CLEAN ANTISEPTIC HAND SANITIZER WIPES- ethyl alcohol cloth
Raw Office Inc, The

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.

CLEAN ANTISEPTIC HAND SANITIZER wipes

Drug Facts

Active Ingredient:

Ethyl Alcohol 75% (v/v)

Purpose

Antimicrobial

Uses

Hand sanitizer to help reduce bacteria on skin

Warnings

■ External use, not oral. Keep out of children's reach.

■ Flammable, keep away from fire and flame.

■ Use with caution if allergic to alcohol.

When using this product

Avoid contacting face, eyes and broken skin. In case of eye contact, flush with plenty of water and seek medical advice.

Stop use and ask doctor if

■ Irritation or redness occurs.

Keep out of children's reach.

If swallowed, get medical help or contact poison control center right away.

Directions

Wet hands thoroughly with product and allow to dry.

Other Information

■ Store below 43°C (110°F)

■ Keep sealed after use

Inactive IngredientsPurified Water, Glycerol, Aloe Vera

Product Use:

Hold and grasp the barrel with both hands then turn the lid counter-clockwise. Take out the wet towel bag, tear the package parallel from the tear hole, pull towel through the centre hole of the barrel lid. Make sure to close lid between uses.

MADE IN CHINA

Packaging

DRUG FACTS LABEL

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Batch No:
Production Date:
Expiry Date:

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MADE IN CHINA
NDC # 75353-002-01 6 28011 26202 8

CLEAN ANTISEPTIC HAND SANITIZER WIPES

ethyl alcohol cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:75353-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75353-002-01	100 in 1 PACKET	08/11/2020	
1		5.4 mL in 1 PACKAGE; 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	part333A	08/11/2020	

Labeler - Raw Office Inc, The (204127000)

Revised: 7/2023 Raw Office Inc, The