

V W- advance hand sanitizer gel
VAL USA Manufacturer 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.

VAL-GEL

Active Ingredient(s)

Alcohol 75% v/v. Purpose: Antiseptic

Purpose

Antiseptic, Hand Sanitizer

Use

Hand Sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

Warnings

For external use only. Flammable. Keep away from heat or flame

Do not use

- in children less than 2 months of age
- on open skin wounds

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

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

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Directions

- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

Other information

- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients


Aloe Barbadosensis leaf Juice, Carbomer, FD&C Blue No.1, FD&C Yellow No.5, Glycerin, Propylene

Glycol, Tocopheryl Acetate Vitamin E), Triethanolamine, Water

Package Label - Principal Display Panel

V&W[®]
since 1984

HADVANCE
HAND
SANITIZER
REFRESHING GEL
KILLS **99.99%** GERMS



Ethyl Alcohol Antiseptic **75%**






KILLS 99.99% ILLNESS
CAUSING GERMS
LEAVES HANDS FEELING SOFT

(500ml / 16 oz)

Drug Facts

Active ingredient Ethyl alcohol 75% v/v.....	Purpose Antiseptic
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Do not use <ul style="list-style-type: none"> ■ In children less than 2 months of age ■ On open skin wounds 	
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Directions <ul style="list-style-type: none"> ■ Place enough product on hands to cover all surfaces. Rub hands together until dry. ■ Supervise children under 6 years of age when using this product to avoid swallowing. 	
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Inactive ingredients Aloe Barbadosensis leaf Juice, Carbomer, FD&C Blue No.1, FD&C Yellow No.5, Glycerin, Propylene Glycol, Tocopheryl Acetate Vitamin E), Triethanolamine, Water	
Questions? call 1-626-839-8069	

Tel: 86-20-3652 0253 Email: info@vw1984.com
www.vw1984.com
Distributed by:Val USA Manufacturer, Inc, 1050 W Central Ave, Unit A, Brea, CA 92821

Made in china

500 mL NDC: 76687-005-13

V W			
advance hand sanitizer gel			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76687-005
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
	Ingredient Name	Basis of Strength	Strength
	ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 mL in 100 mL
Inactive Ingredients			
	Ingredient Name		Strength
	PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
	TROLAMINE (UNII: 9O3K93S3TK)		
	FD&C BLUE NO. 1 (UNII: HBR47K3TBD)		
	FD&C YELLOW NO. 5 (UNII: I753WB2F1M)		

GLYCERIN (UNII: PDC6A3C0OX)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76687-005-01	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2020	
2	NDC:76687-005-02	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2020	
3	NDC:76687-005-03	80 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2020	
4	NDC:76687-005-04	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2020	
5	NDC:76687-005-05	120 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2020	
6	NDC:76687-005-06	150 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2020	
7	NDC:76687-005-07	180 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2020	
8	NDC:76687-005-08	200 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2020	
9	NDC:76687-005-09	240 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2020	
10	NDC:76687-005-10	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2020	
11	NDC:76687-005-11	300 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2020	
12	NDC:76687-005-12	350 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2020	
13	NDC:76687-005-13	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2020	
14	NDC:76687-005-14	650 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2020	
15	NDC:76687-005-15	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2020	
16	NDC:76687-005-16	3800 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2020	
17	NDC:76687-005-17	5000 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2020	
18	NDC:76687-005-18	20000 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2020	
19	NDC:76687-005-19	50000 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2020	
20	NDC:76687-005-20	1000000 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/15/2020	

Labeler - VAL USA Manufacturer Inc (068401328)

Establishment

Name	Address	ID/FEI	Business Operations
Guangzhou Youxing Cosmetics CO.,LTD.		544355953	manufacture(76687-005)

Revised: 5/2020

VAL USA Manufacturer Inc