STERILLIUM RUB FRAGRANCE FREE- alcohol liquid
BODE Chemie GmbH

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

Active Ingredient

Ethyl Alcohol 85% w/w

Purpose

Antiseptic

Uses

surgical hand antiseptic

significantly reduces the number of micro-organisms on the hands and forearms prior to surgery or patient care

Warnings

For external use only.

Flammable, keep away from fire or flame.

Do not use in or near the eyes or on mucous membranes.

When using this product and contact with the eyes occurs, flush immediately with water.

Stop use and ask a doctor if

irritation and redness develop

condition persists for more than 72 hours

Keep out of reach of children.

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

Directions

apply to clean, dry hands

for the first use of the day, use a nail pick

dispense approx. 2 mL into hand, dip fingers of opposite hand into palm, working product under nails and into cuticles

repeat procedure with other hand

with hands still moist spread around the hand and lower 1/3 of the forearm

reapply the product to the hands, paying particular attention to fingers, cuticles, and interdigital spaces

following application, rub hands until dry

hands should remain moist for entire application time, approx. 1.5 minutes.

Inactive Ingredients

Water, n-Propyl Alcohol, Myristyl Alcohol, Methylethylketone, Glycerin.

Bode

Sterillium Rub

Fragrance-Free

Procedural Scrub

Surgical

Hand Antiseptic

85% w/w ethyl alcohol with emollients

Indication: Significantly reduces the number of micro-organisms on the hands and forearms prior to surgery or patient care.

Fast-acting. Dermatologically tested.

Gentle to skin.

Read Drug Facts panel before use.

Lot:

Use by:

Questions? Call 1-800-MEDLINE

NDC 65616-004-06

1000 ml 33.8 fl. oz.

STERILLIUM RUB FRAGRANCE FREE

alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:65616-004
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	85 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
PROPYL ALCOHOL (UNII: 96F264O9SV)	
MYRISTYL ALCOHOL (UNII: V42034O9PU)	
METHYL ETHYL KETONE (UNII: 6PT9KLV9IO)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:65616-004-06	1000 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/28/2012	
2	NDC:65616-004-08	50 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	03/28/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	03/28/2012	04/01/2021

Labeler - BODE Chemie GmbH (316039007)

Establishment

Name	Address	ID/FEI	Business Operations
BODE Chemie GmbH		316039007	manufacture(65616-004)

Revised: 9/2019 BODE Chemie GmbH