

AURO DRI- isopropyl alcohol liquid
Insight Pharmaceuticals LLC

Auro Dri

AURO-DRI ®

Drug Facts

Active ingredient

Isopropyl alcohol 95% in an anhydrous glycerin 5% base

Purpose

Ear drying aid

Use

dries water in the ears and relieves water-clogged ears after

- swimming
- showering
- bathing
- washing the hair

Warnings

Flammable

Keep away from fire or flame

Do not use

in the eyes

Ask a doctor before use if you have

- ear drainage or discharge
- pain, irritation or rash in the ear
- had ear surgery
- dizziness

Stop use and ask a doctor if

irritation (too much burning) or pain occurs

Keep out of reach of children.

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

Directions

- unscrew cap from bottle
- remove foil safety seal from bottle
- affix applicator cap to bottle
- remove red applicator cap tip
- apply 4 to 5 drops in each affected ear

Other information

- recap bottle after each use
- keep away from heat or direct sunlight
- store at room temperature 15° - 25°C (59° - 77°F)
- do not use if foil seal imprinted with "SEALED FOR YOUR PROTECTION" under cap is broken or missing
- keep carton for full drug facts

Inactive ingredient

glycerin

Questions?

1-800-344-7239 Aurodri.com

PRINCIPAL DISPLAY PANEL

29.6 mL Bottle Carton

AURO- DRI®

**ISOPROPYL ALCOHOL 95% in an
ANHYDROUS GLYCERIN 5% base**

- Ear Drying Aid

CLEARs WATER FROM

SWIMMERS' EARS

1 FL OZ (29.6 mL)



B8255

AURO DRI

isopropyl alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63736-327
Route of Administration	AURICULAR (OTIC)		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	95 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	

Product Characteristics

Color	white (Clear)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63736-327-24	24 in 1 CASE	06/14/2010	
1		1 in 1 BOX		
1		29 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:63736-327-12	12 in 1 CASE	06/14/2010	
2		1 in 1 BOX		
2		29 mL in 1 BOTTLE; 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	06/14/2010	

Labeler - Insight Pharmaceuticals LLC (055665422)