EXTRA STRENGTH WITH NATURAL MENTHOL GLACIER MINT COUGH SUPPRESSANT - menthol lozenge
Ricola USA Inc.

EXTRA STRENGTH WITH NATURAL MENTHOL GLACIER MINT

Drug Facts

Purpose

Cough suppressant

Active Ingredient (in each drop)

Menthol, 8.3 mg

Uses

temporarily relieves:

- cough due to minor throat and bronchial irritation occurring with a cold or inhaled irritants

Warnings

Ask doctor before use if you have

- persistent chronic cough such as occurs with smoking, asthma, or emphysema
- cough accompanied by excessive phlegm (mucus)

Stop use and ask a doctor if

- cough persists for more than 1 week, tends to recur, or is accompanied by fever, rash, or persistent headache. These could be signs of a serious condition

Keep out of reach of children.

Directions

- adults and children 6 years and older: dissolve 1 drop slowly in the mouth. Repeat every 2 hours as needed or as directed by a doctor
- children under 6 years: ask a doctor

Other Information

Store in a dry place

Inactive Ingredients

extracts of peppermint and a Ricola herb mixture (elder, horehound, hyssop, lemon balm, linden flowers, mallow, peppermint, sage, thyme, wild thyme), glycerin, invert sugar (fructose/dextrose), natural flavor, starch syrup, sugar

EXTRA STRENGTH WITH NATURAL MENTHOL GLACIER MINT COUGH SUPPRESSANT

menthol lozenge

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63667-503
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	8.3 mg

Inactive Ingredients

Ingredient Name	Strength
SUCROSE (UNII: C151H8M554)	
GLYCERIN (UNII: PDC6A3C0OX)	
INVERT SUGAR (UNII: ED959S6ACY)	
PEPPERMINT (UNII: V95R5KMY2B)	

Product Characteristics

Color	green	**Score**	no score
Shape	OVAL	**Size**	24mm
Flavor	MINT	**Imprint Code**	R
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63667-503-19	19 in 1 BAG; Type 0: Not a Combination Product	05/27/2013	
2	NDC:63667-503-26	26 in 1 BAG; Type 0: Not a Combination Product	05/27/2013	
3	NDC:63667-503-04	4 in 1 PACKAGE; Type 0: Not a Combination Product	05/27/2013	01/01/2016

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	05/27/2013	

Labeler - Ricola USA Inc. (177265261)

Establishment

Name	Address	ID/FEI	Business Operations
Ricola Ag		485393768	manufacture(63667-503)

Revised: 12/2024 Ricola USA Inc.