

**MAGIC POUCH- alcohol**  
**Merci Handy Corporation**

*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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**Magic Pouch, Hand Sanitizer Namaste Hand Sanitizer Glitter Fever Hand Sanitizer Unicorn Edition**

***Drug Facts***

***Active ingredient***

Alcohol 67%

***Purpose***

Antiseptic

***Use***

for handwashing to decrease bacteria on the skin

***Warnings***

**For external use only.**

Flammable, keep away from fire or flame.

**Do not use**

in the eyes. In case of contact, flush eyes with water.

**Stop use and ask a doctor if**

- irritation and redness develop
- condition persists for more than 72 hours

**Keep out of reach of children.**

If swallowed, get medical help or contact a Poison Control Center right away.

***Directions***

wet hands thoroughly with product and allow to dry without wiping

***Other information***

store at a temperature below 110° F (43° C)

### ***Inactive ingredients***

Water (Aqua), Fragrance (Parfum), Aloe Vera Leaf Juice Powder, Glycerin, Propylene Glycol, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aminomethyl Propanol, Mannitol, Microcrystalline Cellulose, Sucrose, Corn (Zea Mays) Starch, Denatonium Benzoate, Tocopheryl Acetate, Maltodextrin, Hydroxypropyl Methylcellulose, Potassium Sorbate, Sodium Benzoate, Hexyl Cinnamal, Benzyl Benzoate, Linalool, Limonene, Ferric Ferrocyanide, D&C Red No. 33, D&C Red No. 30, FD&C Blue No. 1.

### **QUESTIONS OR COMMENTS?**

**(646)-358-3432**

### ***Drug Facts***

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**QUESTIONS OR COMMENTS?**

(646)-358-3432

**Package Labeling:Kit**

# MAGIC POUCH

For you who always looks on the glittery side of life, we've gathered our 3 shiniest hand sanitizers into a magical, reusable holographic pouch...

## Unicorn Edition :

Its scent extracted from the purest rainbows, its holographic label and its multicolored glitter

## Namaste Edition :

Its magical mirror effect label and its soft scent (Very limited edition)

## Glitter Fever :

Its sparkling scent, its golden label and its stardust-sprinkled texture

### Drug Facts

**Active ingredient Purpose**  
Alcohol Denat 67%.....Antiseptic

### Use

■ for handwashing to decrease bacteria on the skin

### Warnings

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**Do not use** in the eyes. In case of contact, flush eyes with water.

**Stop use and ask a doctor if**

■ irritation and redness develop

■ condition persists for more than 72 hours

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

### Directions

wet hands thoroughly with product and allow to dry without wiping

### Other information

store at a temperature below 110° F (43° C)

### Inactive ingredients

### GLITTER FEVER

Aqua (Water), Parfum (Fragrance), Aloe Barbadensis Leaf Juice Powder, Glycerin, Propylene Glycol, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aminomethyl Propanol, Mannitol, Microcrystalline Cellulose, Sucrose, Zea Mays (Corn) Starch, Tocopheryl Acetate, Maltodextrin, Hydroxypropyl Methylcellulose, Potassium Sorbate, Sodium Benzoate, Synthetic Fluorophlogopite, Coumarin, Limonene, Alphalsolethyl Ionone, Cinnamyl Alcohol, Isoeugenol, CI 77891 (Titanium Dioxide), CI 19140 (Yellow 5), CI 77492 (Iron Oxide Yellow), CI 14700 (Red 4).

### NAMASTE

Aqua (Water), Parfum (Fragrance), Aloe Barbadensis Leaf Juice Powder, Glycerin, Propylene Glycol, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aminomethyl Propanol, Mannitol, Microcrystalline Cellulose, Sucrose, Zea Mays (Corn) Starch, Tocopheryl Acetate, Maltodextrin, Hydroxypropyl Methylcellulose, Potassium Sorbate, Sodium Benzoate, Hexyl Cinnamal, Benzyl Benzoate, Linalool, Limonene, CI 77510 (Red 33), CI 73360 (Red 30 Lake), CI 42090 (Blue 1).

