

BENZALKONIUM CHLORIDE- antibacterial hand wipes cloth
Target Corporation

Antibacterial Hand Wipes

Active ingredient

Benzalkonium Chloride 0.13%

Purpose

Antimicrobial

Uses

- for hand sanitizing to decrease bacteria on the skin
- recommended for repeated use

Warnings

For external use only.

Do not use

if you are allergic to any of these ingredients.

When using this product

avoid contact with eyes. In case of contact, flush eyes with water.

Stop use and ask a doctor

if irritation or redness develops, or if condition persists for more than 72 hours.

Keep out of reach of children.

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

Directions

- adults and children 2 years and over: apply to hands; allow to dry without wiping.
- Children under 2 years: ask a doctor before use.
- **To dispense:** Peel back front label slowly. Remove wipes.
- **To reseal:** Firmly rub thumb over label. Dispose of wipe in trash. Do not flush.

Other information

- store below 95°F (35°C)
- keep closed tightly

- may discolor certain fabrics or surfaces.

Inactive ingredients

Water, Phenoxyethanol, Decyl Glucoside, Disodium EDTA, Sodium Benzoate, Potassium Sorbate, Vitamin E, Aloe Barbadosensis Leaf Extract, Citric Acid, Fragrance

antibacterial

hand wipes

plant-based wipes

dermatologist tested

hypoallergenic

25 HAND WIPES

5½ IN X 8 IN

Drug Facts	
Active ingredient	Purpose
Benzalkonium Chloride 0.13%	Antimicrobial
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antibacterial hand wipes

kills 99.9% of common germs**

FRESH SCENT 25 COUNT

plant-based wipe
dermatologist tested
hypoallergenic

up&up

DO NOT FLUSH

25 HAND WIPES
5½ IN X 8 IN (13.9 cm X 20.3 cm)

up & up™ antibacterial hand wipes provide a convenient way to get rid of dirt and germs quickly, leaving your skin feeling fresh, soft and less dry. Use any time to clean and refresh.

Dist. by Target Corporation
Minneapolis, MN 55403
Assembled in U.S.A.
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**Testing based on a subset of FDA proposal published June 17, 1994.

FSC
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The mark of responsible forestry

how2recycle.info
MULTI-LAYER
POUCH

253 04 3911 R02
C-002016-01-021

8 10116 71117 8

40 HAND WIPES

5½ IN X 8 IN

Drug Facts

Active ingredient Benzalkonium Chloride 0.13%...**Purpose** Antimicrobial

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• recommended for repeated use

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up&up **antibacterial** hand wipes

