

NICKELODEON BRIGHT BUBBLE GUM SCENTED HAND SANITIZER 01- benzalkonium chloride liquid
Shenzhen Lantern Science 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.

NS0043TA

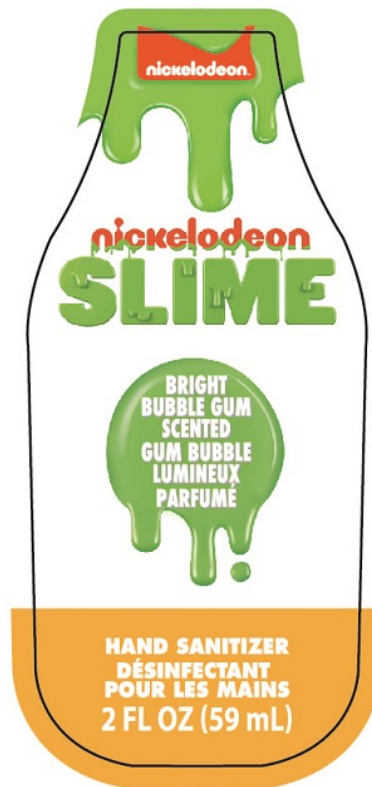
Drug facts

NS0043TA_LB
 2019/09/16 (d)

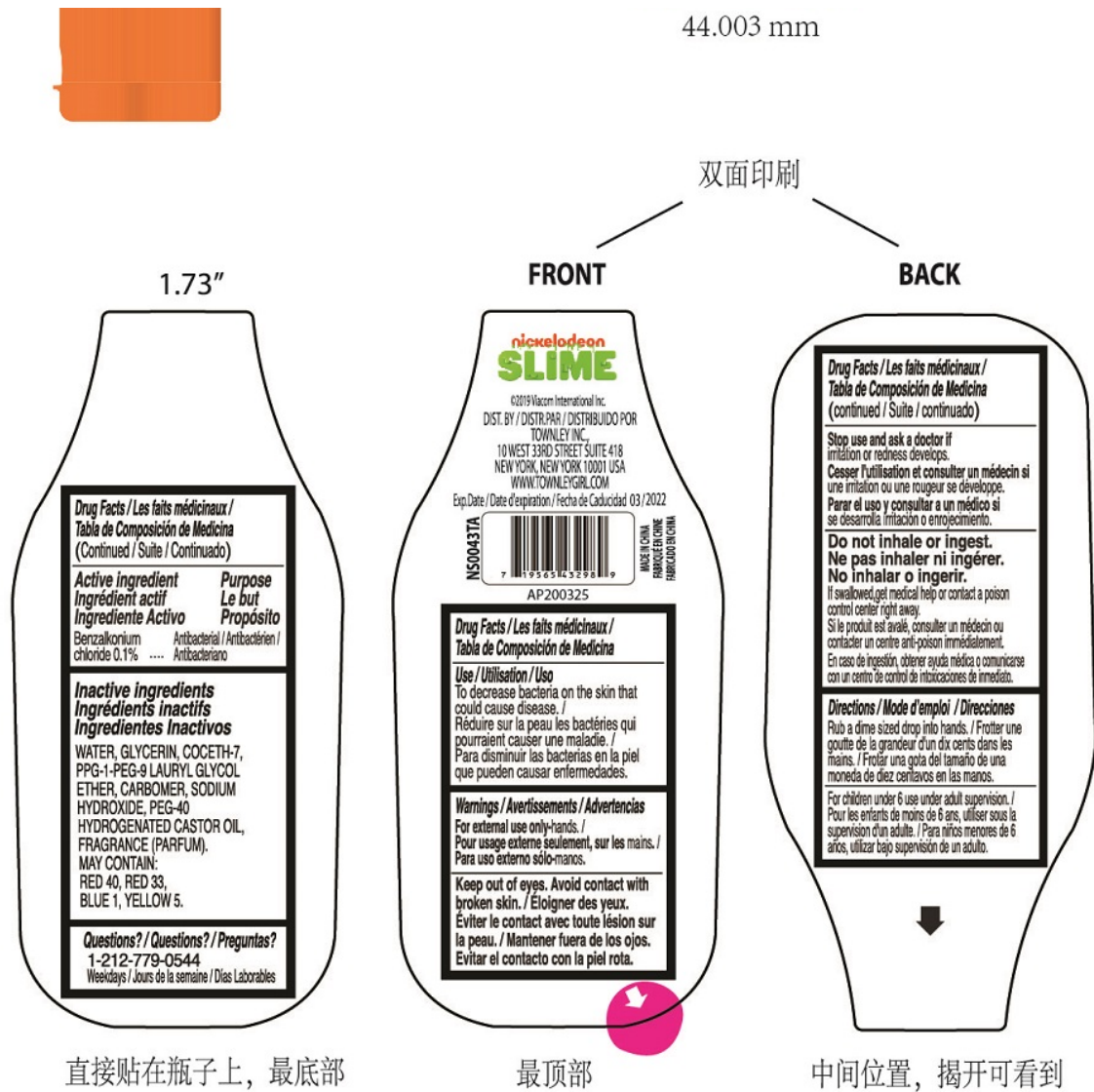
-  DIE LINE—DO NOT PRINT
-  CMYK PROCESS
-  CLEAR ADHESIVE STOCK
DO NOT PRINT
此灰色代表透明区域,不要印刷为灰色

