

BERRY MEDLEY THROAT DROPS- menthol lozenge
Ricola USA Inc.

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	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	1.7 mg

Inactive Ingredients

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

Product Characteristics

Color	yellow (GOLDEN YELLOW)	Score	no score
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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: 12/2024

Ricola USA Inc.