MORE SAVINGS 50 PERCENT ISOPROPYL RUBBING ALCOHOL WITH WINTERGREEN-isopropyl alcohol liquid
D.J.H. 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.

Active ingredient (BY VOLUME)

ISOPROPYL ALCOHOL (50% CONCENTRATE)

Purpose

TOPICAL ANTIMICROBIAL

Uses

- TO DECREASE GERMS IN MINOR CUTS AND SCRAPES.
- HELPS RELIEVE MUSCULAR ACHES DUE TO EXERTION.

Keep Out of Reach of Children

IN CASE OF AN ACCIDENTAL INGESTION, CONTACT A POISON CONTROL CENTER (1-800-222-1222) IMMEDIATELY.

WARNINGS

For external use only.

- FLAMMABLE, KEEP AWAY FROM FIRE AND FLAME
- IF TAKEN INTERNALLY SERIOUS GASTRIC DISTURBANCES WILL RESULT

ASK A DOCTOR BEFORE USE IF YOU HAVE

- DEEP PUNCTURE WOUNDS SERIOUS BURNS

WHEN USING THIS PRODUCT

- DO NOT GET INTO EYES OR MUCOUS MEMBRANES
- USE ONLY IN A WELL-VENTILLATED AREA.

STOP USE AND ASK A DOCTOR IF

- CONDITION PERSISTS OR GET WORSE.

Directions

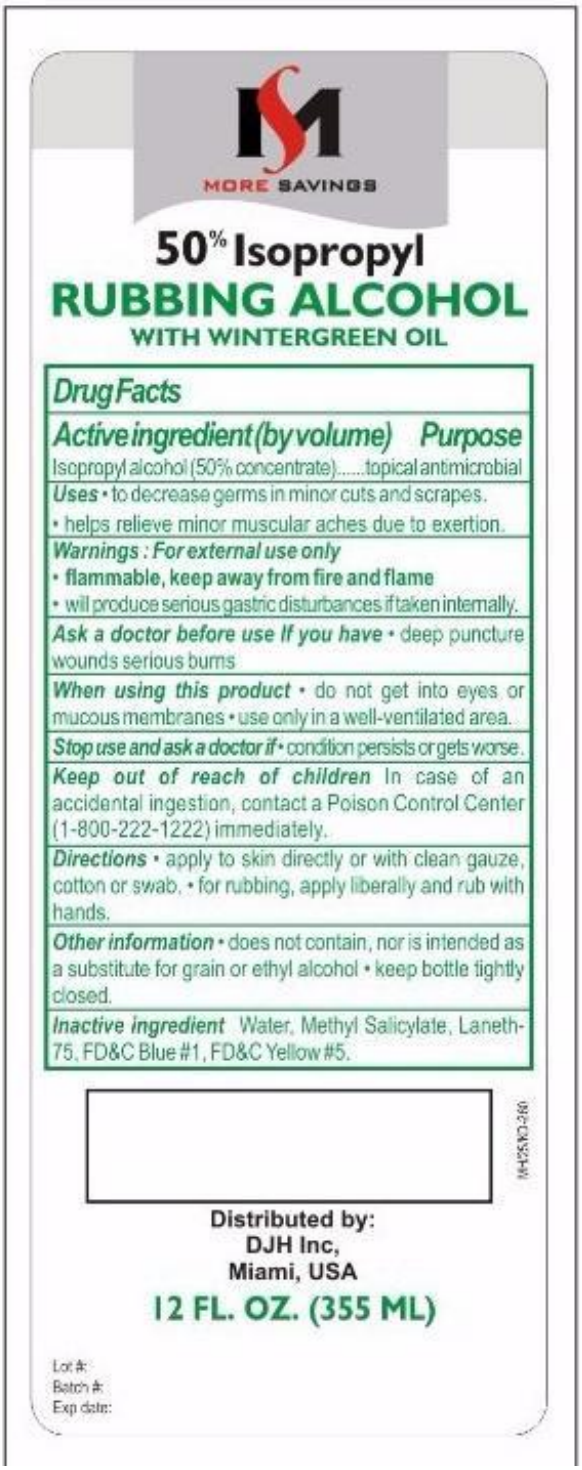
- APPLY TO SKIN DIRECTLY OR WITH CLEAN GAUZE, COTTON OR SWAB.
- FOR RUBBNG, APPLY LIBERALLY AND RUB WITH HANDS.

OTHER INFORMATION

- DOES NOT CONTAIN, NOR IS INTENDED AS A SUBSTITUTE FOR GRAIN OR ETHYL ALCOHOL
- KEEP BOTTLE TIGHTLY CLOSED.

Inactive Ingredient

WATER, METHYL SALICYLATE, LANETH-75, FD&C BLUE #1, FD&C YELLOW #5



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isopropyl alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69811-202
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	50 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
METHYL SALICYLATE (UNII: LAV5U5022Y)	
LANOLIN ALCOHOLS (UNII: 884C3FA9HE)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:69811-202-12	355 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	**Application Number or Monograph Citation**	**Marketing Start Date**	**Marketing End Date**
OTC monograph not final	part333A	05/05/2015	

Labeler - D.J.H. INC (847924271)

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