BABYLISS PRO PROTECT HAND SANITIZER- alcohol gel
Conair Corporation

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.

BABYLISS PRO PROTECT HAND SANITIZER

Drug Facts

Active Ingredient

Ethyl Alcohol 70% V/V

Purpose

Antiseptic

Use

Antibacterial hand sanitizing gel to help kill bacteria and germs as antiseptic on the skin. For use when soap and water are not available.

Warnings

Flammable. Keep away from fire or flame.

For external use only

When using this product

keep out of eyes and mouth. In case of contact with eyes, immediately rinse eyes thoroughly with water.

Stop use and ask doctor if

irritation, rash or allergy occurs. These may be signs of a serious condition.

Keep out of reach of children.

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

Directions

Apply. Put enough product on hands to cover all surfaces. Rub hands together, until hands feel dry. This should take around 20 seconds. Recommended for repeated use. Supervise children under 6 years of age when using this product to avoid swallowing.

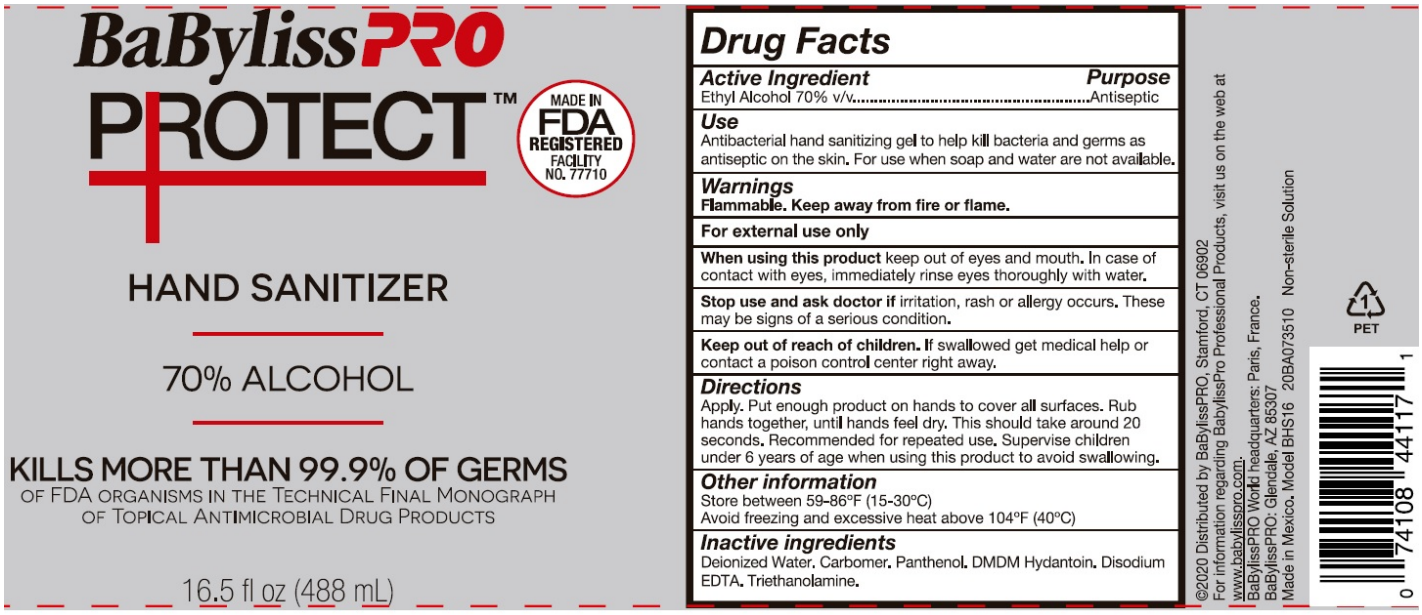
Other information

Store between 59-86°F (15-30°C)
Avoid freezing and excessive heat above 104°F (40°C)

Inactive ingredients

Deionized Water. Carbomer. Panthenol. DMDM Hydantoin. Disodium EDTA. Triethanolamine.

Package Labeling



BABYLISS PRO PROTECT HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:12829-0002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.7 mL in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
PANTHENOL (UNII: WV9CM0O67Z)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
TROLAMINE (UNII: 9O3K93S3TK)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:12829-0002-0	12 in 1 BOX	09/25/2020	
1		488 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	09/25/2020	

Labeler - Conair Corporation (001661222)

Revised: 10/2020 Conair Corporation