3M AVAGARD FOAMING INSTANT HAND ANTISEPTIC- alcohol liquid 3M Health Care

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.

3M™ Avagard™ Foaming Instant Hand Antiseptic

Drug Facts

Active ingredient

Ethyl Alcohol, 70% v/v (62% w/w)

Purpose

Antiseptic

Uses

instant healthcare personnel hand antiseptic to reduce bacteria that potentially can cause disease

- instant hand antiseptic to decrease bacteria on the skin
- recommended for repeated use

Warnings

For external use only.

Flammable, keep away from fire or flame.

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. Apply sufficient amount to thoroughly wet all surfaces of hands and fingers. Rub onto hands until dry.

• Supervise children in the use of this product.

Other information

Store at 20-25°C (68-77°F)

Inactive Ingredients

denatonium benzoate PEG/PPG-18/18 dimethicone, PEG-23M, water

Questions?

call 1-800-228-3957 (Monday to Friday 7AM - 6PM CST)

Made in Canada for **3M Health Care**2510 Conway Ave.St PauL MN 551443M.com/Medical

3M and Avagard are trademarks of 3M. ©2017, 3M.All rights reserved. Patent: 3M.com/Patents 34-8720-7424-9

Principal Display Panel – 1000 mL Cartridge Label

3M NDC 17518-055-00

Avagard[™] Foaming

Instant Hand Antiseptic

Contains: 70% v/v ethyl alcohol

Fragrance-Free

Latex glove and CHG compatible

Directions: Thoroughly wet hands and fingers. Rub until dry.

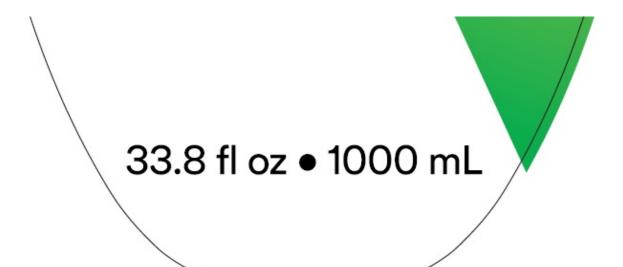
Flammable, keep away from fire or flame, heat, sparks and sources of static discharge.

REF

9322A

33.8 fl oz • 1000 mL





3M™ Avagard™ Foaming Instant Hand Antiseptic kills greater than 99.999% of harmful bacteria in 15 seconds (in vitro) while helping maintain skin condition. Latex glove and CHG compatible.

		d .
Drug Facts		ý
Active ingredient Ethyl Alcohol, 70% v/v (62% w/w)	Purpose Antiseptic	6
	hand antiseptic to reduce bacteria that potentially ntiseptic to decrease bacteria on the skin	7727
Warnings For external use only. Flammable, keep away from fire o	r flames.	7387
When using this product keep out of e	eyes. If contact with eyes occurs, rinse promptly and thoroughly with water.	
Stop use and ask a doctor if signi	ficant irritation or sensitization develops.	
Keep out of reach of children, If swa	owed, get medical help or contact a Poison Control Center right away.	~
	ands. Apply sufficient amount to thoroughly wet all surfaces of until dry. Supervise children in the use of this product.	
Other information Store at 2	0-25°C (68-77°F)	
Inactive ingredients denator PEG-23M, water	nium benzoate, PEG/PPG-18/18 dimethicone,	
Questions? call 1-800-228-395	7 (Monday to Friday 7AM - 6PM CST)	
Made in Canada for J 3M Health Care, 2510 Conway Ave., St. Paul, MN 55144 3M.com/Medical	3M and Avagard are trademarks of 3M. © 2017, 3M. All rights reserved. Patent:3M.com/patents 34-8720-7424-9	4

