DEGREE EVEREST ANTIPERSPIRANT- aluminum chlorohydrate aerosol, spray Conopco Inc. d/b/a Unilever

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.

Degree Everest Dry Spray Antiperspirant

Active Ingredient:

Aluminum Chlorohydrate (23.3%)

Purpose:

anti-perspirant

Uses:

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 heat or store at temperatures above 120 ° F/50° 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.
- 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

• KEEP OUT OF REACH OF CHILDREN

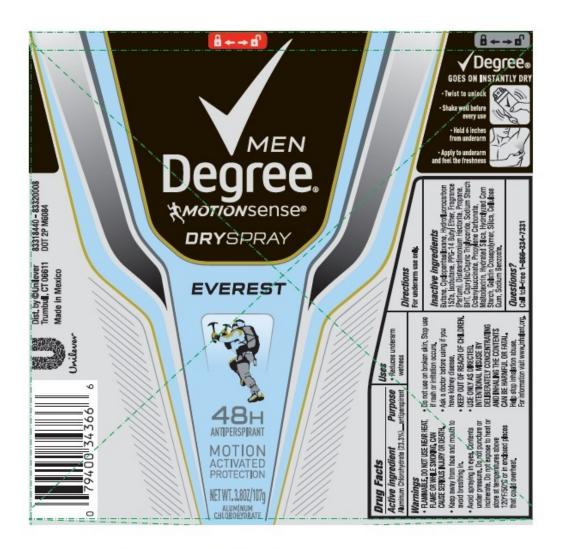
Directions:

For underarm use only.

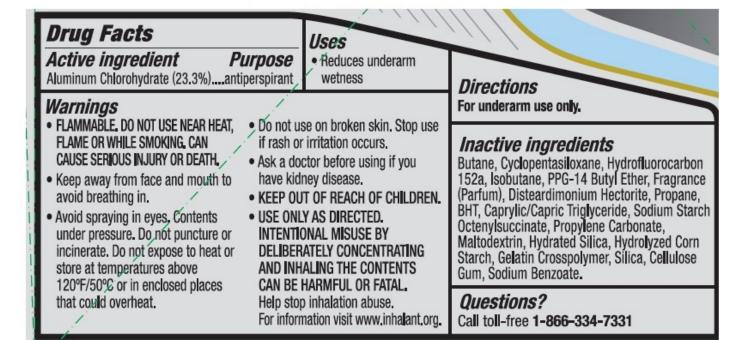
Inactive Ingredients: Butane, Cyclopentasiloxane, Hydrofluorocarbon 152a, Isobutane, PPG-14 Butyl Ether, Fragrance (Parfum), Disteardimonium Hectorite, Propane, BHT, Caprylic/Capric Triglyceride, Sodium Starch Octenylsuccinate, Propylene Carbonate, Maltodextrin, Hydrated Silica, Hydrolyzed Corn Starch, Gelatin Crosspolymer, Silica, Cellulose Gum, Sodium Benzoate.

Questions? Call toll-free 1-866-334-7331

Packaging



DRUG FACTS PANEL



DEGREE EVEREST ANTIPERSPIRANT

aluminum chlorohydrate aerosol, spray

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:64942-1380
Route of Administration	TOPICAL		

l	Active Ingredient/Active Moiety		
l	Ingredient Name	Basis of Strength	Strength
	ALUMINUM CHLOROHYDRATE (UNII: HPN8 MZW13M) (ALUMINUM CHLOROHYDRATE - UNII: HPN8 MZW13M)	ALUMINUM CHLOROHYDRATE	23.3 g in 100 g

Inactive Ingredients		
Ingredient Name	Strength	
BUTANE (UNII: 6LV4FOR43R)		
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)		
1,1-DIFLUOROETHANE (UNII: 0B1U8K2ME0)		
ISOBUTANE (UNII: BXR49TP611)		
PPG-14 BUTYL ETHER (UNII: R199TJT95T)		
DISTEARDIMO NIUM HECTO RITE (UNII: X687XDK09L)		
PROPANE (UNII: T75W9911L6)		
BUTYLATED HYDRO XYTO LUENE (UNII: 1P9 D0 Z171K)		
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)		
PROPYLENE CARBONATE (UNII: 8 D0 8 K3 S51E)		
MALTO DEXTRIN (UNII: 7CVR7L4A2D)		
HYDRATED SILICA (UNII: Y6O7T4G8P9)		
STARCH, CORN (UNII: O8232NY3SJ)		
SILICON DIO XIDE (UNII: ETJ7Z6XBU4)		
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED FORM (UNII: K679 OBS 311)		
SODIUM BENZOATE (UNII: OJ245FE5EU)		

	Packaging				
l	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
l	1 NDC:64942-1380-1	107 g in 1 CAN; Type 0: Not a Combination Product	11/0 1/20 14		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part350	11/0 1/20 14		

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

Revised: 9/2020 Conopco Inc. d/b/a Unilever