PERSONAL CARE FOR MEN QUICK DRY ANTIPERSPIRANT- aluminum chlorohydrate 20.2% spray
Delta Brands Inc

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.

Active Ingredient

Aluminum Chlorohydrate 20.2%

Purpose Antiperspirant

Use reduces underarm wetness

Warnings

Flammable. Do not use near heat, flame or while smoking. Can cause serious Injury or death.
■ keep away from face and mouth to avoid breathing in ■ avoid spraying in eyes. Contents under pressure. Do not puncture or incinerate. Do not expose to heal or store at temperature above 120°F/SO"C or in enclosed places that could overheat ■ do not use on broken skin.

Stop use if rash or irritation occurs

ask a doctor before using if you have kidney disease

Keep out of reach of children.

Use only as directed. Intentional misuse by deliberately concentrating and Inhaling the contents can be harmful or fatal.
Help stop inhalation abuse.
For information visit www.inhalant.org

Directions ■ *hold can 6 inches away from the underarm* ■ *apply to underarms only*

Inactive Ingredients
butane, hydrofluorocarbon 152A, cyclopentasiloxane, alcohol denat., PPG-14 butyl ether, isobutane, propane, disteardimonium, hectorlte, C12-15 alkyl benzoate, helianthus annuus (sunflower) seed oil, dimethiconol, propylene carbonate, octyIdodecanol, fragrance, tocopheryl acetate, BHT

Package Label

PERSONAL CARE FOR MEN QUICK DRY ANTIPERSPIRANT

aluminum chlorohydrate 20.2% spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:20276-906
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALUMINUM CHLOROHYDRATE (UNII: HPN8MZW13M) (ALUMINUM CHLOROHYDRATE - UNII:HPN8MZW13M)	ALUMINUM CHLOROHYDRATE	20.2 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
BUTANE (UNII: 6LV4FOR43R)	
1,1-DIFLUOROETHANE (UNII: 0B1U8K2ME0)	
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	

ALCOHOL (UNII: 3K9958V90M)	
PPG-14 BUTYL ETHER (UNII: R199TJT95T)	
ISOBUTANE (UNII: BXR49TP611)	
PROPANE (UNII: T75W9911L6)	
DISTEARDIMONIUM HECTORITE (UNII: X687XDK09L)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
SUNFLOWER OIL (UNII: 3W1JG795YI)	
DIMETHICONOL (40 CST) (UNII: 343C7U75XW)	
PROPYLENE CARBONATE (UNII: 8D08K3S51E)	
OCTYLDODECANOL (UNII: 461N1O614Y)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:20276-906-34	34 g in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	02/01/2019	
2	NDC:20276-906-45	45 g in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	02/01/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part350	02/01/2019	

Labeler - Delta Brands Inc (102672008)

Revised: 2/2019 Delta Brands Inc