DE LA CRUZ DECOLORIZED IODINE- ethyl alcohol liquid
DLC Laboratories, Inc.

De La Cruz Decolorized Iodine

Drug Facts

Active Ingredients

Alcohol 45 % denatured with ammonia, Ammonium and Potassium Iodides.

Purpose

Antiseptic

Uses

First aid to help prevent infection in minor cuts, scrapes and burns.

Warnings

For external use only.

Ask a doctor before use if you have

deep or puncture wounds, animal bites, serious burns.

Stop use and ask a doctor if:

The condition persists or gets worse, or if using this product for longer than 1 week.

When using this product:

Do not use in the eyes. If contact occurs, flush with large amounts of water while lifting upper and lower lids. Do not apply over large areas of the body.

Keep out of the reach of children.

In case of acidental ingestion give milk, then give a starch solution made by mixing two tablespoonfools of cornstarch or flour to a pint of water. contact a poison Control Center immediately.

Directions

Clean the affected area. Aply a small amount to the affected area 1 to 3 times daily. May be covered with a sterile bandage. If bandaged, let it dry first.

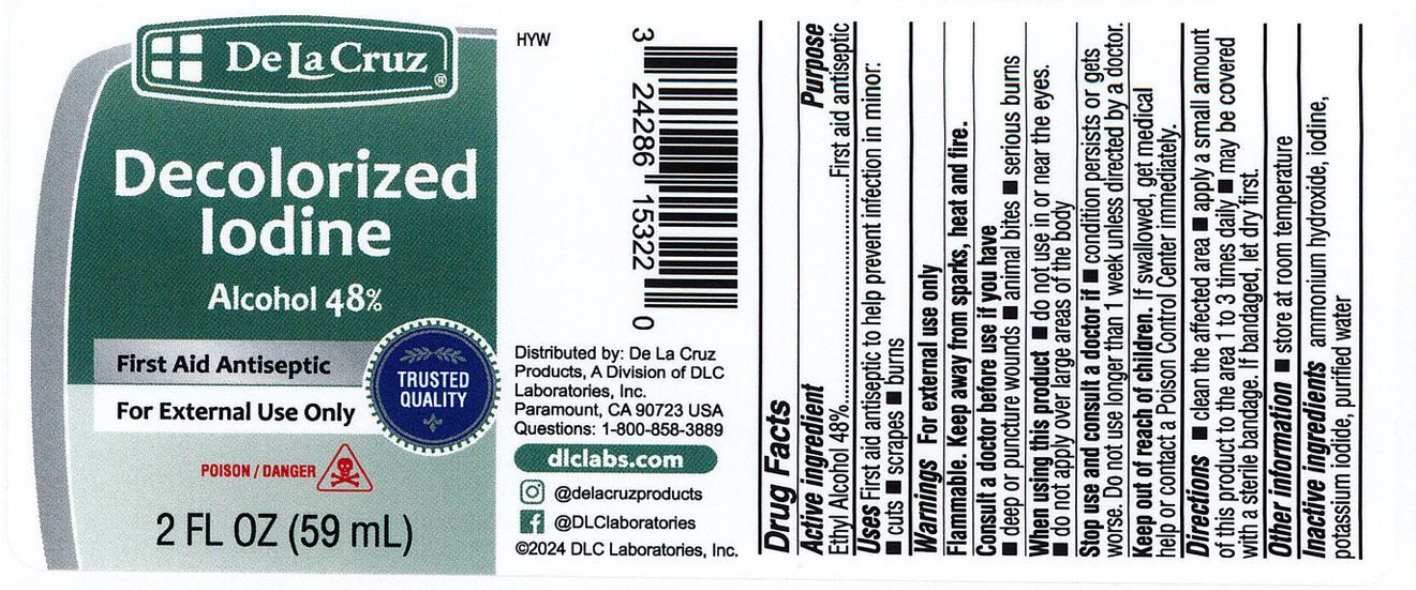
Inactive Ingredient:

Purified Water

Pther Information:

Will stain skin and clothing

Label



DE LA CRUZ DECOLORIZED IODINE

Datos Medicinales

Ingrediente activo **Propósito**
Alcohol 48%....Antiséptico para primeros auxilios

Usos Antiséptico para primeros auxilios; ayuda a prevenir infecciones en:
■ cortaduras menores ■ abrasiones menores ■ quemaduras menores

Advertencias **Para uso externo únicamente**

Inflamable. Manténgase alejado de chispas, calor y fuego.

Consulte con un médico antes de usar si tiene
■ heridas profundas o punzantes
■ mordeduras de animales
■ quemaduras graves

Cuando use este producto
■ no usar en los ojos ni cerca de ellos
■ no aplicar en áreas grandes del cuerpo

Deje de usar y consulte con un médico si
■ la condición persiste o empeora. No usar por más de 1 semana a menos que sea recomendado por un médico.

Manténgase fuera del alcance de los niños.
Si se ingiere, pida ayuda médica o contáctese con un Centro de Control de Envenenamientos inmediatamente.

Instrucciones ■ limpie el área afectada ■ aplique este producto en cantidades pequeñas de 1 a 3 veces al día ■ puede cubrir el área afectada con una venda esterilizada. Antes de vendar, deje que el producto se seque.

Otra información
■ manténgase a temperatura ambiente

Ingredientes inactivos
hidróxido amónico, yodo, yoduro de potasio, agua purificada

Drug Facts

Active ingredient **Purpose**
Alcohol 48%.....First aid antiseptic

Uses First aid antiseptic to help prevent infection in minor: ■ cuts ■ scrapes ■ burns

Warnings **For external use only**

Flammable. Keep away from sparks, heat and fire.

Consult a doctor before use if you have
■ deep or puncture wounds
■ animal bites
■ serious burns

When using this product
■ do not use in or near the eyes
■ do not apply over large areas of the body

Stop use and consult a doctor if
■ the condition persists or gets worse.
Do not use this product longer than 1 week unless directed by a doctor.

Keep out of reach of children.
If swallowed, get medical help or contact a Poison Control Center immediately.

Directions
■ clean the affected area ■ apply a small amount of this product to 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

Inactive ingredients
ammonium hydroxide, iodine, potassium iodide, purified water

Distributed by / Distribuido por:
De La Cruz Products
A Division of DLC Laboratories, Inc.
Paramount, CA 90723 USA
Questions / Preguntas: 1-800-858-3889
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WARNINGS: KEEP OUT OF REACH OF CHILDREN. FLAMMABLE. KEEP AWAY FROM SPARKS, HEAT AND FIRE.

ADVERTENCIAS: MANTÉNGASE FUERA DEL ALCANCE DE LOS NIÑOS. INFLAMABLE. MANTÉNGASE ALEJADO DE CHISPAS, CALOR Y FUEGO.

Drug Facts
Datos Medicinales

Peel Here
Levante Aquí

ethyl alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:24286-1529
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.45 mL in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
AMMONIA (UNII: 5138Q19F1X)	
IODINE (UNII: 9679TC07X4)	
POTASSIUM IODIDE (UNII: 1C4QK22F9J)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:24286-1529-8	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/06/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	03/25/1998	

Labeler - DLC Laboratories, Inc. (093351930)

Registrant - Pharma Nobis, LLC (118564114)

Establishment

Name	Address	ID/FEI	Business Operations
Pharma Nobis, LLC		118564114	manufacture(24286-1529) , analysis(24286-1529) , pack(24286-1529) , label(24286-1529)

Revised: 12/2024 DLC Laboratories, Inc.