DODYS WIPES- antiseptic hand sanitizer wipes cloth
Betone SA de CV

OTC - ACTIVE INGREDIENT SECTION

Benzalkonium Chloride 0.13% (w/w)

OTC - PURPOSE SECTION

Antiseptic

INDICATIONS & USAGE SECTION

- For hand washing to decrease bacteria on the skin.

WARNINGS SECTION

For external use only.

Flammable, keep away from fire or flame.

OTC - STOP USE SECTION

Stop use and ask a doctor if irritation and redness develop; condition persists for more than 72 hours.

OTC - KEEP OUT OF REACH OF CHILDREN SECTION

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

DOSAGE & ADMINISTRATION SECTION

Wet hands thoroughly with product and allow to dry without wiping.

INACTIVE INGREDIENT SECTION

Water, glycerin, polysorbate 20, tocopheryl acetate, PEG-12 dimethicone, methylisothiazolinone, iodopropynyl butylcarbamate, fragrance, citric acid.

STORAGE AND HANDLING SECTION

Store in a cool place.

OTC - DO NOT USE SECTION

Do not use in the eyes

Dodys Box with 24 Packets, each Packet with 48 Wipes NDC 77824-001-02

Dodys Main Packet, 48 Wipe packet NDC 77824-001-01

Dodys Big Size Product NDC 77824-001

DODYS WIPES

antiseptic hand sanitizer wipes cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:77824-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM -	BENZALKONIUM	0.13 mg

UNII:7N6JUD5X6Y)	CHLORIDE	0.13 mg

Inactive Ingredients

Ingredient Name	Strength
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	
PEG-12 DIMETHICONE (UNII: ZEL54N6W95)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77824-001-01	48 in 1 PACKET; Type 0: Not a Combination Product	06/26/2023	
2	NDC:77824-001-02	24 in 1 BOX; Type 0: Not a Combination Product	06/26/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	06/26/2023	

Labeler - Betone SA de CV (813262755)

Establishment

Name	Address	ID/FEI	Business Operations
Betone SA de CV		813262755	manufacture(77824-001) , pack(77824-001) , label(77824-001)

Revised: 12/2023 Betone SA de CV