PREVICARE HYSSOP SANITIZER- alcohol gel
PREVICARE HOLY WATER PLUS HYSSOP SANITIZER- alcohol gel
Previcare Pharmaceutical, Inc.

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.

Previcare Hyssop Sanitizer Gel

Drug Facts

Active Ingredient

Ethyl Alcohol 70% V/V

Purpose

Antiseptic

Uses

Hand sanitizer to help reduce bacteria that potentially cause disease. For use when soap and water are not available.

Warnings

For external use only. Flammable. Keep away from heat or flame. Do not ingest. Do not use if you are allergic to any ingredient.

Keep out of reach of children. If swallowed, get medical help. Do not use on open wounds. When using ths product avoid contact with eyes. **In case of contact with eyes, rinse thoroughly with water. Stop use and ask a doctor if irritation develops.**

Directions

Apply to one palm, rub hands together until dry. Use as part of your daily cleaning routine. Do not rinse with water. Do not dilute the product. Supervise children when using this product.

Other Information

Store at 50-86°F (15-30°C).

Inactive Ingredients

Aqua (purified water), Hyssop and Rosemary Mint Oil, Glycerin, Aloe Barbadensis (Aloe) Leaf Juice, PEG-75 Lanolin, PEG-7 Glyceryl Cocoate (emollient), Myristyl Alcohol, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Triethanolamine.

Questions?

Call 1 (716) 427-7979

Distributed by: **Previcare Pharmaceutical, Inc.**
50 Fountain Plaza • Suite 1400 • Buffalo • NY • 14202 • **USA**

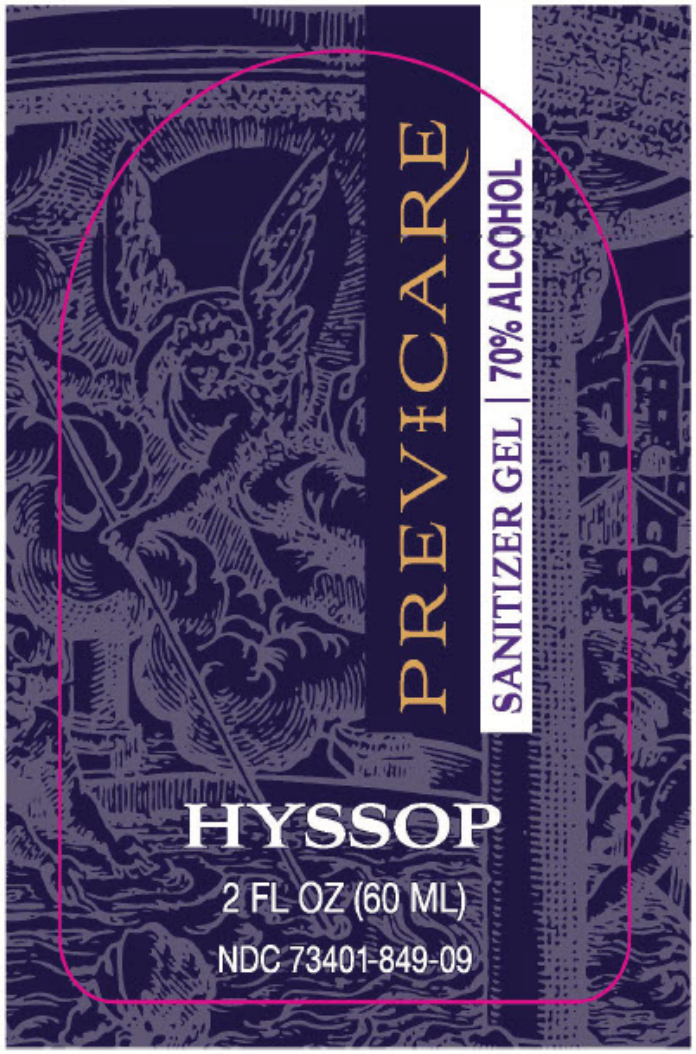
PRINCIPAL DISPLAY PANEL - 60 ML Bottle Label - NDC 73401-849

