GOOD NEIGHBOR PHARMACY EARWAX REMOVAL DROPS- carbamide peroxide liquid

Preferred Pharmaceuticals Inc.

Good Neighbor Pharmacy Earwax Removal Drops

Drug Facts

Active ingredient

Carbamide peroxide 6.5%

Purpose

Earwax removal aid

Uses

for occasional use as an aid to soften, loosen, and remove excessive earwax

Warnings

For use in the ear only

Ask a doctor beofre use if you have

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

When using this product

avoid contact with the eyes

Stop use and ask a doctor if

- you need to use for more than 4 days
- excessive earwax remains after use of this product

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 over 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 may be removed by gently flushing the ear with warm water, using a soft rubber bulb ear syringe
- consult a doctor Children under 12 years:

Other information

- Avoid exposing bottle to excessive heat and direct sunlight
- Store bottle in the outer carton
- Keep cap on bottle when not in use
- Product foams on contact with earwax due to release of oxygen. There may be an associated crackling sound.
- Do not use if tamper-evident safety seal is broken or missing

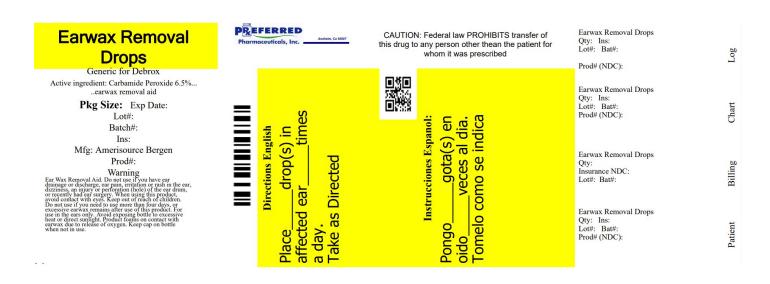
Inactive ingredients

anhydrous citric acid, glycerin, propylene glycol, sodium citrate, sodium lauryl sulfate, water

Questions?

1-800-328-5890

Package Labeling:



GOOD NEIGHBOR PHARMACY EARWAX REMOVAL DROPS

carbamide peroxide liquid

Product I	nformatio	n
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Product Type HUMAN OTC DRUG Item Code (Source) NDC:68788-8586(NDC:46122-557)

Route of Administration TOPICAL

Active Ingredient/Active Moiety Ingredient Name Basis of Strength CARBAMIDE PEROXIDE (UNII: 31PZ 2VAU81) (HYDROGEN PEROXIDE - UNII: BBX060AN9V) CARBAMIDE PEROXIDE (UNII: 31PZ 2VAU81) (HYDROGEN PEROXIDE - UNII: BBX060AN9V) CARBAMIDE PEROXIDE (UNII: 31PZ 2VAU81) (HYDROGEN PEROXIDE - UNII: BBX060AN9V)

Inactive Ingredients		
Ingredient Name	Strength	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)		
GLYCERIN (UNII: PDC6A3C0OX)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)		
SODIUM LAURYL SULFATE (UNII: 368GB5141J)		
WATER (UNII: 059QF0KO0R)		

Packaging					
# Item Code	Package Description	Marketing Start Date	Marketing End Date		
1 NDC:68788- 8586-1	1 in 1 CARTON	02/20/2024			
1	15 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product				

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC Monograph Drug	M014	01/01/2019			

Labeler - Preferred Pharmaceuticals Inc. (791119022)

Registrant - Preferred Pharmaceuticals Inc. (791119022)

Establishment					
Name	Address	ID/FEI	Business Operations		
Preferred Pharmaceuticals Inc.		791119022	RELABEL(68788-8586)		

Revised: 2/2024 Preferred Pharmaceuticals Inc.