**CONFIRM WEIGHTS
 + WARNING
 + INGREDIENTS
 + MAY CONTAINS**

Townley
 389 Fifth Avenue, Suite 1100
 New York, NY 10016
 Call (212) 779-0544 • Fax (212) 779-4192



95.288 mm



Active ingredient purpose

Benzalkonium chloride 0.1% Antibacterial

Use

To decrease bacteria on the skin that could cause disease.

Warnings

For external use only-hands

keep out of eyes- avoid contact with broken skin

Stop use and ask a doctor if irritation or redness develops

Do not inhale or ingest. If swallowed, get medical help or contact a poison control right away.

Direction

Rub a dime sized drop into hands.

For children under 6 use under adult supervision

Inactive ingredients

INACTIVE INGREDIENTS: WATER, GLYCERIN, COCETH-7, PPG-1-PEG-9 LAURYL GLYCOL ETHER, CARBOMER, SODIUM HYDROXIDE, PEG-40 HYDROGENATED CASTOR OIL, FRAGRANCE (PARFUM).

MAY CONTAIN: RED 40, RED 33, BLUE 1, YELLOW 5.

Questions?

1-212-779-0544 weekdays

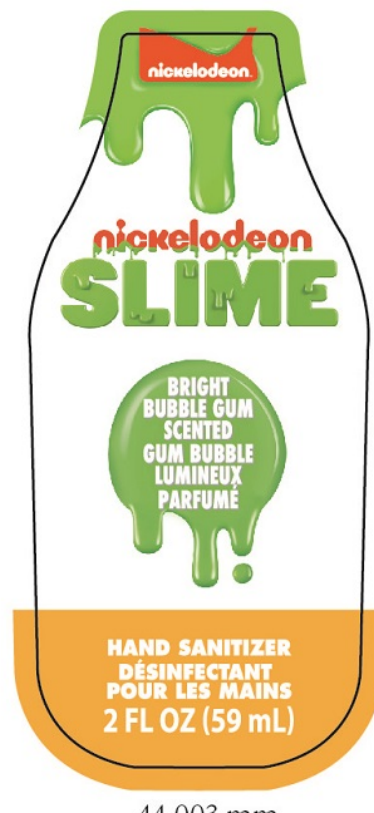
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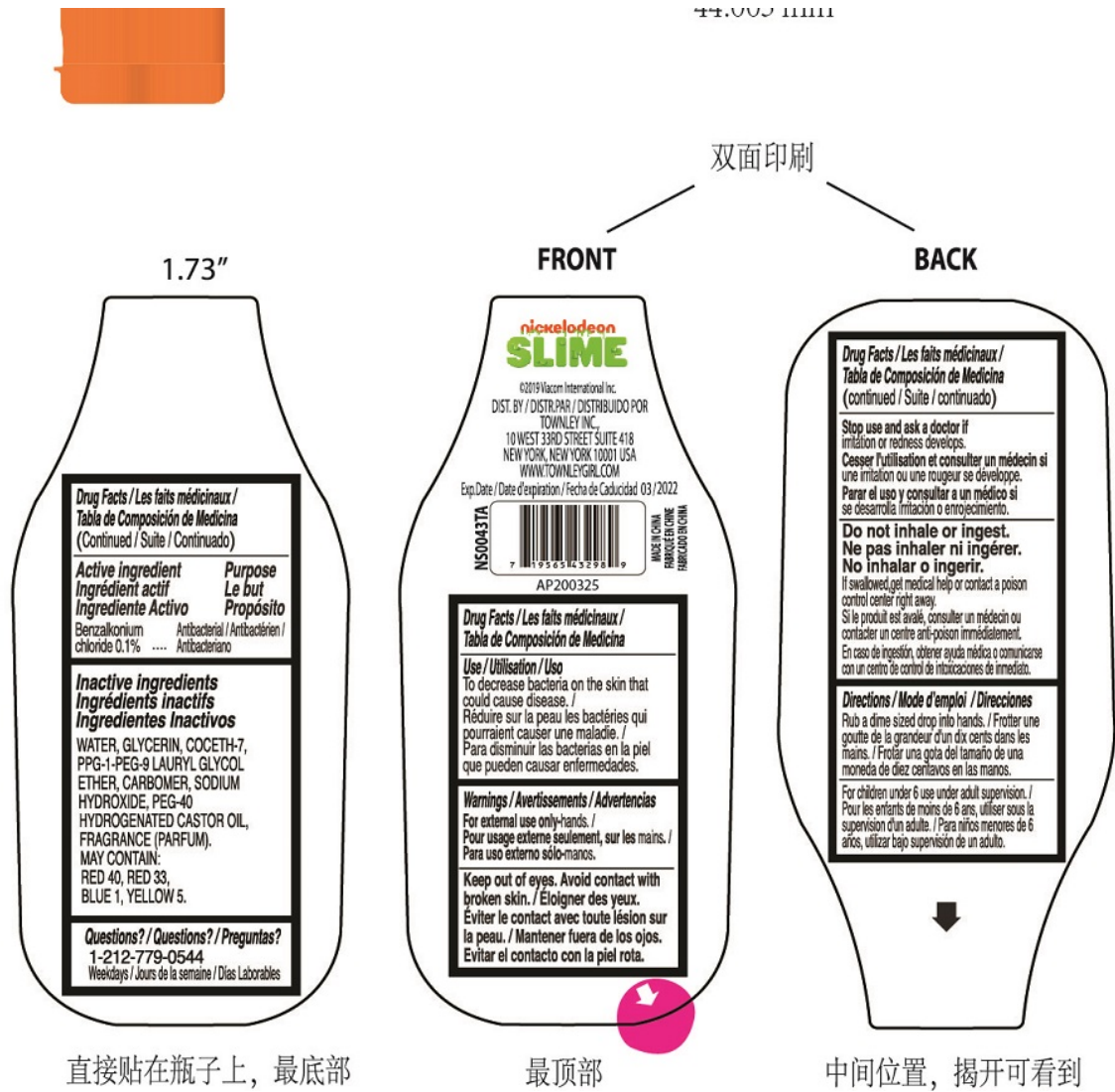
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95.288 mm

44.003 mm



直接贴在瓶子上，最底部

最顶部

中间位置，揭开可看到

NICKELODEON BRIGHT BUBBLE GUM SCENTED HAND SANITIZER 01

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54860-272
