

**SWEET PEA SCENTED ANTIMICROBIAL HAND GEL- ethyl alcohol liquid**  
**Nantong Health & Beyond Hygienic Products Inc.**

*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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**Active ingredient**

Ethyl Alcohol, 65% v/v

**Purpose** Antimicrobial

**Uses** hand sanitizer to help  
reduce bacteria on skin

**Warnings**

**For external use only.**

**Flammable, keep away from  
fire or flame.**

**Do not use** in or near the eyes

**In case of contact**, rinse eyes  
thoroughly with water.

**Stop use and ask a doctor if**  
irritation, excessive redness  
or rash develops.

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

**Directions** Put a dime sized  
drop onto hands and rub  
together briskly until dry.

**Other information**

Store below 110° F (43° C)

**Inactive ingredients**

water, glycerin, propylene  
glycol, fragrance, carbomer,  
triethanolamine, blue 1,  
red 33

**Drug Facts**  
**Active ingredient**  
 Ethyl Alcohol, 65% v/v  
**Purpose** Antimicrobial  
**Uses** hand sanitizer to help reduce bacteria on skin ▼

PEEL UP LABEL FOR  
 ADDITIONAL DRUG FACTS



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196590 1809  
 DISTRIBUTED BY  
 GREENBRIER INTERNATIONAL, INC.  
 500 VOLVO PARKWAY,  
 CHESAPEAKE, VA 23320  
 MADE IN CHINA  
 LOT: 7380738 EXP.: 2020/07

PEEL  
 HERE

**Drug Facts (continued)**

**Warnings**  
 For external use only.

Flammable, keep away  
 from fire or flame.

Do not use in or near the eyes  
 In case of contact, rinse eyes  
 thoroughly with water.

Stop use and ask a doctor if  
 irritation, excessive redness  
 or rash develops.

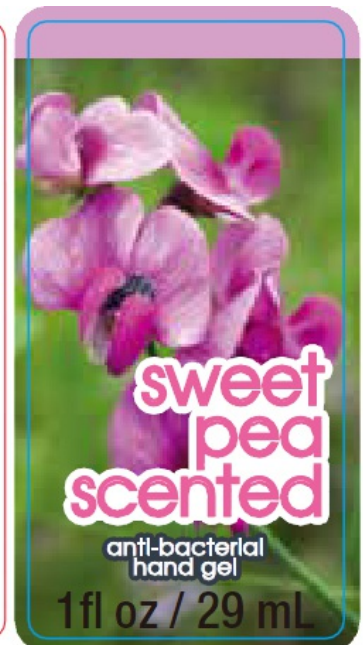
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 water, glycerin, propylene  
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 red 33



## SWEET PEA SCENTED ANTIMICROBIAL HAND GEL

ethyl alcohol liquid

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:43473-824
Route of Administration	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	65 g in 100 mL

### Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
TROLAMINE (UNII: 9O3K93S3TK)	
FD&C BLUE NO. 1 (UNII: HBR47K3TBD)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:43473-824-01	29 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/22/2018	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	03/22/2018	

**Labeler** - Nantong Health & Beyond Hygienic Products Inc. (421280161)

## Establishment

Name	Address	ID/FEI	Business Operations
Nantong Health & Beyond Hygienic Products Inc.		421280161	manufacture(43473-824)

Revised: 3/2018

Nantong Health & Beyond Hygienic Products Inc.