EARWAX REMOVAL AID- earwax removal aid liquid McKesson

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.

Earwax Removal Aid

Active ingredient(s)

Carbamide Peroxide 6.5%

Purpose

Earwax Removal Aid

Use(s)

For occasional use to soften, loosen and remove excessive earwax.

Warnings

Warning

For External Use Only

Do not use

- In the eye
- for more than 4 days

Ask a doctor before use if you have

- ear drainage or discharge
- pain, irritation or rash in the ear
- had ear surgery
- an injury or perforation (hole) in the ear drum

Stop use and ask a doctor if

Stop use and ask a doctor if excessive earwax remains after use

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

FOR USE IN THE EAR ONLY

Adults and children 12 years of age: tilt head sideways and place 5 to 10 drops into ear. Tip of applicator should not enter ear canal. Keep drops in ear for several minutes by keeping head tilted, or placing cotton in the ear. Use twice daily for up to 4 days, if needed, or as directed by a doctor. Any wax remaining after treatment my be removed by gently flushing the ear with warm water, using a soft rubber bulb ear syringe. Children under 12 years of age, consult a doctor.

Other information

Do not use if imprinted seal on cap is broken or missing.

Storage

- Store at controlled room temperature 15 to 30C (59 to 86F)
- KEEP IN A DRY PLACE

Principal Display Panel

Principle display panel:



Inactive Ingredients

anhydrous Glycerin

EARWAX REMOVAL AID

earwax removal aid liquid

Product Information

Product Type

Active Ingredient/Active Moiety					
Ingredient Name			Basis	of Strength	Strength
CARBAMIDE PERO XIDE (UNII: 31PZ2VAU81) (CARBAMIDE PERO XIDE - UNII: 31PZ2VAU81)			CARBA PEROXI		6.5 mg in 100 mL
Inactive Ingredients					
Ingredient Name			Strength		
GLYCERIN (UNII: PDC6A3C0OX)			120.0 mg in 100 mL		
Packaging					
# Item Code	Package Description	Marketing Start Date		Marketing End Date	
1 NDC:62011-0167-1	15 mL in 1 BOTTLE				
Marketing Information					
Marketing Category	Application Number or Monograph Citation		Marketing Sta	rt Date Ma	rketing End Date
OTC MONOGRAPH FINAL	part344		02/06/2012		

Labeler - McKesson (177667227)

Registrant - Continental Manufacturing Chemist, Inc. (005278007)

Revised: 5/2012

McKesson