READYBATH LUXE- benzalkonium chloride cloth
Medline Industries, LP

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.

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180 Readybath LUXE (scented)

Active ingredient

Benzalkonium chloride 0.12%

Purpose

Antiseptic

Uses

- for body cleansing to decrease bacteria on the skin
- helps kill germs that can cause odor

Warnings

For external use only

Do not use

- in the eyes

Ask a doctor before use if you have

- deep or puncture wounds
- animal bites
- serious burns

Stop use and ask a doctor if

- irritation and redness develop
- the condition persists for more than 72 hours or gets worse

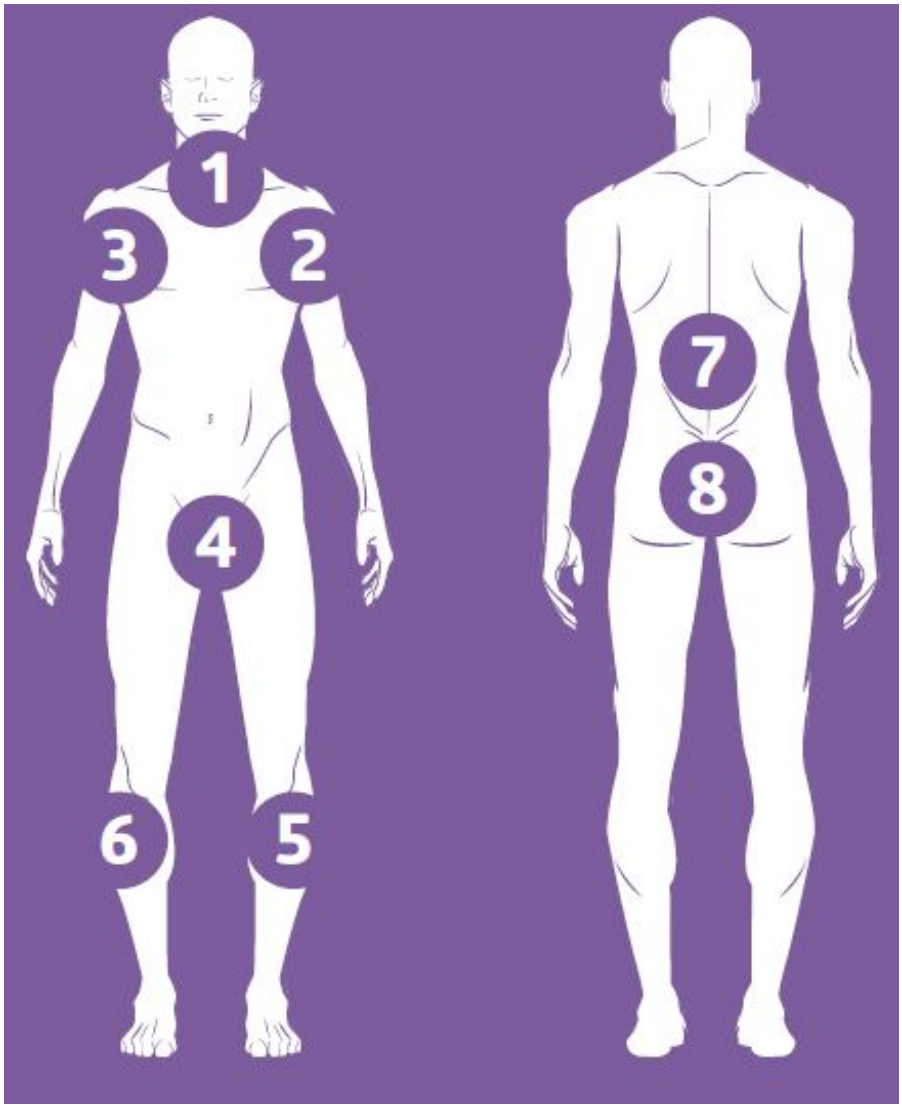
Keep out of reach of children.

If swallowed get medical help or contact a Poison Control Center right away.

