

**SURFACE X CUTIE PALS HAND SANITIZER CARAMEL LATTE- alcohol spray
I WORLD, LLC**

Surface x Cutie Pals Hand Sanitizer, Caramel Latte

Drug Facts

Active ingredient

Ethyl Alcohol 70% v/v

Purpose

Antibacterial

Uses

- For hand-washing to help reduce bacteria on the skin.

Warnings

For external use only.

Flammable, keep away from fire and flames

When using this product

- test on a small area of skin before full use.
- keep out of eyes. In case of contact with eyes, flush thoroughly with water.
- Avoid contact with broken skin.
- Do not inhale or ingest.

Stop use and ask doctor if

- allergy, irritation or redness develops.
- Condition persists for more than 72 hours.

Keep out of reach of children.

- If product is swallowed, get medical help or contact a poison control center right away.

Directions

- Spray hands with product and allow to dry without wiping.
- For children under 6, use only under adult supervision.
- Not recommended for infants.

Other information

- Do not store above 105 °F
- May discolor some fabrics
- Harmful to wood finishes and plastics

Inactive ingredients

WATER (AQUA), FRAGRANCE (PARFUM).

Questions or comments?

1-212- 244-5170 info@amora.beauty

Package Labeling

SFC2-1010-OR

NOTE: MUST APPLY SPOT UV AND EMBOSS to artwork in separate layer "SPOT UV & EMBOSS"

HAND SANITIZER - PUMPKIN

1 fl oz/ 30 mL

BLISTER CARD OUTSIDE

HEAVY CARDSTOCK BLISTER CARD (600 GSM)
(MATTE FINISH)

BLISTER CARD INSIDE

HEAVY CARDSTOCK BLISTER CARD (600 GSM)
(MATTE FINISH)

45ml hand sanitizer 免洗洗手液
酒精浓度: 70%
香型: caramel latte
液体颜色: 透明无色

吸塑纸卡: 哑油
稿件隐藏UV 部分要过UV
反面过PET吸塑油



SURFACE X CUTIE PALS HAND SANITIZER CARAMEL LATTE

alcohol spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:85179-049
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
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:85179-049-00	45 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/22/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	04/22/2026	

Labeler - I WORLD, LLC (830590126)**Registrant** - I WORLD, LLC (830590126)

Revised: 5/2026

I WORLD, LLC