**FRESKARO DECOLORIZED IODINE- alcohol liquid**
**Pharma Nobis, LLC**

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**Freskaro Decolorized Iodine**

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Drug Facts

## Active Ingredient

Alcohol 48%

## Purpose

First Aid Antiseptic

## Uses

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

## Warnings

For External Use Only

### When using the product

Do not use in the eyes or apply over large areas of the bosy. In case of deep or puncture wounds, animal bites or serious burns, consult a doctor.

### Stop use and consult a doctor if

the condition persists or gets worse. Do not use this product for 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. Soap will deactivate the effects of this product.

## Directions

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

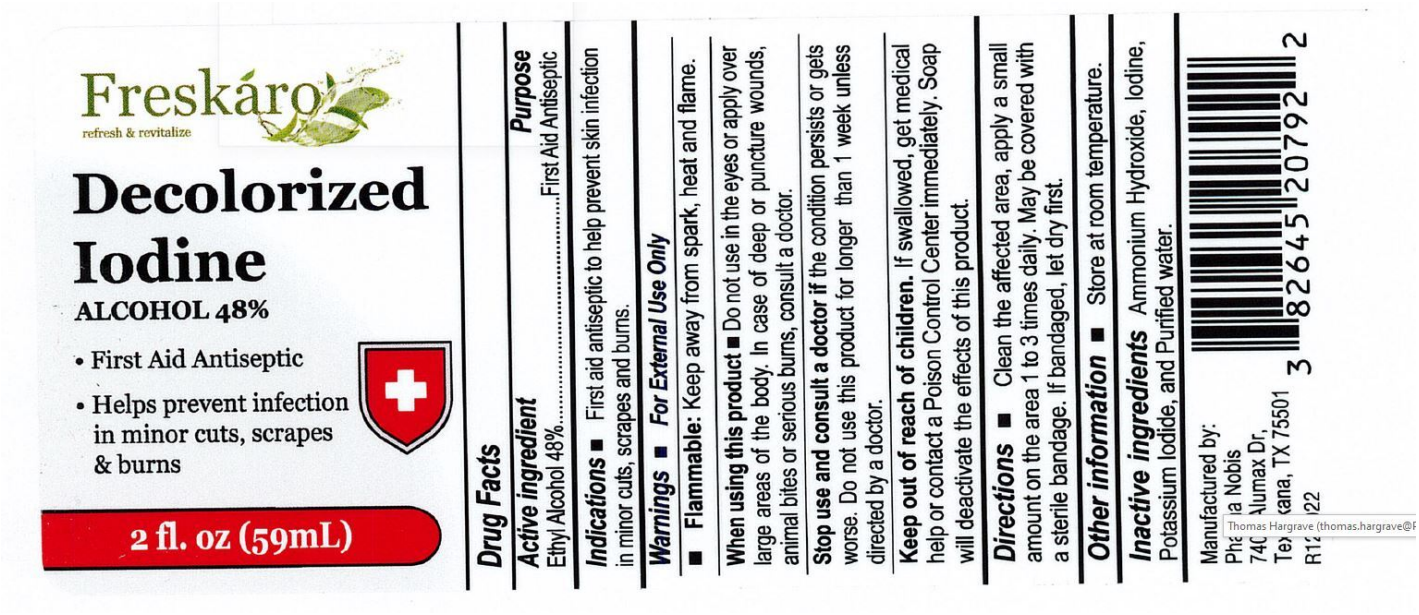
## Inactive Ingredient

Ammonia Hydroxide, Iodine, Potassium Iodide, Purified water.

## Other information

**Flammable:** Keep away from spark, heat and flame

## PRINCIPAL DISPLAY PANEL




# FRESKARO DECOLORIZED IODINE

alcohol liquid

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:82645-207 |
| **Route of Administration** | TOPICAL | | |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **ALCOHOL** (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) | ALCOHOL | 0.48 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:82645-207-92 | 59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | 05/16/2023 | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC Monograph Drug | M003 | 05/16/2023 | |

## Labeler - Pharma Nobis, LLC (118564114)

## Registrant - Pharma Nobis, LLC (118564114)

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| Pharma Nobis, LLC | | 118564114 | analysis(82645-207) , manufacture(82645-207) , pack(82645-207) , label(82645-207) |

Revised: 12/2023 Pharma Nobis, LLC