BATH AND BODY WORKS OCEAN- mens collection antiperspirant deodorant aerosol, spray Bath & Body Works, 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.

DRUG FACTS

Drug Facts

Active ingredient Aluminum Zirconium Trichlorohydrex GLY (16%)

Purpose

Antiperspirant

Use

Reduces underarm wetness

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 the reach of children.** If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Apply to underarms only.

Inactive ingredients

Dimethicone, Stearyl Alcohol, C15-19 Alkane, Diheptyl Succinate, C12-15 Alkyl Benzoate, Hydrogenated Castor Oil, Cetyl Alcohol, Water (Aqua, Eau), Silica, Aluminum Starch Octenylsuccinate, Fragrance (Parfum), Capryloyl Glycerin/Sebacic Acid Copolymer, Tocopheryl Acetate.

Packaging



BATH AND BODY WORKS OCEAN

mens collection antiperspirant deodorant aerosol, spray

| Ρ | roduct Infori | mation | | | | | | | | | | |
|---|----------------------|--|----------------------------|------------|-------------------------|----------------|-----------------------|------------------|--|--|--|--|
| Pı | roduct Type | Type HUMAN OTC DRUG Item Code (Source) | | | | NDC:62670-6127 | | | | | | |
| R | oute of Adminis | stration | TOPICAL | | | | | | | | | |
| | | | | | | | | | | | | |
| Active Ingredient/Active Moiety | | | | | | | | | | | | |
| | | | | | | | Strength | Strength | | | | |
| Aluminum Zirconium Trichlorohydrex GLY (UNII: T27D6T99LH) (ALUMINUM Z IRCONIUM TRICHLOROHYDREX GLY - UNII:T27D6T99LH)Aluminum Zircon Trichlorohydrex GL | | | | | | | | 16 g in 100 g | | | | |
| | | | | | | | | | | | | |
| Inactive Ingredients | | | | | | | | | | | | |
| | | Strength | | | | | | | | | | |
| Di | methicone (UNII: | 92RU3N3Y1O) | | | | | | | | | | |
| | | | | | | | | | | | | |
| P | ackaging | | | | | | | | | | | |
| # | Item Code | Pac | kage Description | Mark | Marketing Start Date | | Marketing End Date | | | | | |
| 1 | NDC:62670- 6127-0 | 77 g in 1 CAN; Product | : Type 0: Not a Combinatio | n 08/13/20 | 08/13/2021 | | | | | | | |
| | | | | | | | | | | | | |
| | | | | | | | | | | | | |
| Μ | larketing l | nformat | ion | | | | | | | | | |
| | | | | | | | | | | | | |

| Marketing | Application Number or Monograph | Marketing Start | Marketing End |
|---------------------|---------------------------------|-----------------|---------------|
| Category | Citation | Date | Date |
| OTC monograph final | part350 | 08/13/2021 | |

Labeler - Bath & Body Works, Inc. (878952845)

Establishment

| Name | Address | ID/FEI | Business Operations |
|--------------------------------------|---------|-----------|----------------------------|
| KDC/ONE Development Corporation, Inc | | 204006464 | manufacture(62670-6127) |

Revised: 8/2021

Bath & Body Works, Inc.