PREVICARE
SANITIZER GEL | 70% ALCOHOL

HYSSOP

2 FL OZ (60 ML)

NDC 73401-849-09



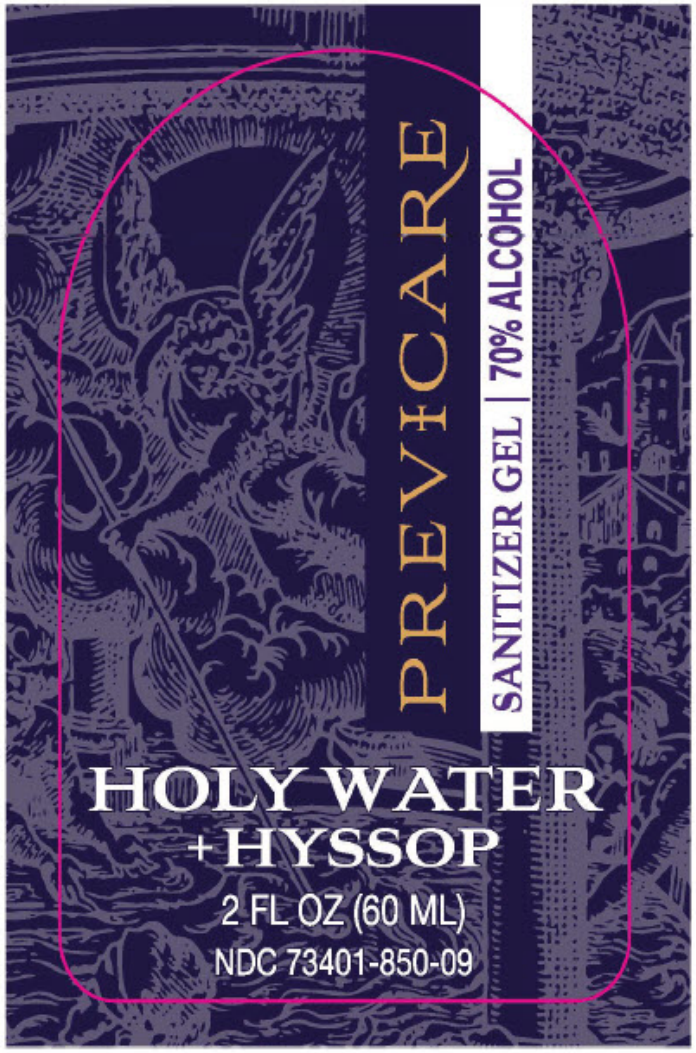
PRINCIPAL DISPLAY PANEL - 60 ML Bottle Label - NDC 73401-850

PREVICARE
SANITIZER GEL | 70% ALCOHOL

HOLY WATER
+ HYSSOP

2 FL OZ (60 ML)

NDC 73401-850-09



PREVICARE HYSSOP SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73401-849
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
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
PEG-75 LANOLIN (UNII: 09179OX7TB)	
MYRISTYL ALCOHOL (UNII: V42034O9PU)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
TROLAMINE (UNII: 9O3K93S3TK)	
PEG-7 GLYCERYL COCOATE (UNII: VNX7251543)	
HYSSOP OIL (UNII: 173D71924B)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73401-849-09	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/15/2021	
2	NDC:73401-849-10	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/17/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part333A	02/15/2021	

PREVICARE HOLY WATER PLUS HYSSOP SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73401-850
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**
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
PEG-75 LANOLIN (UNII: 09179OX7TB)	
MYRISTYL ALCOHOL (UNII: V42034O9PU)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
TROLAMINE (UNII: 9O3K93S3TK)	
PEG-7 GLYCERYL COCOATE (UNII: VNX7251543)	
HYSSOP OIL (UNII: 173D71924B)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73401-850-09	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/15/2021	
2	NDC:73401-850-10	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/17/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part333A	02/15/2021	

Labeler - Previcare Pharmaceutical, Inc. (117180835)

Revised: 2/2021 Previcare Pharmaceutical, Inc.