ANTISEPTIC HAND SANITIZER LAVENDER SCENT- alcohol spray
Two's Company, 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.

ANTISEPTIC HAND SANITIZER SPRAY-LAVENDER SCENT

Drug Facts

Active ingredient

Alcohol 62%

Purpose

Antimicrobial

Uses

• helps to reduce bacteria on the skin

• recommended for repeated use

Warnings

For external use only

Flammable: Keep away from heat and flame

When using this product
• do not use in or near the eyes. In case of eye contact, rinse eyes thoroughly with water
• do not apply to irritated or broken skin.

Stop use and ask a doctor if
• irritation and redness develop
• condition persists for more than 72 hours.

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

Directions

• lift tab and spray a small amount into the palms of your hands and forearms.

• wet the hands thoroughly with product, rub hands together and allow to dry without wiping • no rinsing required.

• children under 6 years of age should be supervised when using this product • not recommended for infants.

Other information

• do not store above 105°F

• may discolor some fabrics or surfaces.

Inactive ingredients

Fragrance, Glycerin, Propylene Glycol, Water

LAVENDER SCENT

BORN TO GARDEN FORCED TO WORK

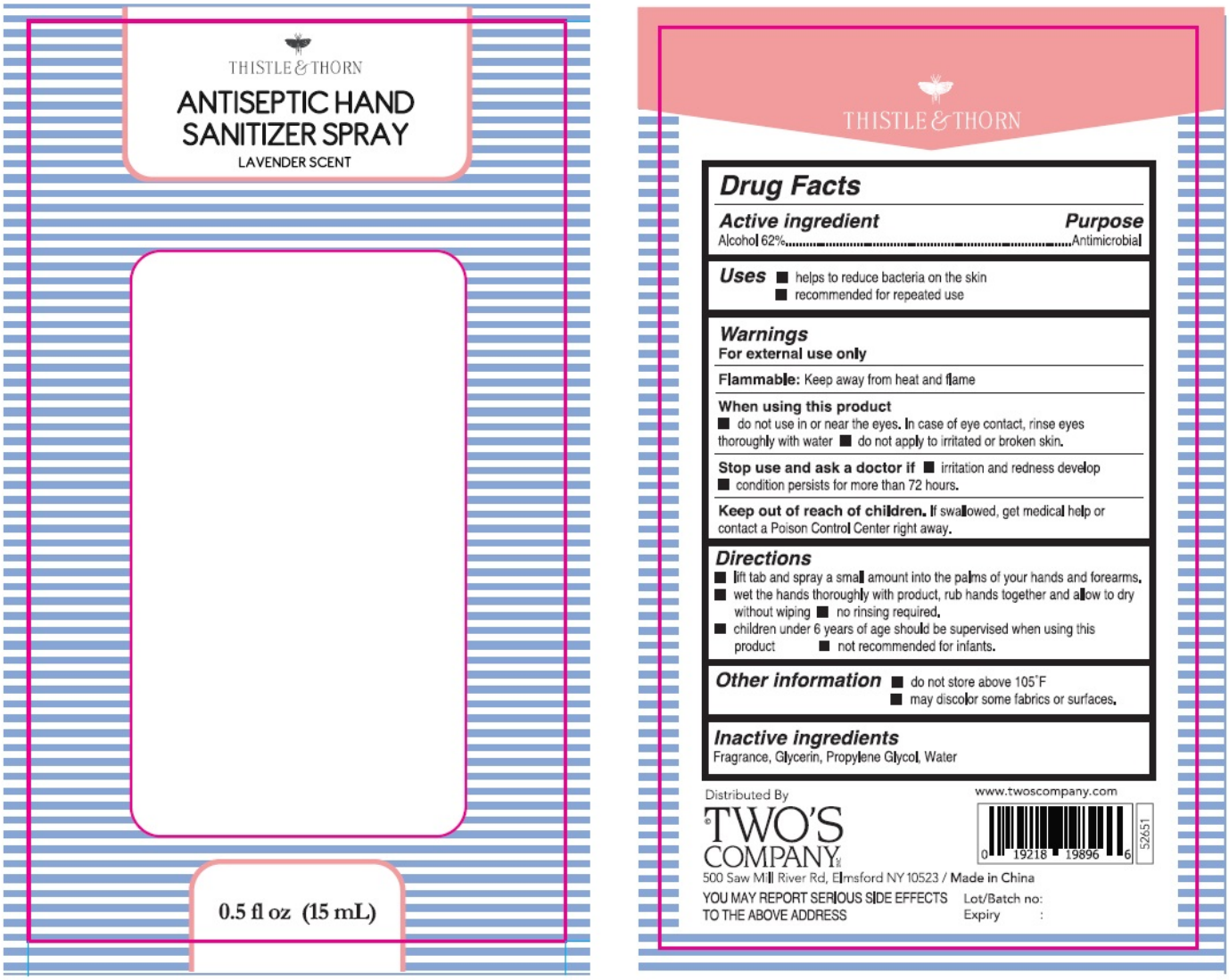
CLEANS AND FRESHENS YOUR HANDS AND REDUCES BACTERIA

Distributed By
TWO'S COMPANY INC.
500 Saw Mill River Rd, Elmsford, NY 10523 / **Made in China**

YOU MAY REPORT SERIOUS SIDE EFFECTS TO THE ABOVE ADDRESS

Packaging

OUTER PACK LABEL



INNER LABEL

ANTISEPTIC HAND SANITIZER LAVENDER SCENT

alcohol spray

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72762-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.62 mL in 1 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72762-001-16	1 in 1 BLISTER PACK	09/01/2018	
1	NDC:72762-001-15	15 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	09/01/2018	

Labeler - Two's Company, Inc. (056307960)

Revised: 1/2019 Two's Company, Inc.