BETADINE DRY POWDER- povidone-iodine aerosol, spray Atlantis Consumer Healthcare, Inc.

BETADINE® Dry Powder Spray

5%Povidone-iodine

Drug Facts

Active ingredient

Povidone-iodine, 5% (0.5% available iodine).

Purpose

First aidantiseptic

Uses

First aidto help prevent infection in minor

- cuts
- scrapes
- burns

Warnings For external use only

Extremely Flammable Do not use near heat, flame or while smoking. Do notpuncture or incinerate. Contents under pressure. Do not store at temperaturesabove 120°F. Intentional misuse by deliberately concentrating and inhaling the contents can be harmful or fatal

Do not use

- in or near eyes
- over large areas of the body
- if you are allergic to povidone-iodine or any other ingredients in this preparation

When using this product

• do not get into eyes. If contact occurs, rinse eyes thoroughlywith water. If irritation persists, consult a doctor.

Ask a doctor before use if you have

- deep or puncture wounds
- serious burns
- animal bites

Stop use and ask a doctor if

- the condition persists or gets worse
- you need to use this product for more than 1 week

Keep out of reachof children.

If swallowed, get medicalhelp or contact a Poison Control Center right away.

Directions

- Clean the affected area
- SHAKE WELL BEFORE USE
- Hold can approximately 6 inches from wound and apply a thinlayer over affected area 1 to 3 daily
- May be covered with a sterile bandage

Other information

• store at 25°C (77°F); excursions permitted between 15°-30°C(59°-86°F)

Inactive ingredients

Isobutane, isopropyl myristate, pentrane, propane

Dist. by: **AvrioHealth L.P. Stamford, CT 06901-3431** 305653-0A

BetadineDry Powder Spray NDC: 67618-165-01



BETADINE DRY POWDER

povidone-iodine aerosol, spray

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:67618-165

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
POVIDONE-IODINE (UNII: 85H0HZU99M) (IODINE - UNII:9679TC07X4)	IODINE	.05 mg in 1 mL

Inactive Ingredients				
Strength				

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1		56.7 mL in 1 CAN; Type 0: Not a Combination Product	06/15/2020		
Marketing Information					
	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
ОТ	C Monograph Drud	M003	06/15/2020		

Labeler - Atlantis Consumer Healthcare, Inc. (118983925)

Registrant - Atlantis Consumer Healthcare, Inc. (118983925)

Revised: 12/2023 Atlantis Consumer Healthcare, Inc.