### UNICORN EDITION

Aqua (Water), Parfum (Fragrance), Aloe Barbadensis Leaf Juice Powder, Glycerin, Propylene Glycol, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aminomethyl Propanol, Mannitol, Microcrystalline Cellulose, Sucrose, Zea Mays (Corn) Starch, Tocopheryl Acetate, Maltodextrin, Hydroxypropyl Methylcellulose, Potassium Sorbate, Sodium Benzoate, Synthetic Fluorophlogopite, Linalool, Limonene, Coumarin, Hydroxycitronellal, Benzyl Benzoate, CI 77891 (Titanium Dioxide), CI 17200 (Red 33), CI 73360 (Red 30 Lake), CI 19140 (Yellow 5).

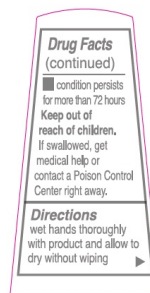
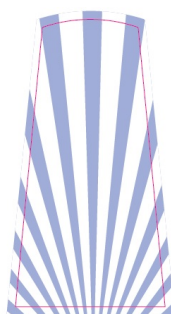


**Distributed by :** MERCI HANDY CORP. 27 WEST 20TH STREET SUITE 800 NEW YORK, NY 10011.  
www.merchihandy.com  
Made in Turkey

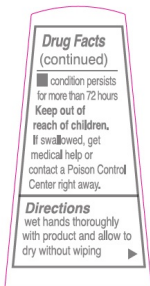
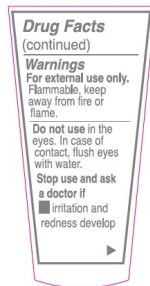


**Made with 67% alcohol and 100% love**  
**98% Natural Ingredients Aloe vera enriched formula**  
**Vegan & Cruelty-free**

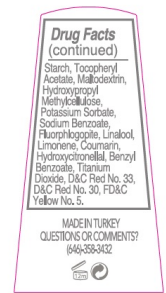
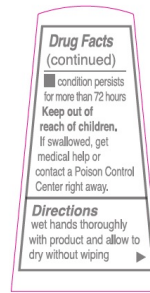
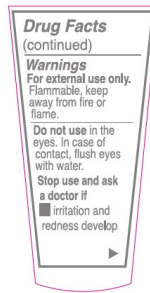
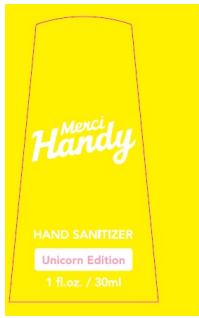
## Package Labeling:NAMASTE



## Package Labeling:GLITTER FEVER



# Package Labeling: UNICORN EDITION



## MAGIC POUCH

alcohol kit

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:72866-011
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### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72866-011-01	1 in 1 KIT	10/15/2020	06/30/2025

### Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	1 BOTTLE	30 mL
Part 2	1 BOTTLE	30 mL
Part 3	1 BOTTLE	30 mL

