MG DERMASAN- n/a liquid
Morgan Gallacher Inc.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

MG DermaSan

Active Ingredient

Benzalkonium Chloride.....0.1 %

Uses

- For hand sanitizing to decrease bacteria on the skin
- Recommended for repeated use

Purpose

Antimicrobial

Warnings

- For external use only.
- When using this product avoid contact with eyes. In case of eye contact, flush eyes with water. Do not ingest.
- Stop use and ask a doctor if irritation or redness develops and conditions persist.

Keep out of reach of children

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

Directions

- Pump a small amount of foam into palm of hand.
- Rub thoroughly over all surfaces of both hands.
- Rub hands together briskly until dry.

Inert Ingredients

- water
- cetrimonium chloride
- laurtrimonium chloride
- dihydroxyethyl cocamine oxide
- glycereth-17 cocoate

- citric acid

MG DermaSan

MG DERMASAN

n/a liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50241-259
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 g

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
LAURTRIMONIUM CHLORIDE (UNII: A81MSI0FIC)	
DIHYDROXYETHYL COCAMINE OXIDE (UNII: 8AR51R3BL5)	
GLYCERETH-17 COCOATE (UNII: 3057VPT0KC)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:50241-259-02	1 in 1 BOX	10/01/2018	
1	NDC:50241-259-01	3756 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		10/01/2018	

Labeler - Morgan Gallacher Inc. (028311595)

Registrant - Morgan Gallacher Inc. (028311595)

Establishment

Name	Address	ID/FEI	Business Operations
Morgan Gallacher Inc.		028311595	manufacture(50241-259) , api manufacture(50241-259) , pack(50241-259)

Revised: 12/2023 Morgan Gallacher Inc.