

**DERMFREE WHOLE BODY DEODORANT- aluminum chlorohydrate 9%whole
body deodorant cream**

Jiangxi Hemei Pharmaceutical Co., Ltd

84010-015

Active Ingredient

Aluminum chlorohydrate 9%

Purpose

Antiperspirant

Use

.Reduces underarm sweat
.24 hour effective protection

Warnings

For external use only

Do not use

on broken skin

When Using

Apply to underarms only

Stop Use

rash or irritation occurs

Ask Doctor

Ask a doctor before use if you have kidney disease

Keep Out Of Reach Of Children

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

Directions

Apply to underarms only

Inactive ingredients

Cetyl alcohol, deionized water, glycerin, GLYCERYL MONOOLEATE, GLYCERYL MONOSTEARATE, GR-270773 PHOSPHOLIPID EMULSION, Silicone Oil, zinc phenolsulfonate etc

PRINCIPAL DISPLAY PANEL

DERMFREE™

ODOR CONTROL Whole Body Deodorant

💧 FOR PITS, PRIVATES
& BEYOND

💧 Invisible cream

3.0 OZ (85g) NET WT

• DRUG FACTS

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GLYCERYL MONOOLEATE, GLYCERYL
MONOSTEARATE, GR-270773 PHOSPHOLIPID
EMULSION, Silicone Oil, zinc phenolsulfonate,
etc.



DISTRIBUTED BY: GERMANY HEALMUSZ
INTERNATIONAL CO., LTD. UNIT 616,
6/F, KAM TEEM INDUSTRIAL BUILDING,
135 CONNAUGHT ROAD WEST,
SAI WAN, H.K. WWW.HEALMUSZ.CN



DERMFREE WHOLE BODY DEODORANT

aluminum chlorohydrate 9% whole body deodorant cream

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:84010-015

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALUMINUM CHLOROHYDRATE (UNII: HPN8MZ W13M) (ALUMINUM CHLOROHYDRATE - UNII:HPN8MZ W13M)	ALUMINUM CHLOROHYDRATE	9 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
GLYCERYL MONOOLEATE (UNII: C4YAD5F5G6)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
WATER (UNII: 059QF0K00R)	
GR-270773 PHOSPHOLIPID EMULSION (UNII: D4B2F53PBH)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
GLYCERIN (UNII: PDC6A3C0OX)	
ZINC PHENOLSULFONATE (UNII: 4O71YT5YB5)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84010-015-01	85 g in 1 BOTTLE; Type 0: Not a Combination Product	04/30/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M019	04/30/2024	

Labeler - Jiangxi Hemei Pharmaceutical Co., Ltd (724892056)

Establishment

Name	Address	ID/FEI	Business Operations
Jiangxi Hemei Pharmaceutical Co., Ltd		724892056	manufacture(84010-015)

Revised: 4/2024

Jiangxi Hemei Pharmaceutical Co., Ltd