

**SHIELD DR.- hand sanitizer liquid spray with peppermint and lemongrass, 70% alcohol liquid
MEDIPHARCO PHARMACEUTICAL JOINT STOCK COMPANY**

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.

Ethyl Alcohol 70%

Purified water, Isopropyl myristate, Fragrance, Yello (C.I. 47005), Blue (C.I. 42090)

If swallowed, get medical attention or contact poison control center immediately

For external use only.

Flammable, keep away from fire or source of ignition.

Hand sanitizer to help reduce bacteria on the skin it could cause disease

Wet hand thoroughly with spray and rub hand together briskly until dry. Children under the age of 6 should be supervised while using this product. Not recommended for infants.

Antiseptic

Store at room temperature 66o F to 77oF (20oC to 25oC)

59ml NDC: 79633-302-59

DRUG FACTS
Ethyl Alcohol 70%.....Antiseptic
Uses: Hand sanitizer to help reduce bacteria on the skin that could cause disease.
Warnings: For External Use Only Flammable, keep away from fire or source of ignition. When using the product, keep out of eyes. In case of eye contact, flush thoroughly with water. Stop use and ask a doctor if irritation and redness develops and persists for more than 72 hours. Keep out of reach of children. If swallowed, get medical attention or contact a poison control center immediately.
Other Information: Store at room temperature 66°F to 77°F (20°C to 25°C) May discolor certain fabrics
Directions: Wet hands thoroughly with spray and rub hands together briskly until dry. Children under the age of 6 should be supervised while using this product. Not recommended for infants.
Inactive Ingredients: Purified water, Isopropyl myristate, Fragrance, Yellow (C.I. 47005), Blue (C.I. 42090)

hand sanitizer liquid spray with peppermint and lemongrass, 70% alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79633-302
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
WATER (UNII: 059QF0K00R)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
JAMMU LEMONGRASS OIL (UNII: K25ZLU1H0O)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
FD&C BLUE NO. 1 (UNII: HBR47K3TBD)	
PEPPERMINT OIL (UNII: AV092KU4JH)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79633-302-59	59 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	08/19/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/19/2020	

Labeler - MEDIPHARCO PHARMACEUTICAL JOINT STOCK COMPANY (555273697)

Establishment

Name	Address	ID/FEI	Business Operations
MEDIPHARCO PHARMACEUTICAL JOINT STOCK COMPANY		555273697	manufacture(79633-302)

Revised: 8/2020

MEDIPHARCO PHARMACEUTICAL JOINT STOCK COMPANY