DO NOT FLUSH

Kills 99.9% of common germs**

40 WIPES PER POUCH

40 HAND WIPES
5½ IN X 8 IN (13.9 cm X 20.3 cm)

antibacterial hand wipes

plant-based wipe dermatologist tested hypoallergenic

up&up

DO NOT FLUSH

Kills 99.9% of common germs**

40 WIPES PER POUCH

antibacterial hand wipes

up&up

Kills 99.9% of common germs**

40 WIPES PER POUCH

DO NOT FLUSH

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C-002016-01-021

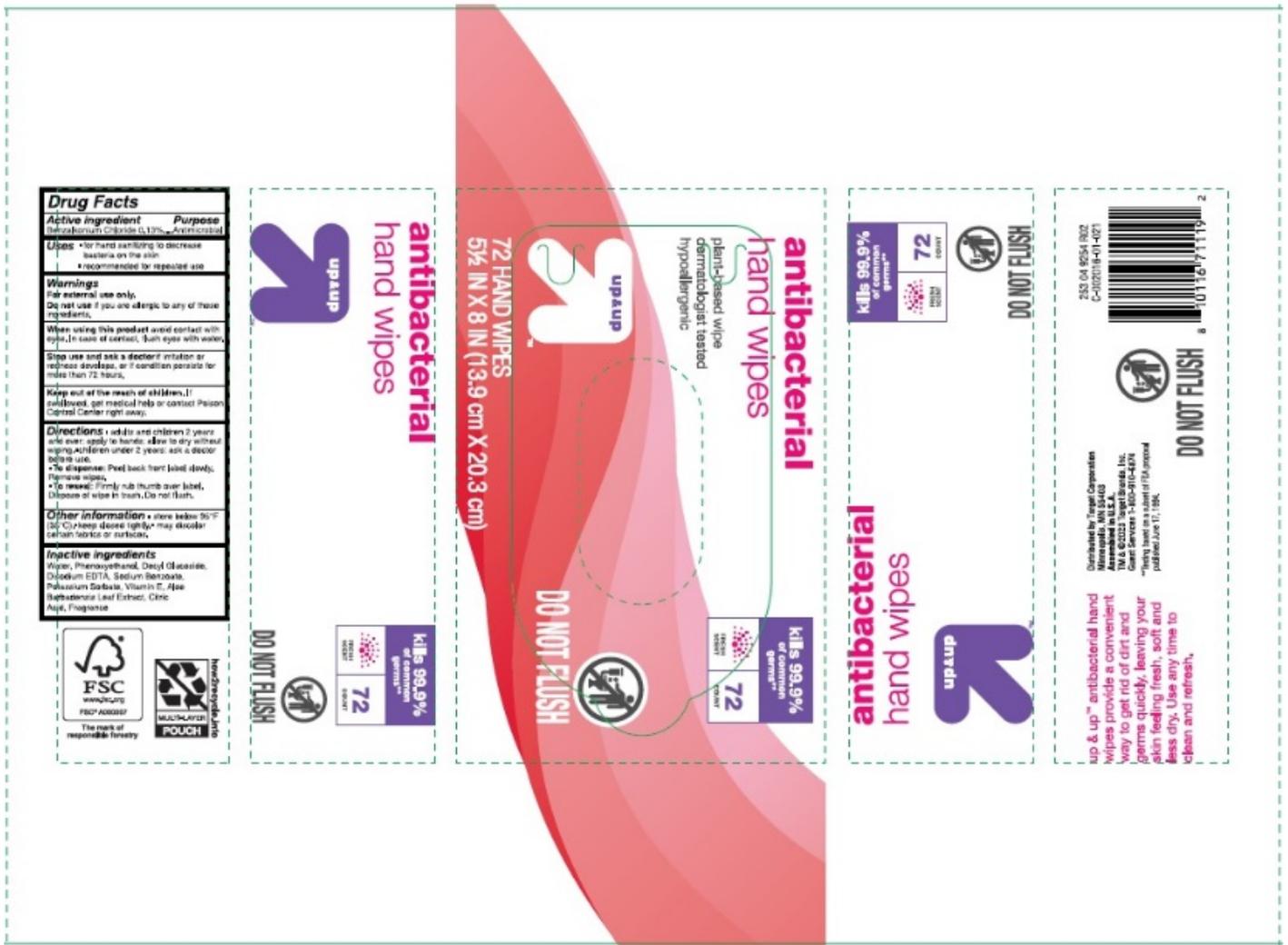
DO NOT FLUSH

Distributed by Target Corporation
Minneapolis, MN 55403
Assembled in U.S.A.
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Guest Services 1-800-910-8874
**Testing based on a subset of FDA protocol published June 17, 1991.

up & up™ antibacterial hand wipes provide a convenient way to get rid of dirt and germs quickly, leaving your skin feeling fresh, soft and less dry. Use any time to clean and refresh.

8 10116 11118 5

72 HAND WIPES
5½ IN X 8 IN



BENZALKONIUM CHLORIDE

antibacterial hand wipes cloth

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82442-250
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

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
POTASSIUM SORBATE (UNII: 1VPU26JZ24)	

PHENOXYETHANOL (UNII: HIE492ZZ3T)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82442-250-40	40 in 1 CONTAINER; Type 0: Not a Combination Product	03/03/2023	
2	NDC:82442-250-72	72 in 1 CONTAINER; Type 0: Not a Combination Product	03/03/2023	
3	NDC:82442-250-25	25 in 1 CONTAINER; Type 0: Not a Combination Product	03/03/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	03/03/2023	

Labeler - Target Corporation (006961700)

Registrant - Target Corporation (006961700)

Revised: 12/2025

Target Corporation