HUGO NATURALS VANILLA PEPPERMINT HAND SANITIZER- ethanol spray
Dm Natural 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.

HUGO Naturals VANILLA PEPPERMINT HAND SANITIZER

Drug Facts

Active Ingredients

Ethanol* 62%

Purpose

Antimicrobial

Uses Helps reduce bacteria on the skin. Recommended for repeated use.

Warnings For external use only. Flammable, keep away from heat or flame. Do not use in or near eyes. In case of contact, rinse eyes thoroughly with water. Stop use and ask a doctor if irritation develops and persists for more than 72 hours.

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

Directions Spray in your palm and rub thoroughly to cover both hands. Briskly rub hands together until dry. Supervise children under 6 when using this product.

Inactive Ingredients: Aqua/Water, Mentha Piperita (Peppermint) Oil, Glycerin, Natural Fragrance Oils, Allantoin, Hamamelis Virginiana (Witch Hazel) Leaf Extract, Lavandula Angustifolia (Lavender) Oil, Aloe Barbadensis Leaf Juice.*

*Certified Organic

Moisturizing

Natural

Sulfate-free

Gluten-free

Soy-free

No Animal Testing

Recyclable

MADE IN THE USA for:

DM Natural Products, Inc.

Chatsworth, CA 91311

HugoNaturals.com

Packaging

HUGO NATURALS VANILLA PEPPERMINT HAND SANITIZER

ethanol spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70279-610
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
PEPPERMINT OIL (UNII: AV092KU4JH)	
GLYCERIN (UNII: PDC6A3C0OX)	

ALLANTOIN (UNII: 344S277G0Z)	
HAMAMELIS VIRGINIANA LEAF (UNII: T07U1161SV)	
LAVENDER OIL (UNII: ZBP1YXW0H8)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70279-610-02	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/29/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	12/29/2015	

Labeler - Dm Natural Products, Inc. (623230898)

Revised: 7/2017 Dm Natural Products, Inc.