Directions

- gently pull label tab until package opening is exposed
- remove cloths one at a time to cleanse body parts in the order of the diagram
- allow skin to dry
- dispose of cloth in waste receptacle

- do not flush cloths in toilet



Other information

- avoid excessive heat, protect from freezing
- washcloth: rayon/polyester

Inactive ingredients

water (aqua), cocamidopropyl PG-dimonium chloride phosphate, glycerin, phenoxyethanol, benzoic acid, dehydroacetic acid, ethylhexylglycerin, disodium EDTA, polysorbate 20, sodium citrate, fragrance, simethicone, aloe barbadensis leaf juice

Manufacturing Information

Manufactured by:

Medline Industries, LP

Three Lakes Drive, Northfield, IL 60093 USA

Made in USA with domestic and foreign materials

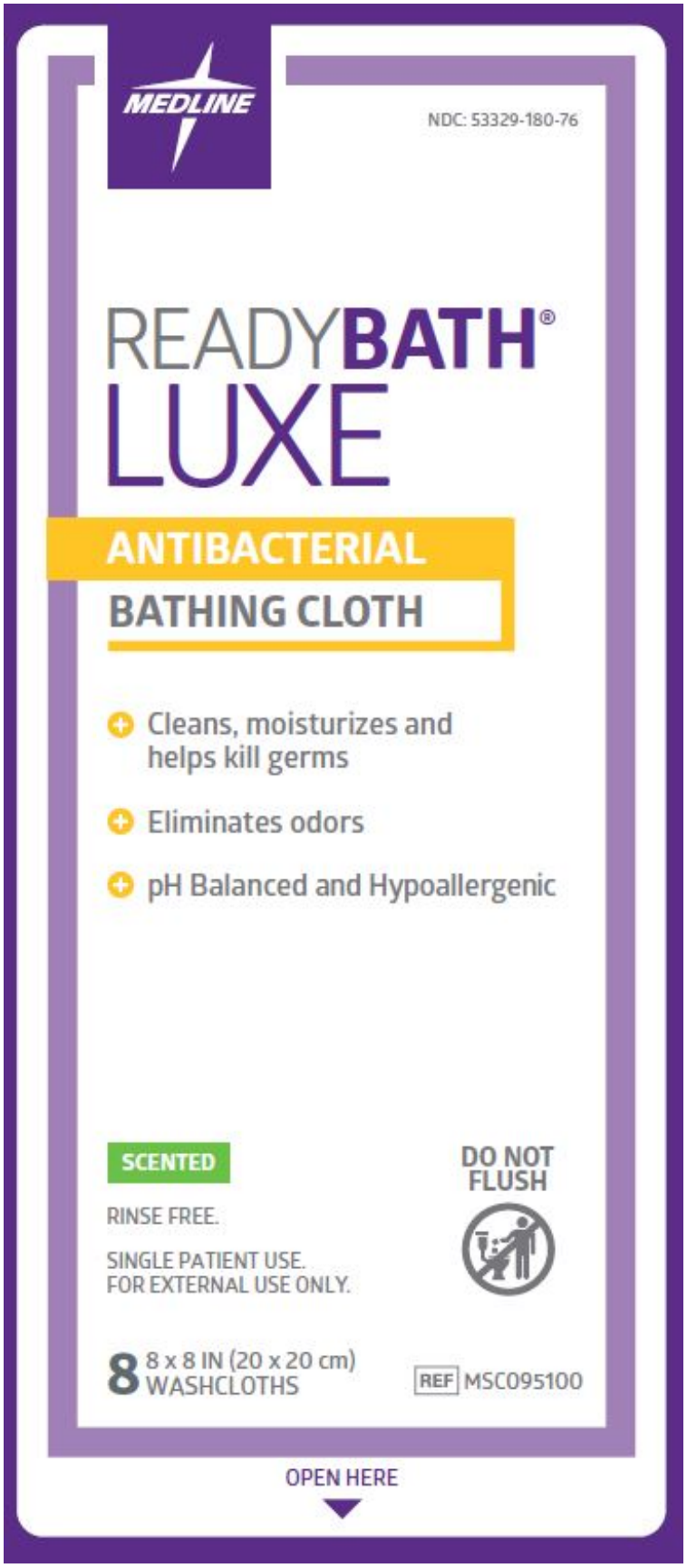
www.medline.com

1-800-MEDLINE (633-5463)

REF: MSC095100

V1 RK21DYN

Package Label



READYBATH LUXE

benzalkonium chloride cloth

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:53329-180
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.12 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
DEHYDROACETIC ACID (UNII: 2KAG279R6R)	
COCAMIDOPROPYL PG-DIMONIUM CHLORIDE PHOSPHATE (UNII: H2KVQ74JM4)	
BENZOIC ACID (UNII: 8SKN0B0MIM)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53329-180-76	129 g in 1 PACKAGE; Type 0: Not a Combination Product	05/01/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/01/2016	

Labeler - Medline Industries, LP (025460908)

Registrant - Medline Industries, LP (025460908)

Revised: 3/2022 Medline Industries, LP