NATURAL CHERRY HONEY HERB THROAT DROPS- menthol lozenge
Ricola Ag

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.

NATURAL CHERRY HONEY HERB THROAT DROPS

Drug Facts

Purpose

Oral anesthetic

Active Ingredient (in each drop)

Menthol, 2.1 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

extract of a Ricola herb mixture (elder, horehound, hyssop, lemon balm, linden flowers, mallow, peppermint, sage, thyme, wild thyme), honey, malic acid, natural cherry concentrate, natural color, natural flavors, starch syrup, sugar

NATURAL CHERRY HONEY HERB THROAT DROPS

menthol lozenge

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:54305-517
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	2.1 mg

Inactive Ingredients

Ingredient Name	Strength
MALIC ACID (UNII: 817L1N4CKP)	
HONEY (UNII: Y9H1V576FH)	
CHERRY (UNII: BUC5I9595W)	
SUCROSE (UNII: C151H8M554)	

Product Characteristics

Color	red	**Score**	no score
Shape	RECTANGLE	**Size**	19mm
Flavor	CHERRY (CHERRY HONEY)	**Imprint Code**	R
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54305-517-09	9 in 1 PACKAGE; Type 0: Not a Combination Product	01/01/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part356	01/01/2022	

Labeler - Ricola Ag (480227248)

Establishment

Name	Address	ID/FEI	Business Operations
Ricola Ag		480227248	manufacture(54305-517)

Revised: 1/2023 Ricola Ag