EXTRA STRENGTH WITH NATURAL MENTHOL GLACIER MINT COUGH SUPPRESSANT - menthol lozenge Ricola Ag

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

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	e (Source)	NDC:54	305-617
Route of Administration	ORAL				
Active Ingredient/Active	Moiety				
Active Higheric/Active Profety					
Ingredient Name			Basis of Stren	gth	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:54305-617- 19	19 in 1 BAG; Type 0: Not a Combination Product	05/27/2013	
2	NDC:54305-617- 26	26 in 1 BAG; Type 0: Not a Combination Product	05/27/2013	
3	NDC:54305-617- 04	4 in 1 PACKAGE; Type 0: Not a Combination Product	05/27/2013	01/01/2015

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

Labeler - Ricola Ag (480227248)

Establishment				
Name	Address	ID/FEI	Business Operations	
Ricola Ag		485393768	manufacture(54305-617)	

Revised: 12/2024 Ricola Ag