

**SANI-MAXX ANTIBACTERIAL WIPES- benzalkonium chloride cloth**  
**Tv Direct Llc**

*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.*

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**Sani-Maxx ANTI-BACTERIAL WET WIPES**

**Drug Facts**

**Active Ingredients**

Benzalkonium Chloride 0.13%

**Purpose**

Antiseptic/Hand & Skin Sanitizer

**USES** Sanitizing Hand Wipes to help decrease bacteria on the skin. Recommended for single use.

**WARNINGS:**

**Do not freeze. For external use only. Store at room temperature.**

**Do not use ears, eyes or mouth.**

- Avoid contact with eyes. In case of contact, flush eyes with water.
- Stop use and ask a doctor if redness or irritation develop and persist for more than 72 hours.
- Keep out of reach of children.
- Children should be supervised when using this product.
- If swallowed, get medical help or contact Poison Control Centre immediately.

**DIRECTIONS FOR USE:**

Wipe liberally over the hands & body, rub into skin until dry. Recommended for single use.

**Inactive Ingredients:**

Water (Aqua), Glycerol, Propanediol, Aloe Vera Extract, Amino acid moisturizer, Hexadecylpyridinium Chloride, Chlorphenesin, Vitamin E, Dodecyl polyglucoside, Citric acid, Sodium citrate.

**Questions?**

Call +1 212.268.2121 (Mon-Fri 9am to 5pm)

- Kill 99.9% of Germs and Bacteria.
- Easily Removes Dirt.
- Alcohol & Bleach Free.

Aloe Vera

Vitamin E

Anti-bacterial

Spulance Nonwoven

**CAUTION: KEEP OUT OF THE REACH OF CHILDREN**

Please dispose of used wipes responsibly.

Do not dispose used wipes down the toilet.

No Alcohol . No Bleach

DISTRIBUTED BY: TV DIRECT LLC, 385 FIFTH AVENUE, SUITE 809

NEW YORK, NY10016, USA

Wipe Dimensions: 5.9 in. X 7.87 in. (14.9cm x 19.9cm)

2020 TV Direct LLC.

MADE IN CHINA

Packaging

SANI-MAXX ANTIBACTERIAL WIPES				
benzalkonium chloride cloth				
Product Information				
Product Type		HUMAN OTC DRUG	Item Code (Source)	
Route of Administration		TOPICAL	NDC:78915-852	
Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)			BENZALKONIUM CHLORIDE	0.13 g in 100 mL
Inactive Ingredients				
Ingredient Name				Strength
WATER (UNII: 059QF0K00R)				
GLYCERIN (UNII: PDC6A3C00X)				
PROPANEDIOL (UNII: 5965N8W85T)				
ALOE VERA LEAF (UNII: ZY81Z83H0X)				
BETAINE (UNII: 3SCV180C9W)				
CETYLPYRIDINIUM CHLORIDE ANHYDROUS (UNII: 6BR7T22E2S)				
CHLORPHENESIN (UNII: I670DAL4SZ)				
.ALPHA.-TOCOPHEROL (UNII: H4N855PNZ1)				
LAURYL GLUCOSIDE (UNII: 76LN7P7UCU)				
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)				
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78915-852-80	80 in 1 BAG	08/12/2020	
1		0.491 mL in 1 PATCH; Type 0: Not a Combination Product		
2	NDC:78915-852-70	70 in 1 CANISTER	06/09/2020	
2		0.468 mL in 1 PATCH; Type 0: Not a Combination Product		
3	NDC:78915-852-50	50 in 1 BAG	06/09/2020	
3		0.28 mL in 1 PATCH; 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	06/09/2020	

