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 6585 Drug Facts and Label

Drug Facts Box OTC-Active Ingredient Section

Ethyl Alcohol 62% v/v

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, DEA-C8-18 perfluoroalkylethyl phosphate, propylene glycol, fragrance

Han-I-Foam 6585 50 ml

6585Z1Q610.jpg Han-I-Foam 50ml



 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

Drug Facts

Active Ingredient

Purpose Ethyl Alcohol 62% v/v.....Antiseptic

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

Warnings

Flammable, keep away from fire and flames For external use only

When using this product

- do not get into eves
- if contact occurs, rinse eyes thoroughly with water

Stop use and ask a doctor if

■ irritation and redness develop

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

Directions ■ 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

Inactive Ingredients water, DEA-C8-18 perfluoroalkylethyl phosphate, propylene glycol, fragrance

6585Z1Q610.1211

HAN-I-FOAM

alcohol liquid

Product	Information

Product Type	HUMAN OTC DRUG LABEL	Item Code (Source)	NDC:63533-855

Route of Administration TOPICAL **DEA Schedule**

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength		
ALCOHOL (ALCOHOL)	ALCOHOL	0.62 mL in 1 mL		

Inactive Ingredients

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Ingredient Name	Strength	
WATER		
PROPYLENE GLYCOL		
DIETHANOLAMINE BIS(C8-C18 PERFLUOROALKYLETHYL)PHOSPHATE		

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63533-855-06	1 in 1 BOX		
1		800 mL in 1 BAG		

NDC:63533-855-17	532 mL in 1 BOTTLE, PLASTIC
NDC:63533-855-24	118 mL in 1 BOTTLE, PLASTIC
NDC:63533-855-01	1200 mL in 1 CARTRIDGE
NDC:63533-855-03	350 mL in 1 CARTRIDGE
NDC:63533-855-05	540 mL in 1 BOTTLE, PLASTIC
NDC:63533-855-07	700 mL in 1 BAG
NDC:63533-855-09	2000 mL in 1 CARTRIDGE
NDC:63533-855-10	1000 mL in 1 CARTRIDGE
NDC:63533-855-11	1000 mL in 1 BOTTLE, PLASTIC
NDC:63533-855-12	1000 mL in 1 BAG
NDC:63533-855-13	800 mL in 1 BAG
NDC:63533-855-14	3785 mL in 1 BOTTLE, PLASTIC
NDC:63533-855-15	946 mL in 1 BOTTLE, PLASTIC
NDC:63533-855-28	149 mL in 1 BOTTLE, PLASTIC
NDC:63533-855-27	800 mL in 1 CARTRIDGE
NDC:63533-855-55	208200 mL in 1 DRUM
NDC:63533-855-08	1 in 1 BOX
	1000 mL in 1 BAG
NDC:63533-855-16	236 mL in 1 BOTTLE, PLASTIC
NDC:63533-855-18	50 mL in 1 BOTTLE, PLASTIC
	NDC:63533-855-24 NDC:63533-855-01 NDC:63533-855-05 NDC:63533-855-07 NDC:63533-855-07 NDC:63533-855-10 NDC:63533-855-11 NDC:63533-855-12 NDC:63533-855-13 NDC:63533-855-14 NDC:63533-855-15 NDC:63533-855-15 NDC:63533-855-28 NDC:63533-855-27 NDC:63533-855-55 NDC:63533-855-08

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	09/01/2009	

Labeler - Momar Incorporated (003266616)

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

Establishment			
Name	Address	ID/FEI	Business Operations
ABC Compounding Co., Inc.		003284353	manufacture(63533-855)

Revised: 11/2012 Momar Incorporated