

HAN I FOAM- alcohol liquid
Momar Incorporated

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.

Han I Foam 6007 Drug Facts and Label

Drug Facts Box OTC-Active Ingredient Section

Ethyl Alcohol 62%

Drug Facts Box OTC-Purpose Section

Antiseptic

Drug Facts Box OTC-Indications & Usage Section

for hand-washing to decrease bacteria on the skin, only when water is not available

Drug Facts Box OTC-Warnings Section

FLAMMABLE, keep away from fire and flames

For external use only

Drug Facts Box OTC-When Using Section

do not get into eyes

if contact occurs, rinse eyes thoroughly with water

Drug Facts Box OTC-Stop Use Section

irritation and redness develop

Drug Facts Box OTC-Keep Out of Reach of Children Section

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

Drug Facts Box OTC-Dosage & Administration Section

press pump twice to deliver two squirts (about a quarter size) of foaming product onto the palm of your hand

rub hands together and allow to dry without wiping

Drug Facts Box OTC-Inactive Ingredient Section

water, PEG-10 acrylate, perfluorohexylethyl acrylate copolymer, propylene glycol

Han I Foam 6007



- Contains Skin Conditioners
- Kills 99.99% of common germs that cause illness in as little as 15 seconds

Net Contents: 50 ml (1.7 oz.)

Sold by: **MOMAR**® INCORPORATED
 1830 Ellsworth Industrial Drive, N.W., Atlanta, Georgia 30318
 A Cleaning Force Throughout the World Since 1947
 Providing Products of Momar Research in the U.S.A., Europe,
 Africa, and Australia. 800-556-3967 • www.momar.com

6007Z1Q610.021420 Batch No: XXXX

Drug Facts

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Inactive Ingredients

Water; PEG-10 Acrylate, Perfluorohexylethyl Acrylate Copolymer; Perfluorohexylethyl Alcohol; Propylene Glycol



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Han I Foam 6007

HAN I FOAM

alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63533-007
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.62 mL in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PEG-10 ACRYLATE/PERFLUOROHXYLETHYL ACRYLATE COPOLYMER (UNII: D76Z87928N)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63533-007-17	532 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/10/2020	
2	NDC:63533-007-18	50 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/10/2020	
3	NDC:63533-007-14	3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/10/2020	
4	NDC:63533-007-10	1000 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	08/10/2020	
5	NDC:63533-007-11	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/10/2020	
6	NDC:63533-007-12	1000 mL in 1 BAG; Type 0: Not a Combination Product	08/10/2020	
7	NDC:63533-007-08	1 in 1 BOX	08/10/2020	
7		1000 mL in 1 BAG; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC mono graph not final	part333E	08/10/2020	

Labeler - Momar Incorporated (003266616)

Registrant - ABC Compounding Co., Inc. (003284353)

Establishment

Name	Address	ID/FEI	Business Operations
ABC Compounding Co., Inc.		003284353	manufacture(63533-007)

Revised: 8/2020

Momar Incorporated