ISOPROPYL RUBBING ALCOHOL 70 PERCENT WITH WINTERGREEN- isopropyl alcohol solution
JC SALES

JC Sales (as PLD) - NuValu ISOPROPYL RUBBING ALCOHOL, 70% with wintergreen oil (72520-317)

Active ingredient

ISOPROPYL ALCOHOL 50%

Purpose

FIRST AID ANTISEPTIC

Uses

FIRST AID TO HELP THE RISK OF INFECTION IN MINOR CUTS AND SCRAPES.

WARNINGS

For external use only.

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

ASK A DOCTOR BEFORE USE IF YOU HAVE

- DEEP PUNCTURE WOUNDS, ANIMAL BITES OR SERIOUS BURNS

WHEN USING THIS PRODUCT

- DO NOT GET INTO EYES
- DO NOT APPLY OVER LARGE AREAS OF THE BODY.
- DO NOT USE LONGER THAN ONE WEEK UNLESS DIRECTED BY A DOCTOR.

STOP USE AND ASK A DOCTOR IF

- CONDITION PERSISTS OR GET WORSE.

KEEP OUT OF REACH OF CHILDREN. IN CASE OF INGESTION, GET MEDICAL HELP OR CONTRACT A POISON CONTROL CENTER (1-800-222-1222) IMMEDIATELY.

Directions

- CLEAN THE AFFECTED AREA.
- APPLY A SMALL AMOUNT OF THIS PRODUCT ON THE AREA 1 TO 3 TIMES DAILY
- MAY BE COVERED WITH A STERILE BANDAGE
- IF BANDAGED, LET DRY FIRST.

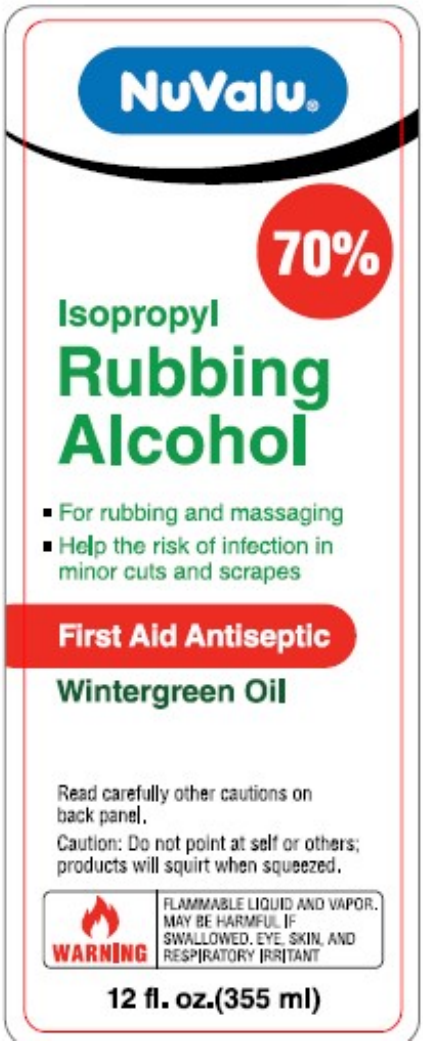
OTHER INFORMATION

- STORE AT ROOM TEMPERATURE
- DOES NOT CONTAIN, NOR IS INTENDED AS A SUBSTITUTE FOR GRAIN OR ETHYL ALCOHOL

Inactive Ingredients

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

Front



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ISOPROPYL RUBBING ALCOHOL 70 PERCENT WITH WINTERGREEN

isopropyl alcohol solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72520-317
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
		Basis of

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
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
LANETH-75 (UNII: QQ4C4Q56ST)	
WATER (UNII: 059QF0KO0R)	
METHYL SALICYLATE (UNII: LAV5U5022Y)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72520-317-12	355 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/08/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	08/08/2024	

Labeler - JC SALES (610969578)

Revised: 8/2024 JC SALES