

**CLINICAL STRENGTH MIRACLEDRY ANTIPERSPIRANT DEODORANT AM-
aluminum zirconium octachlorohydrex gly liquid
Highland Consumer Products**

Clinical Strength MiracleDry Antiperspirant Deodorant AM

Drug Facts

Active Ingredient

Aluminum Zirconium Octachlorohydrex Gly Anhydrous 18%
Antiperspirant

Uses

Treats excessive perspiration and reduces underarm wetness and odor.

Warnings

For external use only

Do not use

on irritated skin.

Keep out of reach of children

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

Stop use and contact a doctor

if rash or irritation develops

Ask your doctor

before using if you have kidney disease

Directions

Apply to underarms only. Use in the morning every day. Apply few strokes to underarms after tipping bottle to get maximum effect.

Inactive Ingredients

Water, PPG-11 Stearyl Ether, Cyclopentasiloxane, Steareth-2, Steareth-20, Isopropyl

Myristate, Glycine, Steareth-21

Storage

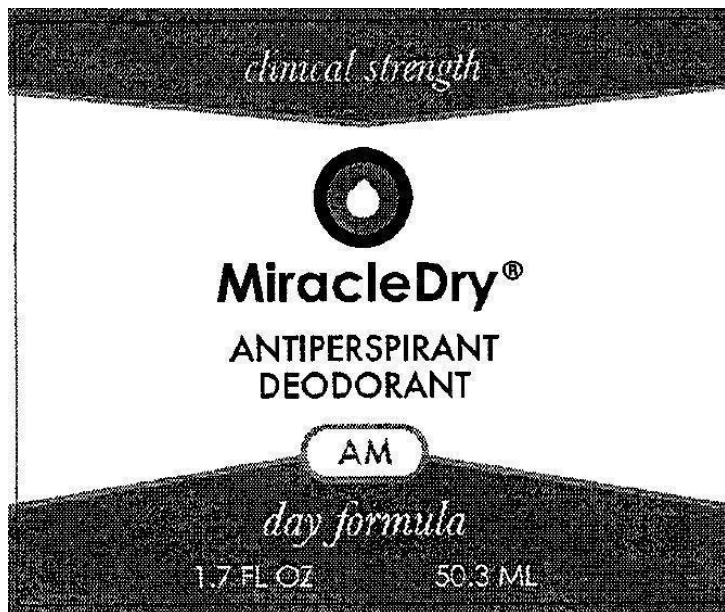
Keep at room temperature

Questions?

www.miracledry.com

clinical strength Miracle Dry ANTIPERSPIRANT DEODORANT AM day formula 1.7 FL OZ 50.3 ML

Product Label



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Storage Keep at room temperature	
Questions? www.miracledry.com	

Dist. by Highland Consumer Products, Inc. Franklin Lakes, NJ 07417

CLINICAL STRENGTH MIRACLEDRY ANTIPERSPIRANT DEODORANT AM

aluminum zirconium octachlorohydrate gly liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:53897-000
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALUMINUM ZIRCONIUM OCTACHLOROXYDREX GLY (UNII: P9D3YP29MY) (ALUMINUM ZIRCONIUM OCTACHLOROXYDREX GLY - UNII:P9D3YP29MY)	ALUMINUM ZIRCONIUM OCTACHLOROXYDREX GLY	18 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
PPG-11 STEARYL ETHER (UNII: S4G2J0Y0LG)	
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	
STEARETH-2 (UNII: V56DFE46J5)	
STEARETH-20 (UNII: L0Q8IK9E08)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
GLYCINE (UNII: TE7660XO1C)	
STEARETH-21 (UNII: 53J3F32P58)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53897-000-01	50.3 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/19/2017	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M019	10/15/2012	

Labeler - Highland Consumer Products (078647762)

Establishment			
Name	Address	ID/FEI	Business Operations
Englewood Lab, Inc.		172198223	manufacture(53897-000)

Revised: 12/2023

Highland Consumer Products