VAX-MAN ANTISEPTIC HAND WIPES- alcohol cloth
Vaxman Group 2015 Ltd.

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.

Vax-Man Antiseptic Hand Wipes

Drug Facts

Active ingredients

Alcohol 70% v/v

Purpose

Antiseptic

Uses

- sanitizing wipe to help reduce bacteria that potentially can cause disease
- for use when soap and water are not available
- safe for repeated use

Warnings

For external use only.

Flammable. Keep away from heat or flame.

Do not use

as a diaper wipe

When using this product

avoid contact with eyes. If contact occurs, flush thoroughly with water.

Discontinue use

if irritation or redness develops. If condition persists for more than 72 hours contact a doctor.

Keep out of reach of children

unless under adult supervision. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Open the plastic cap. Open reusable sticker label. Pull up first wipe. Reseal sticker label and close plastic cap. Discard wipe in trash receptacle after use. Do not flush.

Other information

Store in a cool, dry place, away from direct sunlight

Contact: 415 Ocean Parkway Apt 2g. Brooklyn NY 11218

Inactive ingredients

Water, Aloe Barbadensis Leaf Juice, Glycerin, Potassium Sorbate, Sodium Benzoate, Fragrance, Lactic Acid.

Company Information

Vaxman Group 2015 Ltd

Haifa Israel

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www.vaxmangroupcom

Made in Israel

Product Packaging - 200 wipe canister

Vax-Man

Antiseptic Hand Wipes

For a better world

70% Alcohol

200 WIPES

VAX-MAN ANTISEPTIC HAND WIPES
alcohol cloth

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:79566-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
LACTIC ACID (UNII: 33X04XA5AT)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79566-001-02	50 in 1 POUCH	07/10/2020	
1		240 mL in 1 POUCH; Type 0: Not a Combination Product		
2	NDC:79566-001-04	100 in 1 POUCH	07/10/2020	
2		480 mL in 1 POUCH; Type 0: Not a Combination Product		
3	NDC:79566-001-03	80 in 1 POUCH	07/10/2020	
3		384 mL in 1 POUCH; Type 0: Not a Combination Product		
4	NDC:79566-001-01	1 in 1 PACKET	07/10/2020	
4		4.8 mL in 1 PACKET; Type 0: Not a Combination Product		
5	NDC:79566-001-05	80 in 1 CANISTER	07/10/2020	
5		150 mL in 1 CANISTER; Type 0: Not a Combination Product		
6	NDC:79566-001-06	100 in 1 CANISTER	07/10/2020	
6		150 mL in 1 CANISTER; Type 0: Not a Combination Product		
7	NDC:79566-001-07	140 in 1 CANISTER	07/10/2020	
7		210 mL in 1 CANISTER; Type 0: Not a Combination Product		
8	NDC:79566-001-08	200 in 1 CANISTER	07/10/2020	
8		420 mL in 1 CANISTER; Type 0: Not a Combination Product		
9	NDC:79566-001-09	400 in 1 CONTAINER	07/10/2020	
9		840 mL in 1 CONTAINER; Type 0: Not a Combination Product		

10	NDC:79566-001-10	500 in 1 CONTAINER	07/10/2020	
10		1050 mL in 1 CONTAINER; Type 0: Not a Combination Product		
11	NDC:79566-001-11	750 in 1 CONTAINER	07/10/2020	
11		1575 mL in 1 CONTAINER; 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	part333A	07/10/2020	

Labeler - Vaxman Group 2015 Ltd. (531964346)

Revised: 7/2020 Vaxman Group 2015 Ltd.