LA BODIES PURITA ANTI-BACTERIAL HAND SANITIZER- alcohol gel
SAMSON PHARMACEUTICAL

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.

LA Bodies™ Purita™ Anti-Bacterial Hand Sanitizer

Drug Facts

Active Ingredients

Ethyl Alcohol 62%

Purpose

Antimicrobial

Inactive Ingredients

Water (Aqua), Propylene Glycol, Aloe Barbadensis Leaf Juice, Glycerin, Tocopheryl Acetate, Carbomer, Triethanolamine, Fragrance (Parfum)

Use

Hand Sanitizer to help reduce bacteria on the skin

Warnings

- **Flammable. Keep away from fire or flame. For external use only.**
- **When using this product** do not use in or near eyes. In case of contact, rinse eyes thoroughly with water.
- **Stop use and ask doctor if** irritation or rash appears and lasts.
- **Keep out of reach of children.** If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Place product on hands. Rub until dry.

Manufactured by:
Samson Pharmaceuticals Inc.
Commerce, CA 90040

PRINCIPAL DISPLAY PANEL - 59 ml Bottle Label

KILLS 99.99%
of GERMS

MADE IN
U.S.A.

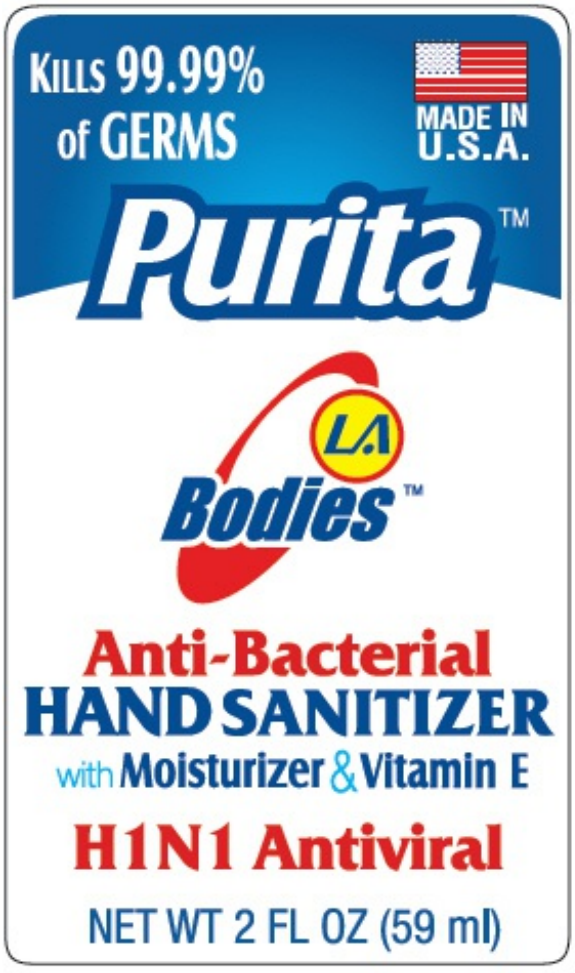
Purita™

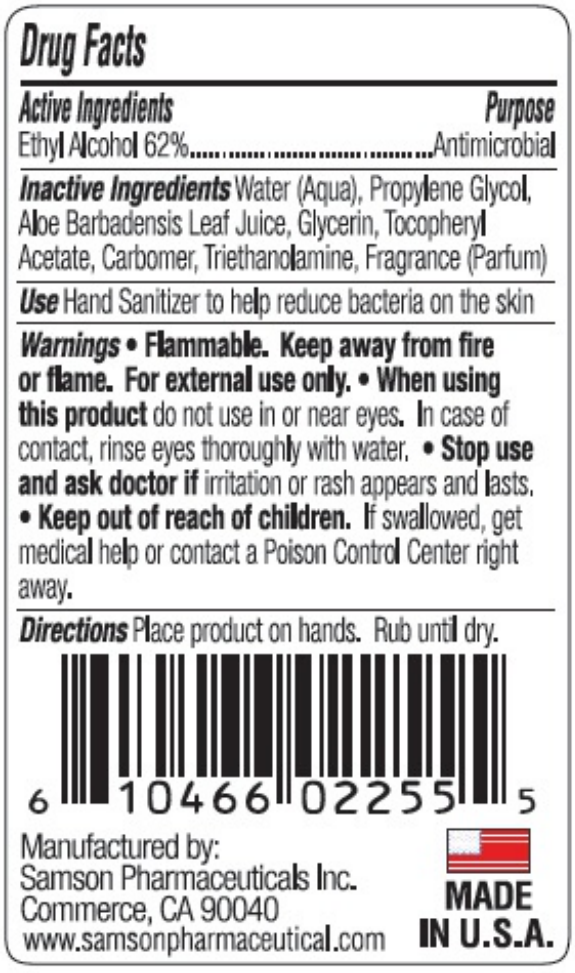
LA
Bodies™

Anti-Bacterial
HAND SANITIZER
with **Moisturizer** & **Vitamin E**

H1N1 Antiviral

NET WT 2 FL OZ (59 ml)





LA BODIES PURITA ANTI-BACTERIAL HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:20146-4009

Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Alcohol (UNII: 3K9958V90M) (Alcohol - UNII:3K9958V90M)	Alcohol	620 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
water (UNII: 059QF0KO0R)	
propylene glycol (UNII: 6DC9Q167V3)	
aloe vera leaf (UNII: ZY81Z83H0X)	
glycerin (UNII: PDC6A3C0OX)	
.alpha.-tocopherol acetate (UNII: 9E8X80D2L0)	
carbomer homopolymer type c (allyl pentaerythritol crosslinked) (UNII: 4Q93RCW27E)	
trolamine (UNII: 9O3K93S3TK)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:20146-4009-1	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/2015	
2	NDC:20146-4009-2	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/2015	
3	NDC:20146-4009-3	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/01/2015	

Labeler - SAMSON PHARMACEUTICAL (088169581)

Establishment

Name	Address	ID/FEI	Business Operations
SAMSON PHARMACEUTICAL		088169581	MANUFACTURE(20146-4009)