REFRESHIFY ANTISEPTIC HAND WIPES- alcohol
Emes Commerce Group Inc.

Refreshify Antiseptic Hand Wipes

Drug Facts

Active ingredient

Ethyl alcohol 75%

Purpose

Antiseptic

Use

for hand washing to decrease bacteria on the skin.

Warnings

For external use only
Flammable, keep away from fire or flame.

Do not use
- in the eyes

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.

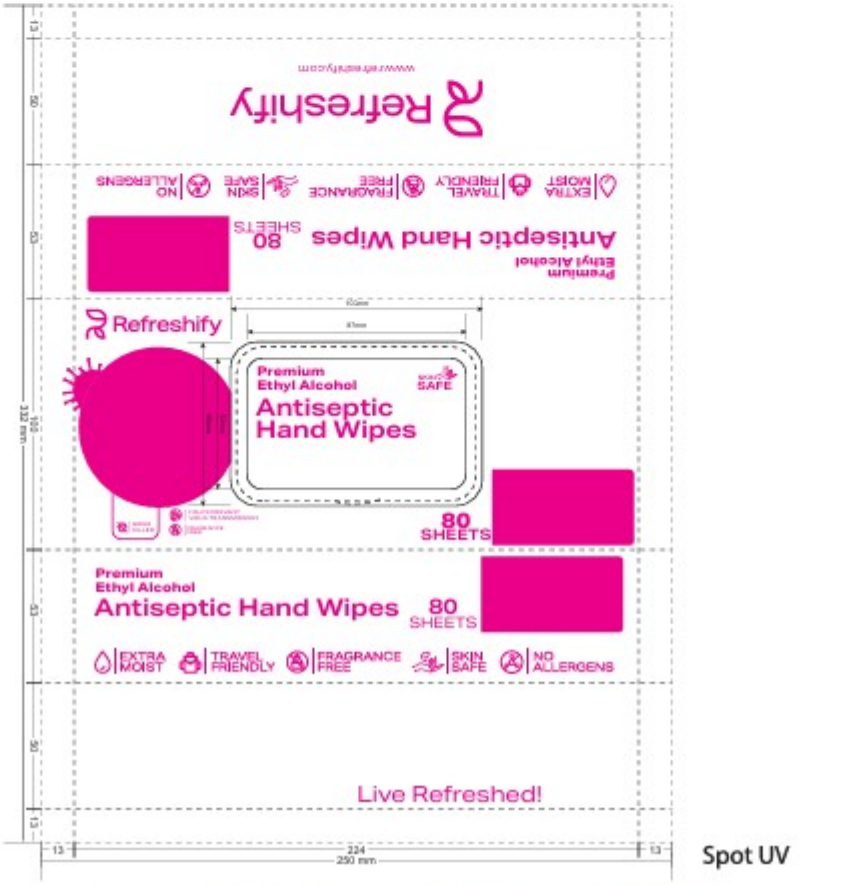
Directions

Wet hands thoroughly with wipe and allow to dry without wiping.

