TARGET UP AND UP PAIN RELIEVING CLEANSING- benzalkonium chloride and lidocaine hydrochloride spray
TARGET CORPORATION

Target Up and Up Pain Relieving Cleansing Spray

Active Ingredients

Benzalkonium Cl 0.13%

Lidocaine HCL 2.5%

Purpose

First aid antiseptic

Pain relieving spray

Uses

first aid to help prevent bacterial contamination or skin infection and for temporary relief of pain and itching associated with minor:

- cuts
- scrapes
- minor burns
- sunburn
- minor skin irritations

Warnings

For external use only

Ask a doctor before use if you have

- deep or puncture wounds
- animal bites
- serious burns

When using this product

- do not use in or near the eyes
- do not apply over large areas of the body or in large quantities
- do not apply over raw surfaces or blistered areas

Stop use and ask a doctor if

- condition worsens

- symptoms persist for more than 7 days, or clear up and occur again within a few days

Keep out of reach of children.

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

Directions

- adults and children 2 years and older:
- clean the affected area; apply a small amount on the area 1-3 times daily
- may be covered with a sterile bandage
- if bandaged, llet dry first
- children under 2 years, ask a doctor

Other information

- avoid excessive heat
- store at room temperature

Inactive ingredients

alcohol, disodium EDTA, fragrance, propylene glycol, purified water, sodium hydroxide

Label

TARGET UP AND UP PAIN RELIEVING CLEANSING

benzalkonium chloride and lidocaine hydrochloride spray

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-252
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 mL
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	2.5 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

EDETATE DISODIUM (UNII: 7FLD91C86K)	
ALCOHOL (UNII: 3K9958V90M)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-252-21	150 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/20/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	11/20/2023	

Labeler - TARGET CORPORATION (006961700)

Revised: 1/2024

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