

NICE CHERRY COUGH DROPS- menthol lozenge

Leosons International

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.

Nice Cherry Cough Drops (Best)

Drug Facts

Active Ingredient & Purpose

Active Ingredient (in each lozenge)

Menthol 5.0 mg

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Menthol 5.0 mg.....

Purposes

Cough Suppressant / Oral Anesthetic

Uses

temporarily suppresses/relieves:

- cough due to minor throat and bronchial irritation from a cold or inhaled irritants
- occasional minor sore throat pain

Warnings

Ask a doctor before use if you have

- chronic cough that lasts or occurs with smoking, asthma, chronic bronchitis or emphysema
- cough that occurs with too much phlegm (mucus)

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When using this product

- do not use more than directed, taking more may have a laxative effect

Stop use and ask a doctor if

- cough lasts more than 7 days, comes back, or occurs with fever, rash or headache that lasts. These could be signs of a serious condition
- sore throat is severe, lasts for more than 2 days, occurs with or is followed by fever, headache, rash, swelling, nausea or vomiting

- sore mouth systems last more than 7 days, or irritation, pain or redness continues or worsens

Keep out of reach of children.

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Directions

- dissolve a lozenge slowly in mouth
- may be repeated every hour for coughs as needed or as directed by a doctor
- may be repeated every 2 hours for sore throat
- do not take more than 10 lozenges per day

adults and children 6 years and older

5 mg

children under 6 years

ask a doctor

Inactive Ingredients

Acesulfame Potassium, Eucalyptus Oil, FD&C Blue No. 1, FD&C Red No. 40, Isomalt, Natural and Artificial Cherry Flavor, Other Natural and Artificial Flavors, Tartaric Acid

Questions?

Call 1-855-452-9500 or email at info@leosonsintl.com

Packaging



NICE CHERRY COUGH DROPS

menthol lozenge

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69626-0042
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	5 mg

Inactive Ingredients

Ingredient Name	Strength
ISOMALT (UNII: S870P5502W)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
TARTARIC ACID (UNII: W4888119H)	

ACESULFAME POTASSIUM (UNII: 23OV73Q5G9)

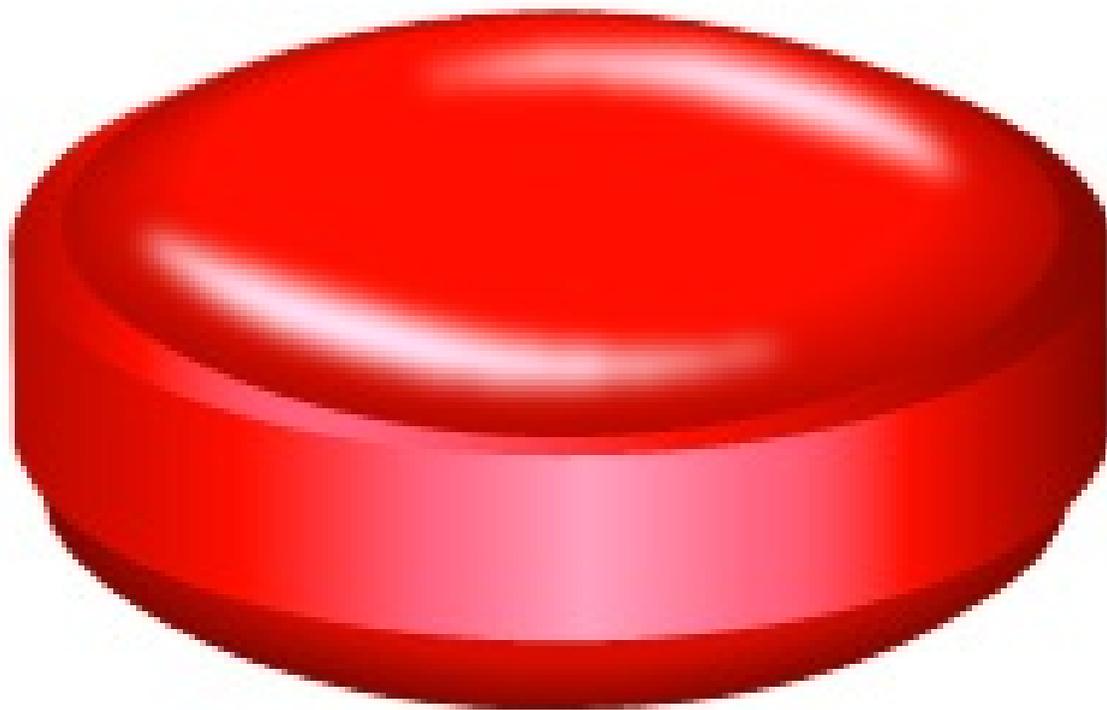
EUCALYPTUS OIL (UNII: 2R04ONI662)

Product Characteristics

Color	red	Score	no score
Shape	ROUND	Size	17mm
Flavor	CHERRY	Imprint Code	N
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69626-0042-4	24 in 1 BLISTER PACK; Type 0: Not a Combination Product	11/08/2019	



Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	11/08/2019	

