DR PURE ANTISEPTIC LIQUID HAND BERRIES MIX SCENTED- benzalkonium chloride liquid
Industrias Bernal Canton S.A. de C.V.

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.

Dr Pure Antiseptic Liquid Hand Soap, Berries Mix Scented

Drug Facts

Active ingredient

Benzalkonium chloride 0.1%

Purpose

Antiseptic

Use

For hand washing to decrease bacteria on the skin

Warnings

For external use only.

Do not use

in the eyes.

Stop use and ask a doctor if

- irritation and redness develop
- condition persists for more than 72 hours

Keep out of reach of children.

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

Directions

- Wet hands and forearms.
- Apply palmful to hands and forearms.
- Scrub thoroughly.
- Rinse and repeat

Inactive ingredients

Water, Sodium Lauryl Sulfate, Sodium Chloride, Cocamidopropyl Betaine, DMDM Hydantoin, Glycerol, Aloe Barbadensis Leaf Juice Extract, Fragrance, Color Red No. 40 (Cl 16035), Red No. 3 (CI 45430) and Yellow No. 5 (Cl 19140).

Package Labeling:

The unique formula of Dr. Pure True Sanitizer liquid hand soap combines antiseptic protection with moisturizers and leaves your hands feeling soft and protected. / La fórmula única del jabón líquido para manos Dr. Pure True Sanitizer combina su protección antiséptica con humectantes y deja las manos suaves y protegidas.

Drug Facts / Información Sobre El Fármaco

Active ingredient / Ingrediente activo ***Purpose / Propósito***
Benzalkonium chloride 0.1% / Cloruro de benzalconio al 0.1%.....Antiseptic / Antiséptico

Use / Empleo
For hand washing to decrease bacteria on the skin / Para lavarse las manos con el fin de disminuir las bacterias en la piel.

Warnings / Advertencias
For external use only. / Sólo para uso externo.

Do not use in the eyes. / **No lo aplique** en los ojos.

Stop use and ask a doctor if / Deje de usarlo y consulte a un médico si:
- irritation and redness develop / se presenta irritación y enrojecimiento.
- condition persists for more than 72 hours / Si la condición persiste por más de 72 horas consulte a su médico.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away. **Manténganse fuera del alcance de los niños.** En caso de ingestión, busque ayuda médica o pónganse en contacto con el Centro de Control de Envenenamientos inmediatamente.

Directions
- Wet hands and forearms. / Humedezca las manos y los antebrazos.
- Apply palmful to hands and forearms. / Aplique un poco en las manos y los antebrazos.
- Scrub thoroughly. / Frote bien.
- Rinse and repeat. / Enjuague y repita.

Inactive ingredients / Ingredientes inactivos
Water, Sodium Lauryl Sulfate, Sodium Chloride, Cocamidopropyl Betaine, DMDM Hydantoin, Glycerol, Aloe Barbadensis Leaf Juice Extract, Fragrance, Color Red No. 40 (CI 16035), Red No. 3 (CI 45430) and Yellow No. 5 (CI 19140). / Agua, Lauril Sulfato de Sodio, Cloruro de Sodio, Cocamidopropil Betaina, DMDM Hidantoína, Glicerol, Extracto de Zumo de Hojas de Aloe Barbadensis, Fragancia, Color Rojo No. 40 (CI 16035), Rojo No. 3 (CI 45430) y Amarillo No. 5 (CI 19140).

Made in Mexico / Hecho en México

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Dr. Pure true sanitizer®

ANTISEPTIC LIQUID HAND SOAP / JABÓN LÍQUIDO ANTISÉPTICO PARA MANOS
Benzalkonium Cloride 0.1% / Cloruro de Benzalconio al 0.1%

BERRIES MIX SCENTED
CON AROMA A FRUTOS ROJOS

16.9 fl oz (1.06 pt) 500 mL

DR PURE ANTISEPTIC LIQUID HAND BERRIES MIX SCENTED

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:78239-005
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM -	BENZALKONIUM	1 mg

UNII:7N6JUD5X6Y)	CHLORIDE	in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
FD&C RED NO. 3 (UNII: PN2ZH5LOQY)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78239-005-16	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/09/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	01/09/2021	

Labeler - Industrias Bernal Canton S.A. de C.V. (812827340)

Revised: 6/2021 Industrias Bernal Canton S.A. de C.V.