

**ANTIPERSPIRANT BLUSH LILLY- aluminum chlorohydrate aerosol, spray**  
**Universal Distribution Center LLC**

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**DRY SPRAY ANTIPERSPIRANT BLUSH LILLY**

**DRUG FACTS**

**ACTIVE INGREDIENT**

Aluminum Chlorohydrate (23.3%)

**PURPOSE**

Antiperspirant

**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 use** on broken skin. Stop use if rash or irritation occurs.
- **Ask a doctor** before use 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.

**DIRECTIONS:**

Apply to underarms only.

**INACTIVE INGREDIENTS**

Cyclopentasiloxane, Butane, Propane, Isobutane, Hydrofluorocarbon 152A, PPG-14 Butyl Ether, C12-15 Alkyl Benzoate, Dimethiconol, Disteardimonium Hectorite, Helianthus Annuus (Sunflower) Seed Oil, Octyldodecanol, Fragrance (Parfum), BHT, Propylene Carbonate, Tocopheryl Acetate.

**Compare to Dove®  
Advanced Care Beauty Finish**

**48 HOURS**

Quick-Dry  
No Residue

**SWEAT PROTECTION**

- *Shake well before each use*
- *Hold can 6 inches from underarm and spray evenly*
- *Allow to dry before dressing for soft, smooth-feeling skin*

Made in China

Mfd for and Distributed By:  
**Universal Distribution Center LLC**  
330 Applegarth Road,  
Monroe Township, NJ 08831  
**[www.universaldc.com](http://www.universaldc.com)**

\*This product is not manufactured, distributed, or endorsed by Unilever, the owner of the registered trademark Dove® Advanced Beauty Finish.

**Packaging**



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# DRY SPRAY

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Advanced Care Beauty Finish



**BLUSH LILLY**

**48** | Quick-Dry  
HOURS | No Residue

**SWEAT PROTECTION**  
**ANTIPERSPIRANT**

NET WT. 3.5 OZ (99 g)

## ANTIPERSPIRANT BLUSH LILLY

aluminum chlorohydrate aerosol, spray

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:52000-446
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>ALUMINUM CHLOROHYDRATE</b> (UNII: HPN8MZ W13M) (ALUMINUM CHLOROHYDRATE - UNII:HPN8MZ W13M)	ALUMINUM CHLOROHYDRATE	23.3 g in 100 g

## Inactive Ingredients

Ingredient Name	Strength
<b>CYCLOMETHICONE 5</b> (UNII: 0THT5PCI0R)	
<b>BUTANE</b> (UNII: 6LV4FOR43R)	
<b>PROPANE</b> (UNII: T75W9911L6)	
<b>ISOBUTANE</b> (UNII: BXR49TP611)	
<b>1,1-DIFLUOROETHANE</b> (UNII: 0B1U8K2ME0)	
<b>PPG-14 BUTYL ETHER</b> (UNII: R199TJT95T)	
<b>ALKYL (C12-15) BENZOATE</b> (UNII: A9EJ3J61HQ)	
<b>DIMETHICONOL (40 CST)</b> (UNII: 343C7U75XW)	
<b>DISTEARDIMONIUM HECTORITE</b> (UNII: X687XDK09L)	
<b>SUNFLOWER OIL</b> (UNII: 3W1JG795YI)	
<b>OCTYLDODECANOL</b> (UNII: 461N1O614Y)	
<b>BUTYLATED HYDROXYTOLUENE</b> (UNII: 1P9D0Z171K)	
<b>PROPYLENE CARBONATE</b> (UNII: 8D08K3S51E)	
<b>.ALPHA.-TOCOPHEROL ACETATE</b> (UNII: 9E8X80D2L0)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52000-446-35	99 g in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	05/04/2026	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M019	05/04/2026	

**Labeler** - Universal Distribution Center LLC (019180459)

Revised: 2/2026

Universal Distribution Center LLC