HAND SANITIZER WIPE- alcohol cloth HAND SANITIZER- alcohol liquid Duro-Last

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.

This is a hand sanitizer manufactured according to the Temporary Policy for Preparation of Certain Alcohol-Based Hand Sanitizer Products During the Public Health Emergency (CoViD-19); Guidance for Industry.

The hand sanitizer is manufactured using only the following United States Pharmacopoeia (USP) grade ingredients in the preparation of the product (percentage in final product formulation) consistent with World Health Organization (WHO) recommendations:

- a. Alcohol (ethanol) (USP or Food Chemical Codex (FCC) grade) (80%, volume/volume (v/v)) in an aqueous solution denatured according to Alcohol and Tobacco Tax and Trade Bureau regulations in 27 CFR part 20.
- b. Glycerol (1.45% v/v).
- c. Hydrogen peroxide (0.125% v/v).
- d. Sterile distilled water or boiled cold water.

The firm does not add other active or inactive ingredients. Different or additional ingredients may impact the quality and potency of the product.

Active Ingredient(s)

Alcohol 80% v/v. Purpose: Antiseptic

Purpose

Antiseptic, Hand Sanitizer

Use

Hand Sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

Warnings

For external use only. Flammable. Keep away from heat or flame

Do not use

- in children less than 2 months of age
- on open skin wounds

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition. Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

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Directions

- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

Other information

- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients

glycerin, hydrogen peroxide, purified water USP

Package Label - Principal Display Panel



DURO-LAST® HAND SANITIZER

Alcohol Antiseptic 80% Topical Solution Non-Sterile Solution 1 Gallon (128 FL OZ)



Drug Facts

Active ingredient(s)

Purpose

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.Antiseptic

Use(s)

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Inactive ingredients glycerin, hydrogen peroxide, purified water USP



Manufactured By: **Anvil Paints & Coatings, Inc.** 1255 Starkey Rd Largo, FL 33771 www.anvilpaints.com

3.8L NDC: 77378-532-01

2.5gal NDC: 77378-533-01



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DURO-LAST® SANITIZING WIPES

Alcohol Antiseptic 80% Topical Solution Non-Sterile Solution 2.5 Gallon (320 FL OZ)

Net Content: 75 Wipes - 8" x 18"



Drug Facts

Active ingredient(s)

Sanitizing Wipes used to help reduce bacteria that pot water are not available. Intended to be used on washa

Luse only. Flammable. Keep away from he Do not use

On children less than 2 months of age

On open skin wounds
When using this product keep out of eyes, ears, and thoroughly with water.

Stop use and ask a doctor if irritation or rash occurs,

Keep out of reach of children. If swallowed, get med

- Use this product on washable hard, nonporous surface For soiled surfaces: Use wipe to first remove grime at to sanitize surface.
 To sanitize: use an individual wipe on nonporous surface.
- Discard used wipe in the trash, wipes are not meant t

Other information Store between 15-30°C (59-86°F)

Avoid freezing and excessive heat above 40°C (104°F)

Inactive ingredients glycerin, hydrogen peroxide, pur



Manufactured B Anvil Paints & Coati

5.0gal NDC: 77378-533-02



DURO-LAST® SANITIZING WIPES

Alcohol Antiseptic 80% Topical Solution Non-Sterile Solution 5 Gallon (640 FL OZ)

Net Content: 75 Wipes - 12" x 24"



Drug Facts

Active ingredient(s) Alcohol 80% v/v.....

Purpose

Sanitizing Wipes used to help reduce bacteria that potentially can cause disease. For use when soap and water are not available. Intended to be used on washable hard, nonporous surfaces

Warnings

For external use only. Flammable. Keep away from heat or flame.

Do not use

On children less than 2 months of age

• On open skin wounds

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition. Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center

Directions

- Use this product on washable hard, nonporous surfaces.
 For soiled surfaces: Use wipe to first remove grime and debris. Discard and use additional wipe
- to sanitize surface.

 To sanitize: use an individual wipe on nonporous surface until visibly wet. Let air dry.
- Discard used wipe in the trash, wipes are not meant to be re-used.

Other information
- Store between 15-30°C (59-86°F)

· Avoid freezing and excessive heat above 40°C (104°F)

Inactive ingredients glycerin, hydrogen peroxide, purified water USP



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HAND SANITIZER WIPE

alcohol cloth

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:77378-533(NDC:76559-533)

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	80 mL in 100 mL

Inactive Ingredients	
Ingredient Name	Strength
GLYCERIN (UNII: PDC6 A3C0 OX)	1.45 mL in 100 mL
HYDRO GEN PERO XIDE (UNII: BBX060AN9V)	0.125 mL in 100 mL
WATER (UNII: 059QF0KO0R)	

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:77378-533- 01	9464 mL in 1 CONTAINER; Type 0: Not a Combination Product	05/06/2020	
2 NDC:77378-533- 02	18927 mL in 1 CONTAINER; Type 0: Not a Combination Product	05/06/2020	

Marketing Infor	mation		
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/06/2020	

HAND SANITIZER

alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77378-532(NDC:76559-532)
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	80 mL in 100 mL

Inactive Ingredients	
Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	1.45 mL in 100 mL
HYDRO GEN PERO XIDE (UNII: BBX060AN9V)	0.125 mL in 100 mL
WATER (UNII: 059QF0KO0R)	

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Packaging		

#	Item Code	Package Description	Date	Date
1	NDC:77378-532- 01	3800 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/06/2020	
N	Tarketing In	formation		
	Tarketing In		Aarketing Start Date	Marketing End Date
		ory Application Number or Monograph Citation M	Marketing Start Date	Marketing End Date

Labeler - Duro-Last (065173064)

Revised: 5/2020 Duro-Last