# BABY DOVE- 8h moisturization hand sanitizer fragrance free gel Conopco Inc. d/b/a/ Unilever

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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# **Baby Dove 8h Moisturization Hand Sanitizer Fragrance Free**

# BABY DOVE 8H MOISTURIZATION HAND SANITIZER FRAGRANCE FREE - Ethyl Alcohol gel

Baby Dove 8h Moisturization Hand Sanitizer

# **Drug Facts**

# Active ingredient

Alcohol (61%)

# **Purpose**

**Antiseptic** 

#### Uses

Hand sanitizer to help reduce bacteria on the skin

## **Warnings**

For external use only

Flammable. Keep away from fire or flame, Do not store in car.

**Avoid contact with eyes,** in case of contact, rinse eyes thoroughly with water immediately.

**If irritation develps,** discontinue use and consult a doctor.

# Keep out of reach of children except under adult supervision.

If swallowed, get medical help or contact a poison control center right away.

#### **Directions**

- Not recommended for use on infants (younger than 12 months).
- Children under 6 years of age should be supervised when using this product.
- Place enough product in your palm to thoroughly cover your hands and rub hands togeter briskly until dry.
- for babies 1 year\*, rub their hands in yours and do not let them touch their eyes right after rubbing

Allow to dry without wiping or rinsing

## Other information

Store below 105°F (40°C). May discolor fabrics or surfaces.

# Inactive ingredients

Water (Aqua), Glycerin, Butylene Glycol, Dimethicone, Glycine Soja (Soybean) Oil, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Echium Plantagineum Seed Oil, Aminomethyl Propanol, Hydrogenated Lecithin, Tocopherol, Xanthan Gum, Stearic Acid, Citric Acid, Benzyl Alcohol

#### Questions or Comments?

Call 1-800-761-DOVE(3683) or visit us at www.babydove.com

# **Packaging**





8h moisturization hand sanitizer fragrance free gel

# Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:64942-2036 Route of Administration TOPICAL

ctive Ingredient/Active Moiety			
Ingredient Name	<b>Basis of Strength</b>	Strength	
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	610 mg in 1 mL	

Inactive Ingredients		
Ingredient Name	Strength	
BENZYL ALCOHOL (UNII: LKG8494WBH)		
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)		
WATER (UNII: 059QF0KO0R)		
GLYCERIN (UNII: PDC6A3C0OX)		
DIMETHICONE (UNII: 92RU3N3Y10)		
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)		
CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)		
TOCOPHEROL (UNII: ROZB2556P8)		
SOYBEAN OIL (UNII: 241ATL177A)		
HYDROGENATED SOYBEAN LECITHIN (UNII: H1109Z9J4N)		
XANTHAN GUM (UNII: TTV12P4NEE)		
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)		
ECHIUM PLANTAGINEUM SEED OIL (UNII: PIB7XBU8XW)		
STEARIC ACID (UNII: 4ELV7Z65AP)		

P	ackaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
	NDC:64942- 2036-1	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/15/2022		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	07/15/2022		

# Labeler - Conopco Inc. d/b/a/ Unilever (001375088)

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