SEVEN AROMA DR- alcohol liquid Happy Seven Co., 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.

Active ingredient

Alcohol 70.0%

Purpose

Antiseptic

Uses

■ It is used by attaching the solution container to the automatic sprayer.

Warnings

Flammable, keep away from fire/flame

For external use only

Do not use

- in children less than 2 months of age
- on open skin wounds

When using this product • do not get into eyes. In case of contact, rinse eyes thoroughly with water

Stop use and ask a doctor if

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

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

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
- supervise children under 6 years of age when using this product to avoid swallowing

Other Information

- store between 1-30°C (33.8-86°F)
- avoid freezing and excessive heat above 40°C (104°F)
- may discolor certain fabrics or surfaces

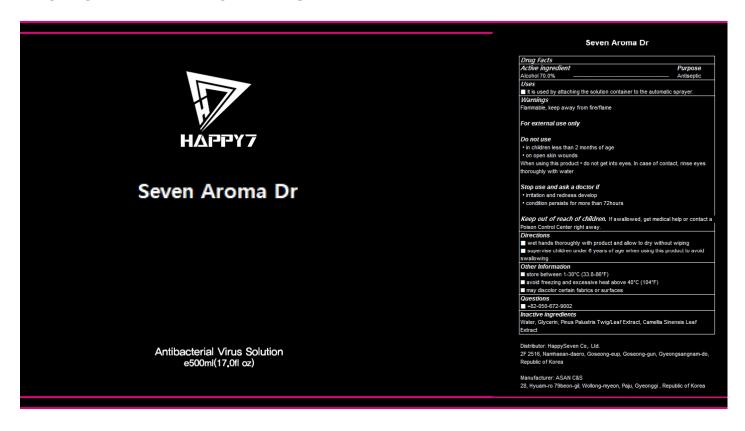
Questions

■ +82-050-672-9002

Inactive ingredients

Water, Glycerin, Pinus Palustris Twig/Leaf Extract, Camellia Sinensis Leaf Extract

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL



SEVEN AROMA DR alcohol liquid				
Product Information				
Product Type	HUMAN OTC DRUG	Item C	ode (Source)	NDC:83225-100
Route of Administration	TOPICAL			
Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength

Alcohol (UNII: 3K9958V90M) (ALCOHOL - UNII:3K995	BV90M) Alcohol	70 mL in 100 mL
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Inactive Ingredients				
Ingredient Name	Strength			
Water (UNII: 059QF0KO0R)				
Glycerin (UNII: PDC6A3C0OX)				
GREEN TEA LEAF (UNII: W2ZU1RY8B0)				

l	Packaging				
	# Item Code Package Descriptio		Package Description	Marketing Start Date	Marketing End Date
	1	NDC:83225- 100-01	500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	01/02/2023	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/02/2023	

Labeler - Happy Seven Co., Ltd. (695089951)

Registrant - Happy Seven Co., Ltd. (695089951)

Establishment				
Name	Address	ID/FEI	Business Operations	
ASAN C&S		631139649	manufacture(83225-100)	

Revised: 1/2023 Happy Seven Co., Ltd.