3M AVAGARD FOAMING INSTANT HAND ANTISEPTIC

alcohol liquid

Product Infor	mation					
Product Type		HUMA	HUMAN OTC DRUG Item Code (So		de (Source)	NDC:17518-055
Route of Admin	stration	ТОРЮ	CAL			
Active Ingred	ient/Acti	vo Moietv				
Active Ingree	ic no Acu	Ingredient N	ame		Basis of Strength	Strength
Alcohol (UNII: 3K9958V90M) (Alcohol - UNII:3K9958V90M)					Alcohol	548 mg in 1 mL
Inactive Ingre	dients					
Ingredient Name						Strength
Denatonium Ben	,					
PEG/PPG-18/18 D						
POLYETHYLENE	OXIDE 10	00000 (UNII: HZ	58M6D839)			
Water (UNII: 0590	DO IZO O DI					
	(FUKOUR)					
	(FOKOOR)					
Packaging	(FOKOUR)	Pack	age Description		Marketing Start Date	Marketing End Date
Packaging # Item Code			age Description			-
Packaging # Item Code 1 NDC:17518-055 02	- 25 in 1 C	CASE 1 1 BOTTLE, PUI	age Description MP; Type 0: Not a Co	o mbinatio n	Date	-
Packaging H Item Code NDC:17518-055 02 NDC:17518.055	25 in 1 C 50 mL i Product	CASE 1 1 BOTTLE, PUI		o mbinatio n	Date	-
Packaging Item Code I NDC:17518-055 I NDC:17518-055	 25 in 1 0 50 mL is Product 12 in 1 0 500 mL Product 	CASE 1 BOTTLE, PUI CASE in 1 BOTTLE, PU			Date 05/27/2011	-
Packaging Item Code I NDC:17518-055 I NDC:17518-055	25 in 1 C	CASE n 1 BOTTLE, PUI CASE in 1 BOTTLE, PU	MP; Type 0: Not a Co UMP; Type 0: Not a (Combinatio n	Date 05/27/2011	-
Packaging # Item Code 1 NDC:17518-055 1 NDC:17518-055 2 NDC:17518-055 3 NDC:17518-055	25 in 1 C	CASE n 1 BOTTLE, PUI CASE in 1 BOTTLE, PU ASE L in 1 CARTRIDG	MP; Type 0: Not a Co	Combinatio n	Date 05/27/2011 05/27/2011	-
Packaging # Item Code 1 NDC:17518-055 1 NDC:17518-055 2 NDC:17518-055 3 NDC:17518-055	 25 in 1 0 50 mL ii Product 12 in 1 0 500 mL 500 mL 12 in 1 0 500 mL 10 0 m 	CASE n 1 BOTTLE, PUI CASE in 1 BOTTLE, PU ASE L in 1 CARTRIDG	MP; Type 0: Not a Co UMP; Type 0: Not a (Combinatio n	Date 05/27/2011 05/27/2011	-
Packaging Item Code NDC:17518-055 02 NDC:17518-055 01 NDC:17518-055 01 NDC:17518-055 01 NDC:17518-055 01 NDC:17518-055 01	 25 in 1 0 50 mL i Product 12 in 1 0 500 mL product 5 in 1 C. 1000 m Product 	CASE n 1 BOTTLE, PUI CASE in 1 BOTTLE, PU ASE L in 1 CARTRIDG	MP; Type 0: Not a Co UMP; Type 0: Not a (Combinatio n	Date 05/27/2011 05/27/2011	-
Packaging # Item Code 1 NDC:17518-055 1 NDC:17518-055 2 NDC:17518-055 3 NDC:17518-055	 25 in 1 C 50 mL i 50 mL i Product 12 in 1 C 500 mL Product 5 in 1 C 1000 m Product 	CASE n 1 BOTTLE, PUI CASE in 1 BOTTLE, PU ASE L in 1 CARTRIDO	MP; Type 0: Not a Co UMP; Type 0: Not a (Combination mbination	Date 05/27/2011 05/27/2011	-

Labeler - 3M Health Care (006173082)