

**HUSTLE BUTTER ANTIBACTERIAL FOAMING- benzalkonium chloride soap
H.B.D. DIST. INC.**

HUSTLE BUTTER ANTIBACTERIAL FOAMING SOAP

Drug Facts

Active ingredient

Benzalkonium Chloride 0.10%
w/v

Purpose

Antibacterial

Uses

- for washing to decrease bacteria on the skin • recommended for repeated use

Warnings

For external use only

Do not use • in the eyes • if you are allergic to any ingredients

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 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 the skin • apply palmful to hands • scrub thoroughly on the skin • rinse and repeat

Other information

avoid exposure to direct sunlight and excessive heat.

Inactive ingredients

Aloe Barbadensis Leaf Juice, Coceth-7, PPG-1-PEG-9 Lauryl Glycol Ether, PEG-40

Hydrogenated Castor Oil, Soytrimonium Chloride, Fragrance (Parfum), Propylene Glycol, Glycerin, Water (Aqua), Chamomilla Recutita (Matricaria) Flower Extract, Camellia Sinensis Leaf Extract, Arnica Montana Flower Extract, Hamamelis Virginiana (Witch Hazel) Leaf Extract, Symphytum Officinale Leaf Extract, Rosmarinus Officinalis (Rosemary) Oil, Methylchloroisoethiazolinone, Methylisothiazolinone, Benzyl Benzoate, Citronellol, Coumarin, Geraniol, Hexyl Cinnamal, Hydroxycitronellal

Questions or comments?

+1-212-505-3222

You may also report serious side effects to this phone number.

Mon - Fri 9AM-5PM ET

PROFESSIONAL TATTOO CARE

Distributed by H.B.D. Dist. Inc.
132 Crosby Street, New York, NY
10012

Contact Us: info@hustlebutter.com

[www. hustlebutter.com](http://www.hustlebutter.com)

Certified Vegan.

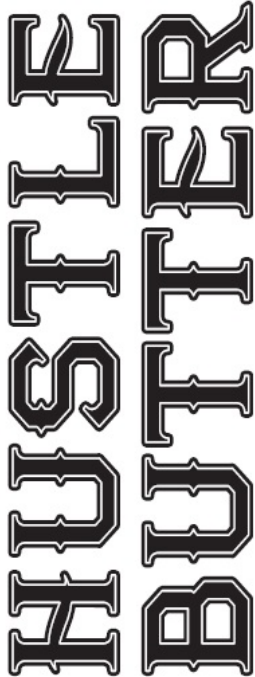
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Packaging



PROFESSIONAL TATTOO CARE



ANTIBACTERIAL FOAMING SOAP

7 FL OZ / 207 mL

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0 51497 09649 6

HUSTLE BUTTER ANTIBACTERIAL FOAMING

benzalkonium chloride soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72014-200
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.1 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

COCETH-7 (UNII: 58Y261JLH5)
PPG-1-PEG-9 LAURYL GLYCOL ETHER (UNII: 5R8J43K25L)
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)
TRIMETHYLAMINE HYDROCHLORIDE (UNII: UW38SRM77K)
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)
GLYCERIN (UNII: PDC6A3C0OX)
WATER (UNII: 059QF0KO0R)
CHAMOMILE (UNII: FGL3685T2X)
GREEN TEA LEAF (UNII: W2ZU1RY8B0)
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)
HAMAMELIS VIRGINIANA LEAF (UNII: T07U1161SV)
COMFREY LEAF (UNII: DG4F8T839X)
ROSEMARY OIL (UNII: 8LGU7VM393)
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)
BENZYL BENZOATE (UNII: N863NB338G)
.BETA.-CITRONELLOL, (R)- (UNII: P01OUT964K)
COUMARIN (UNII: A4VZ22K1WT)
GERANIOL (UNII: L837108USY)
.ALPHA.-HEXYLCINNAMALDEHYDE (UNII: 7X6O37OK2I)
HYDROXYCITRONELLAL (UNII: 8SQ0VA4YUR)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72014-200-07	207 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/30/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	05/30/2024	

Labeler - H.B.D. DIST. INC. (031127044)

Revised: 5/2024

H.B.D. DIST. INC.