DOVE- sensitive 48h antiperspirant deodorant stick
Conopco Inc. d/b/a/ Unilever

Dove Sensitive 48h Antiperspirant Deodorant

DOVE SENSITIVE 48H ANTIPERSPIRANT DEODORANT - aluminum zirconium tetrachlorohydrex gly stick

Drug Facts

Active ingredient

Aluminum Zirconium Tetrachlorohydrex GLY (15.2 %)

Purpose

antiperspirant

Uses

reduces underarm wetness

48 Hour Protection

Warnings

- **For external** use only.
- **Do not use** on broken skin **.**
- **Ask a doctor before use if you have** kidney disease.
- **Stop use** if rash or irritation occurs.
- **Keep out of reach of children.**

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

Directions

apply to underarms only

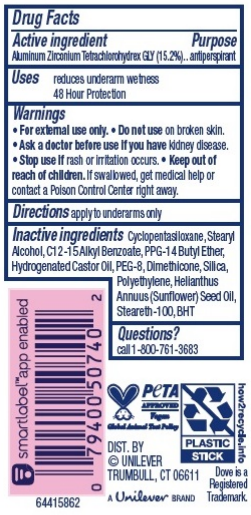
Inactive ingredients

Cyclopentasiloxane, Stearyl Alcohol, C12-15 Alkyl Benzoate, PPG-14 Butyl Ether, Hydrogenated Castor Oil, PEG-8, Dimethicone, Fragrance (Parfum), Silica, Polyethylene, Helianthus Annuus (Sunflower) Seed Oil, Steareth-100, BHT

Questions?

Call 1-800-761-3683

Packaging



DOVE

sensitive 48h antiperspirant deodorant stick

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:64942-2341
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALUMINUM ZIRCONIUM TETRACHLOROHYDREX GLY (UNII: 8O386558JE) (ALUMINUM ZIRCONIUM TETRACHLOROHYDREX GLY - UNII:8O386558JE)	ALUMINUM ZIRCONIUM TETRACHLOROHYDREX GLY	15.2 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
HYDROGENATED CASTOR OIL (UNII: ZF94AP8MEY)	
HIGH DENSITY POLYETHYLENE (UNII: UG00KM4WR7)	
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
PPG-14 BUTYL ETHER (UNII: R199TJT95T)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
SUNFLOWER OIL (UNII: 3W1JG795YI)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
STEARETH-100 (UNII: 4OH5W9UM87)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:64942-2341-1	74 g in 1 CONTAINER; Type 0: Not a Combination Product	12/01/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M019	12/01/2024	

Labeler - Conopco Inc. d/b/a/ Unilever (001375088)

Revised: 8/2024 Conopco Inc. d/b/a/ Unilever