

AVAGARD D- alcohol lotion
3M ESPE Dental Products

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, 61% w/w

Purpose

Antiseptic

Uses

instant healthcare personnel hand antiseptic

- reduces bacteria that potentially can cause disease
- recommended for repeated use

Warnings

☐Flammable, keep away from fire or flame.

For external use only

When using this product

- **Keep out of eyes.** If contact with eyes occurs, rinse promptly and thoroughly with water.

Stop use and ask a doctor if significant irritation, or sensitization develops

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. Wet hands thoroughly with product and allow to dry.

Other information

- Store at 20-25°C (68-77°F)

Inactive ingredient

beheneth-10, behenyl alcohol, C20-40 pareth-24, cetyl palmitate, diisopropyl dimer dilinoleate, dimethicone, glycerin, polyethylene glycol, squalane, water

Principal Display Panel

NDC 48878-0231-7

3M ESPE

Avagard™ D

Instant Hand Antiseptic

With Moisturizers

Contains: 61% w/w ethyl alcohol

Destroys Bacteria. Not Your Skin.

Flammable, keep away from fire or flame, heat,

sparks and sources of static discharge.

REF 2317

500 ml • 16 fl oz

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3M™ Avagard™ D Instant Hand Antiseptic helps maintain skin integrity. Latex glove and CHG compatible. 3M, ESPE and Avagard are trademarks of 3M or 3M Deutschland GmbH. © 3M 2018. All rights reserved. Patent No. US 6,090,395; US 6,534,069; US 6,623,744; US 7,081,246

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Made in USA for
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34-8723-7973-9-A

AVAGARD D

alcohol lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48878-0231
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Alcohol (UNII: 3K9958V90M) (Alcohol - UNII:3K9958V90M)	Alcohol	530.7 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
Beheneth-10 (UNII: 313S43DM16)	
Docosanol (UNII: 9G1OE216XY)	
Cetyl Palmitate (UNII: 5ZA2S6B08X)	
Dimethicone (UNII: 92RU3N3Y1O)	
Glycerin (UNII: PDC6A3C0OX)	
Polyethylene Glycol, unspecified (UNII: 3WJQ0SDW1A)	

Squalane (UNII: GW89575KF9)	
Water (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48878-0231-7	12 in 1 CASE	04/28/2011	
1		500 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	10/01/1999	

Labeler - 3M ESPE Dental Products (801390852)

Revised: 4/2020

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