HAND WIPES ANTIBACTERIAL- benzalkonium chloride cloth
Hannaford Brothers Company

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.

Drug Facts

Active Ingredient

Benzalkonium Chloride 0.115%

Purpose

Antibacterial

Uses

For handwashing to decrease bacteria on the skin

Warnings

For external use only

Do not use

- in the eyes
- if you are allergic to any of the ingredients

When using this product

if eye contact occurs, rinse eyes thoroughly with water

Stop use and ask a doctor

if irritation and redness develop and persist 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

- **Storage.** Store at room temperature
- **Dispensing.** Remove seal band. Lift tab at front of lid. Pull up corner of center sheet and thread through dispenser slit in lid. Close lid to retain moisture
- **Use.** Apply wipe thouroughly to hands as desired. Allow to dry without wiping.
- **Disposal.** Dispose of used wipes in trash recepacle after use. Do not flush.

Inactive Ingredients

Water, Aloe Barbadensis Leaf Extract, Disodium Cocoamphodiacetate, SD Alcohol 40-B,

Methylchloroisothiazolinine, Methylisothiazolinone, Tocopheryl Acetate, Disodium EDTA, Citric acid, Fragrance

Package Label

HAND WIPES ANTIBACTERIAL

benzalkonium chloride cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:41268-600

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 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
DISODIUM COCOAMPHODIACETATE (UNII: 18L9G3U51M)	
ALCOHOL (UNII: 3K9958V90M)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
EDETATE SODIUM (UNII: MP1J8420LU)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41268-600-40	40 in 1 PACKAGE; Type 0: Not a Combination Product	12/01/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	12/01/2017	

Labeler - Hannaford Brothers Company (006949556)

Registrant - Rockline Industries (966920881)

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