

SIGNATRY ANTIBACTERIAL PLUM FOAM HANDWASH- benzalkonium chloride liquid
GOJO Industries, Inc.

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.

Signatry Antibacterial Plum Foam Handwash

Active ingredient

Benzalkonium Chloride 0.5%

Purpose

Antimicrobial

Uses

- Handwash to help decrease bacteria on the skin
- Recommended for repeated use

Warnings

For external use only

When using this product do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash appears and lasts

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

Directions

- Wet hands
- Apply a small amount of product and work into a lather
- Rinse well and dry hands completely

Inactive Ingredients

Water (Aqua), Propanediol, Glycerin, Cocamidopropyl Betaine, PEG-80 Sorbitan Laurate, Citric Acid, Ethylhexylglycerin, Lauramine Oxide, Polyquaternium-10, Trisodium Ethylenediamine Disuccinate, Fragrance (Parfum), Phenoxyethanol



**ANTIBACTERIAL PLUM
FOAM HANDWASH**

**JABÓN ESPUMOSO PARA
MANOS ANTIBACTERIAL
DE CIRUELAS**



S8712

Made in U.S.A. for: Hecho en los
E.E.U.U. por: Signatry, Inc. Akron, OH
44309 1-800-321-9647
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reserved. Todos los derechos
reservados.

8712-640-SGY-F

700 mL (23.6 US/ÉU FL OZ)

Drug Facts

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Ethylenediamine Disuccinate,
Fragrance (Parfum), Phenoxyethanol

Datos Farmacológicos

Ingrediente activo Propósito
Cloruro de benzalconio 0.5%.....Antimicrobiano

Usos • Lavado de manos empleado
para disminuir la cantidad de
bacterias en la piel
• Recomendado para uso reiterado ▶

Datos Farmacológicos (continuado)

Advertencias

Sólo para uso externo

Al utilizar este producto, evitar el
contacto con los ojos o con la zona
alrededor de los ojos. En caso de contacto,
enjuagar completamente los ojos con agua.

**Dejar de usar el producto y consultar a
un médico** si aparece y persiste una
irritación o erupción cutánea

**Mantener fuera del alcance de los
niños.** En caso de ingestión, de inmediato
acudir a un médico o ponerse en contacto
con un centro para el control de tóxicos.

Modo de uso • Mojarse las manos
• Aplicar una pequeña cantidad del
producto y frotar las manos hasta producir
una espuma abundante • Enjuagar bien
• Secarse las manos completamente

Ingredientes inactivos:
Agua, Propanediol, Glicerina, Cocamidopropil
betaina, Laurato de PEG-80 sorbitan, Acido
citrnico, Ethylhexil Glicerina, Oxido de lauramina,
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disuccinato, Fragancia, Fenoxietanol



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Poliquaternio-10, Trisódico etilendiamina
disuccinato, Fragancia, Fenoxietanol

SIGNATRY ANTIBACTERIAL PLUM FOAM HANDWASH

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:21749-545
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Benzalkonium Chloride (UNII: F5UM2KM3W7) (BENZALKONIUM -	Benzalkonium	0.5 mg

UNII:7N6JUD5X6Y)	Chloride	in 100 mL
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Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	
PROPANEDIOL (UNII: 5965N8W85T)	
Glycerin (UNII: PDC6A3C0OX)	
Cocamidopropyl Betaine (UNII: 5OCF3O11KX)	
PEG-80 Sorbitan Laurate (UNII: 239B50Y732)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
Ethylhexylglycerin (UNII: 147D247K3P)	
Lauramine Oxide (UNII: 4F6FC4MI8W)	
POLYQUATERNIUM-10 (10000 MPAS AT 2%) (UNII: PI1STR9QYH)	
TRISODIUM ETHYLENEDIAMINE DISUCCINATE (UNII: YA22H34H9Q)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21749-545-08	236 mL in 1 PACKAGE; Type 0: Not a Combination Product	06/14/2017	
2	NDC:21749-545-97	700 mL in 1 PACKAGE; Type 0: Not a Combination Product	06/14/2017	
3	NDC:21749-545-89	1200 mL in 1 PACKAGE; Type 0: Not a Combination Product	06/14/2017	
4	NDC:21749-545-90	1250 mL in 1 PACKAGE; Type 0: Not a Combination Product	06/14/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	06/14/2017	

Labeler - GOJO Industries, Inc. (004162038)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		036424534	manufacture(21749-545)