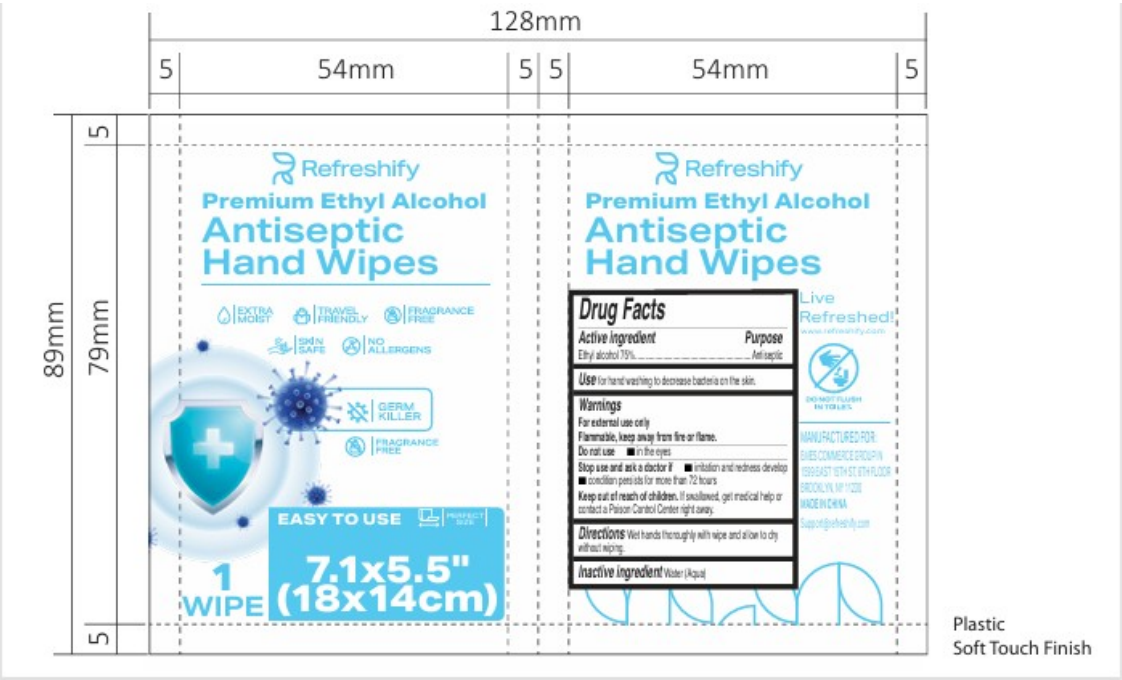
Inactive ingredient

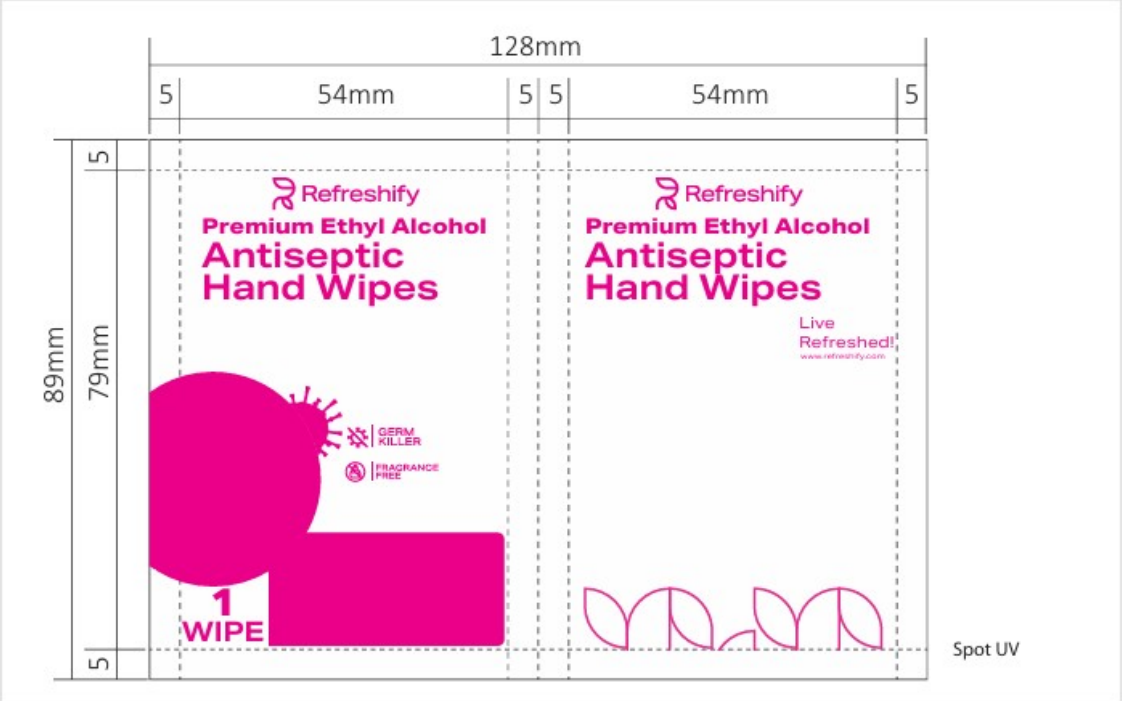
Water (Aqua)

Drug Part 80 count wipes



Drug part label 1 wipe





Kit label

REFRESHIFY ANTISEPTIC HAND WIPES
alcohol kit

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:87031-000

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:87031-000-00	1 in 1 KIT	08/11/2025	

Quantity of Parts		
Part #	**Package Quantity**	**Total Product Quantity**
Part 1	4 PATCH	17.2 mL
Part 2	4 PATCH	17.2 mL

Part 1 of 2

REFRESHIFY ANTISEPTIC HAND WIPES

alcohol cloth

Product Information	
Item Code (Source)	NDC:87031-001
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**
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:87031-001-01	80 in 1 BAG		
1		4.3 mL in 1 PATCH; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	08/11/2025	

Part 2 of 2

REFRESHIFY ANTISEPTIC HAND WIPES

alcohol cloth

Product Information

Item Code (Source)	NDC:87031-002
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
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:87031-002-02	1 in 1 BAG		
1		4.3 mL in 1 PATCH; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	08/11/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	08/11/2025	

Labeler - Emes Commerce Group Inc. (026815145)

Registrant - Emes Commerce Group Inc. (026815145)

Revised: 8/2025 Emes Commerce Group Inc.