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.1 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	0.15 mL in 100 mL
COCETH-7 (UNII: 58 Y261JLH5)	0.68 mL in 100 mL
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	0.1 mL in 100 mL

MYRISTICA FRAGRANS FRUIT (UNII: GOO22BTU71)	0.3 mL in 100 mL
FD&C RED NO. 40 (UNII: WZB9127XOA)	0.1 mL in 100 mL
FD&C BLUE NO. 1 (UNII: HBR47K3TBD)	0.1 mL in 100 mL
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	0.1 mL in 100 mL
WATER O-18 (UNII: 7QV8F8BYNJ)	96 mL in 100 mL
CARBOMER 934 (UNII: Z135WT9208)	0.24 mL in 100 mL
GLYCERIN (UNII: PDC6A3C0OX)	0.68 mL in 100 mL
PPG-1-PEG-9 LAURYL GLYCOL ETHER (UNII: 5R8J43K25L)	0.68 mL in 100 mL
SODIUM HYDROXIDE (UNII: 55X04QC32I)	0.12 mL in 100 mL

Product Characteristics

Color	orange	Score	
Shape		Size	
Flavor	BUBBLE GUM	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54860-272-01	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/25/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC mono graph not final	part333A	03/25/2020	

Labeler - Shenzhen Lantern Science Co.,Ltd. (421222423)

Registrant - Lantern Beauty America,INC. (117371139)

Establishment

Name	Address	ID/FEI	Business Operations
Shenzhen Lantern Science Co.,Ltd.		421222423	manufacture(54860-272)

Revised: 5/2020

Shenzhen Lantern Science Co.,Ltd.