EAR WAX REMOVAL DROPS PREFERRED PLUS PHARMACY- carbamide peroxide - 6.5% solution/ drops

Kinray

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.

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 ear wax.

□ Warnings

Do not use if you have

- eardrainage, discharge, ear pain, irritation
- rashin the ear, or are dizzy
- injuryor perforation (hole) of the ear drum or after ear surgery

When using this product

- do not use for more than four days
- avoid contact with the eyes. If accidental contact with the eyes occurs, flush eyes with water and consult a doctor
- if excessive earwax remains after the use of this product, consult a doctor

Example 2 IKeep out of the 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 agel**:
- 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 earwax remaining after treatment may be removed by gently flushing the ear with warm water, using a soft rubber bulb ear syringe.
- When the ear canal is irrigated, the tip of the ear syringe should not obstruct the flow of water leaving the ear canal.
- **Children under 12 years** : consult a doctor.

Other information

- Protect from heat and direct sunlight
- Keep cap on bottle when not in use.
- Lot No. and EXP date: see label, bottom container or box.

Inactive ingredients

Citric Acid, Glycerin, Propylene Glycol, Sodium Citrate, Sodium Lauryl Sulfate, Tartaric Acid

Principal Display Panel Bottle Label 0.5 FL OZ



Purpose
Carbamide peroxide, 6.5% Earwax removal aid
Use for occasional use as an aid to soften, loosen and remove excessive earwax

Warnings
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away. If accidental contact with the eyes occurs, flush eyes with water and consult a doctor.

Directions

For use in the ear only.
Adults and children over 12 years of age:

 till 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 ear
 use twice daily for up to 4 days if needed, or as directed by doctor
 any earwax remaining after treatment may 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

See carton for full labeling

Kinray, Inc. Whitestone, NY 11357

Distributed By:



Manufactured by: Sheffield Pharmaceuticals New London, CT 06320 USA

Questions

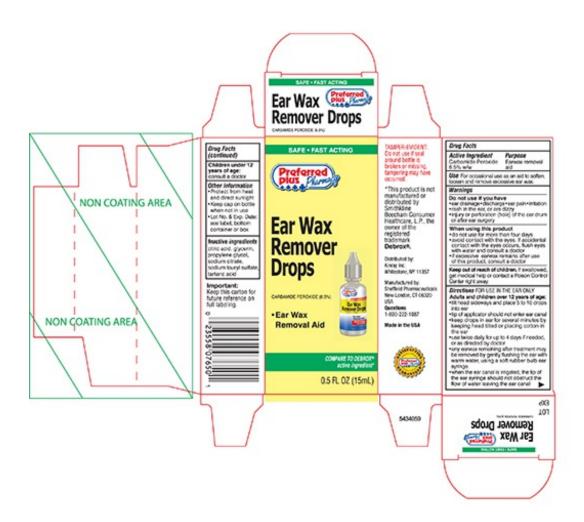
1-800-222-1087

#1613590 Rev 2/08

Preferred Plus Pharmacy Ear Wax Removal drops

Carbamide Peroxide 6.5% 0.5 FL OZ (15ml)

Principal Display Panel - Carton label 0.5 FL OZ



Preferred Plus Pharmacy

Ear wax Remover Drops

Carbamide Peroxide 6.5%

Ear wax removal aid

0.5 FL OZ (15ml)

EAR WAX REMOVAL DROPS PREFERRED PLUS PHARMACY

carbamide peroxide - 6.5% solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:61715-171
Route of Administration	TOPICAL		

l	Active Ingredient/Active Moiety				
ı	Ingredient Name	Basis of Strength	Strength		
	CARBAMIDE PERO XIDE (UNII: 31PZ2VAU81) (HYDROGEN PERO XIDE - UNII:BBX060AN9V)	CARBAMIDE PEROXIDE	65 mg in 1 mL		

T .*	T .	1
Inactiva	Ingra	diante
Inactive	Ingic	uiciits

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
ANHYDRO US CITRIC ACID (UNII: XF417D3PSL)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
TARTARIC ACID (UNII: W4888I119H)	

1	Packaging				
#	t Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:61715-171- 74	1 in 1 CARTON	06/09/2014		
1		15 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product			

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part344	06/09/2014	

Labeler - Kinray (012574513)

Registrant - Sheffield Pharmaceuticals LLC (151177797)

Establishment				
Name	Address	ID/FEI	Business Operations	
Sheffield Pharmaceuticals LLC		151177797	manufacture(61715-171)	

Revised: 11/2017 Kinray