REDI WIPES- benzalkonium chloride patch
Radienz Living, LLC

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.

Redi Wipes Antibacterial Hand Wipes

Active ingredients

Benalkonium chloride (0.115%)

Purpose

Antimicrobial

Uses

- For hand sanitizing to decrease bacteria on the skin.
- Recommended for repeated use.

Warnings

For external use only.

Do Not Use

Do not use over large areas of the body or if you are allergic to any of these ingredients.

When using this product

Avoid contact with eyes. If contact occurs, rinse eyes thoroughly with water.

Stop use and ask a doctor if

If irritation or redness develops and 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:

- To use, peel back label, pull out wipe and firmly reseal label to maintain package moisture. Do not remove label.

- Unfold cloth and rub thoroughly over all surfaces of both hands. Use only once and throw away. After cleansing, dispose of wipe in a trash receptacle. Rub hands together briskly to dry.

Inactive ingredients

Water, Behentrimonium Chloride, Dihydroxypropyl Peg-5 Linoleammonium Chloride, Glycereth-2 Cocoate, Aloe Barbadensis Leaf Extract, Tocopheryl Acetate (Vitamin E), Phenoxyethanol, Potassium Sorbate, Fragrance

Other Information

*Kills 99.9% of germs that may cause illness, including E. Coli and MRSA in just 15 seconds.

**This product is not manufactured or distributed by Edgewell Personal Care, owner of the registered trademark, Wet Ones®.

DISTRIBUTED BY: U.S. NONWOVENS CORP. 100 EMJAY BLVD., BRENTWOOD, NY 11717

Principal Display Panel

SANITIZING REDI WIPES™

Compare to Wet Ones®**

ANTIBACTERIAL HAND WIPES

FRESH SCENT

ALCOHOL FREE

KILLS 99.9% OF GERMS*

15 Soft Wipes

7 IN x 6 IN

(17.78 cm x 15.24 cm)

REDI WIPES

benzalkonium chloride patch

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:52553-004
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.115 mg in 100 mg

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
BEHENTRIMONIUM CHLORIDE (UNII: X7GNG3S47T)	
DIHYDROXYPROPYL PEG-5 LINOLEAMMONIUM CHLORIDE (UNII: 0Y0NQR2GH1)	
GLYCERETH-2 (UNII: 0UY833L6XU)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52553-004-15	15 in 1 PACKAGE	03/16/2021	
1		0.115 mg in 1 PATCH; Type 1: Convenience Kit of Co-Package		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/16/2021	

Labeler - Radienz Living, LLC (117451061)

Registrant - Radienz Living, LLC (080453184)

Establishment

Name	Address	ID/FEI	Business Operations
Radienz Living, LLC		080453184	manufacture(52553-004)

Revised: 4/2021 Radienz Living, LLC