

**HCS ROLL-ON ANTIPERSPIRANT- aluminum chlorohydrate liquid
Concordance Healthcare Solutions, LLC.**

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

Aluminum Chlorohydrate 4.5%

Purpose

Antiperspirant

Use

Reduces underarm perspiration.

Warnings

For external use only

Do not use on broken skin. Discontinue use if irritation and redness develop. If condition persists for more than 72 hours consult a doctor.

Ask a doctor before use if you have kidney disease.

Keep out of reach of children. If swallowed get medical help and contact Poison Control Center right away.

Directions

Apply to underarms only.

Inactive ingredients

Ceteareth-24, Cetearyl Alcohol, Fragrance, Isopropyl Palmitate, Magnesium Aluminum Silicate, Methylparaben, Paraffinum Liquidum, Propylparaben, Sodium Benzoate, Water

Principal Display Panel - 44 mL Bottle, Dispensing Label

HCS

NDC 83506-004-01

Clean Fresh Scent

Roll-On

Antiperspirant/

Deodorant

- Fights Wetness & Odor

1.5 FL. OZ. (44ml)



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Reorder No. HCS0069



(01)00810132042195

Manufactured for: Concordance Healthcare Solutions, LLC.
Tiffin, OH 44883 · 1-800-447-0225
Made in Malaysia · #901-HCS0069 Rev. 2

HCS ROLL-ON ANTIPERSPIRANT

aluminum chlorohydrate liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83506-004
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Aluminum Chlorohydrate (UNII: HPN8MZ W13M) (Aluminum Chlorohydrate -	Aluminum	45 mg

UNII:HPN8MZ W13M)		Chlorohydrate	in 1 mL	
Inactive Ingredients				
Ingredient Name			Strength	
Water (UNII: 059QF0KO0R)				
Mineral Oil (UNII: T5L8T28FGP)				
Cetostearyl Alcohol (UNII: 2DMT128M1S)				
Isopropyl Palmitate (UNII: 8CRQ2TH63M)				
Magnesium Aluminum Silicate (UNII: 6M3P64V0NC)				
Methylparaben (UNII: A2I8C7HI9T)				
Propylparaben (UNII: Z8IX2SC1OH)				
Sodium Benzoate (UNII: OJ245FE5EU)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83506-004-01	44 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	06/06/2023	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part350	06/06/2023		

Labeler - Concordance Healthcare Solutions, LLC. (080220245)

Establishment			
Name	Address	ID/FEI	Business Operations
Ken Prima Cosmeceuticals SDN BHD		865792209	MANUFACTURE(83506-004)

Revised: 6/2023

Concordance Healthcare Solutions, LLC.