at a time) slowly in the mouth. Do not bite or chew. Repeat every 2 hours as needed or as directed by a doctor
- children under 6 years: ask a doctor

**Other Information**

protect from heat and moisture

Inactive Ingredients

bilberry juice concentrate, black currant juice concentrate, citric acid, extract of Ricola herb mixture (lemon balm, peppermint, thyme, hyssop, sage, elder, linden, mallow, horehound, wild thyme), glucose syrup, natural color, natural flavors, peppermint oil, raspberry juice concentrate, sugar



BERRY MEDLEY THROAT DROPS

menthol lozenge

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63667-436
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name		Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)		1.7 mg

Inactive Ingredients	
Ingredient Name	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
SUCROSE (UNII: C151H8M554)	

Product Characteristics		
Color	yellow (GOLDEN YELLOW)	Score
		no score

Shape	OVAL	Size	24mm	
Flavor	LEMON (LEMON MINT)	Imprint Code	R	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63667-436-45	45 in 1 BAG; Type 0: Not a Combination Product	05/17/2021	
2	NDC:63667-436-19	19 in 1 BAG; Type 0: Not a Combination Product	05/17/2021	
3	NDC:63667-436-10	10 in 1 BAG; Type 0: Not a Combination Product	05/17/2021	
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug		M022	05/17/2021	

**Labeler -** Ricola USA Inc. (177265261)

## Establishment

Name	Address	ID/FEI	Business Operations
Ricola Ag		480227248	manufacture(63667-436)

Revised: 11/2023

Ricola USA Inc.