## Part 1 of 3

### HAND SANITIZER, NAMASTE

alcohol gel

### Product Information

<b>Item Code (Source)</b>	NDC:72866-002
<b>Route of Administration</b>	TOPICAL

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	670 mg in 1 mL

## Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>ALOE VERA LEAF</b> (UNII: ZY81Z83H0X)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>AMINOMETHYLPROPANOL</b> (UNII: LU49E6626Q)	
<b>MANNITOL</b> (UNII: 3OWL53L36A)	
<b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)	
<b>SUCROSE</b> (UNII: C151H8M554)	
<b>STARCH, CORN</b> (UNII: O8232NY3SJ)	
<b>DENATONIUM BENZOATE</b> (UNII: 4YK5Z54AT2)	
<b>.ALPHA.-TOCOPHEROL ACETATE</b> (UNII: 9E8X80D2L0)	
<b>MALTODEXTRIN</b> (UNII: 7CVR7L4A2D)	
<b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO)	
<b>POTASSIUM SORBATE</b> (UNII: 1VPU26JZZ4)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	
<b>.ALPHA.-HEXYLCINNAMALDEHYDE</b> (UNII: 7X6O37OK2I)	
<b>BENZYL BENZOATE</b> (UNII: N863NB338G)	
<b>LINALOOL, (+/-)-</b> (UNII: D81QY6I88E)	
<b>LIMONENE, (+)-</b> (UNII: GFD7C86Q1W)	
<b>FERRIC FERROCYANIDE</b> (UNII: TLE294X33A)	
<b>D&amp;C RED NO. 33</b> (UNII: 9DBA0SBB0L)	
<b>D&amp;C RED NO. 30</b> (UNII: 2S42T2808B)	
<b>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72866-002-30	30 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	10/15/2020	

## Part 2 of 3

### HAND SANITIZER, GLITTER FEVER

alcohol gel

## Product Information

<b>Item Code (Source)</b>	NDC:72866-009
<b>Route of Administration</b>	TOPICAL

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>ALCOHOL</b> (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	670 mg in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>ALOE VERA LEAF</b> (UNII: ZY81Z83H0X)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED)</b> (UNII: 59TL3WG5CO)	
<b>AMINOMETHYLPROPANOL</b> (UNII: LU49E6626Q)	
<b>MANNITOL</b> (UNII: 3OWL53L36A)	
<b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)	
<b>SUCROSE</b> (UNII: C151H8M554)	
<b>CORN</b> (UNII: 0N8672707O)	
<b>DENATONIUM BENZOATE</b> (UNII: 4YK5Z54AT2)	
<b>.ALPHA.-TOCOPHEROL ACETATE</b> (UNII: 9E8X80D2L0)	
<b>MALTODEXTRIN</b> (UNII: 7CVR7L4A2D)	
<b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO)	
<b>POTASSIUM SORBATE</b> (UNII: 1VPU26JZZ4)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	
<b>COUMARIN</b> (UNII: A4VZ22K1WT)	
<b>LIMONENE, (+)-</b> (UNII: GFD7C86Q1W)	
<b>ISOMETHYL-.ALPHA.-IONONE</b> (UNII: 9XP4LC555B)	
<b>CINNAMYL ALCOHOL</b> (UNII: SS8YOP444F)	
<b>ISOEUGENOL</b> (UNII: 5M0MWY797U)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	
<b>FD&amp;C YELLOW NO. 5</b> (UNII: I753WB2F1M)	
<b>FERRIC OXIDE RED</b> (UNII: 1K09F3G675)	
<b>FD&amp;C RED NO. 4</b> (UNII: X3W0AM1JLX)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72866-009-30	30 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			



### Part 3 of 3

## HAND SANITIZER, UNICORN EDITION

alcohol gel

### Product Information

**Item Code (Source)** NDC:72866-003

**Route of Administration** TOPICAL

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	670 mg in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
MANNITOL (UNII: 3OWL53L36A)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SUCROSE (UNII: C151H8M554)	
STARCH, CORN (UNII: O8232NY3SJ)	
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
LIMONENE, (+)- (UNII: GFD7C86Q1W)	
COUMARIN (UNII: A4VZ22K1WT)	
HYDROXYCITRONELLAL (UNII: 8SQ0VA4YUR)	
BENZYL BENZOATE (UNII: N863NB338G)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
D&C RED NO. 30 (UNII: 2S42T2808B)	

### Packaging

Marketing Start      Marketing End

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72866-003-30	30 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	10/15/2020	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	10/15/2020	06/30/2025

**Labeler** - Merci Handy Corporation (118006306)

Revised: 11/2022

Merci Handy Corporation