ANTISEPTIC- isopropyl alcohol liquid
ReliaMed

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.

Drug Facts

Active Ingredients

Isopropyl Alcohol 71.5%

Purpose

Antiseptic

Uses

For preparation of the skin prior to injection or venipuncture

Warnings

For external use only.

Flammable. Keep away from fire or flame.

Do not use with electrocautery procedures or in or near the eyes. For external use only.

Stop use and ask a doctor if irritation or redness develop and persists for more than 72 hours

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

Directions

- clean the area
- start at the venipuncture site and apply in a circular fashion using friction

Inactive ingredients

purified/deionized water, ethyl ester of PVMA/MA copolymer, ethyl alcohol SDA-40B, acetyl tributyl citrate, chloroxylenol

PRINCIPAL DISPLAY PANEL – packet label

REORDER NO. ZA50075

LATEX FREE

NDC 52641-700-02

ReliaMed

I.V.
Antiseptic Wipes

For preparation of the skin prior

to injection or venipuncture

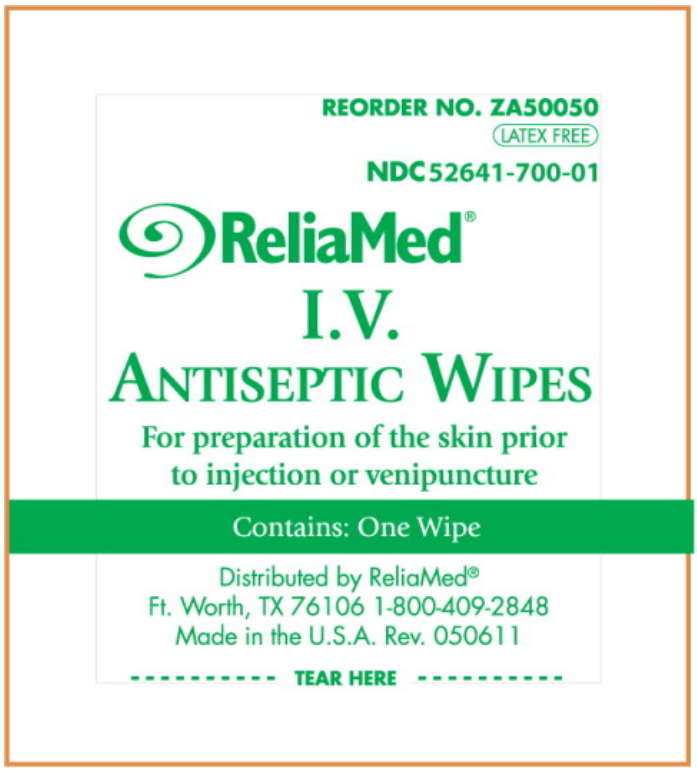
Contains: One Wipe

Distributed by ReliaMed

Ft. Worth, TX 76106 1-800-409-2848

Made in the U.S.A. Rev 050611

TEAR HERE



PRINCIPAL DISPLAY PANEL – box label

NEW

ReliaMed

REORDER NO. ZA50075

LATEX FREE

NDC 52641-700-02

I.V. Antiseptic Wipes

For preparation of the skin prior to injection or venipuncture

75 Individual Wipes

- Protect skin from adhesives
- Non-irritating to intact skin
- Thick, soft application pad

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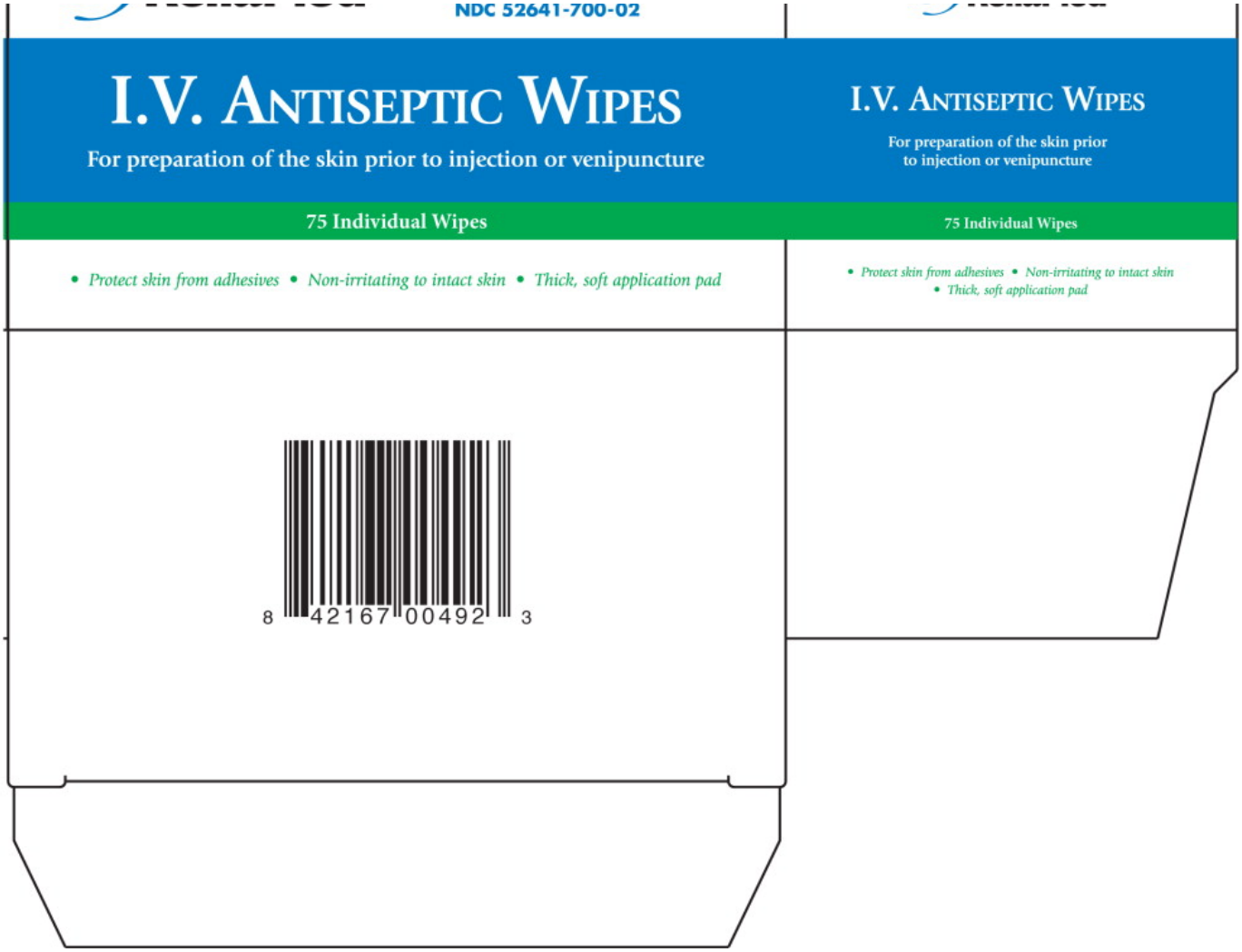
ReliaMed®



ANTISEPTIC

isopropyl alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:52641-700
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
isopropyl alcohol (UNII: ND2M416302) (isopropyl alcohol - UNII:ND2M416302)	isopropyl alcohol	0.56118 g in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
water (UNII: 059QF0KO0R)	
alcohol (UNII: 3K9958V90M)	
acetyltributyl citrate (UNII: 0ZBX0N59RZ)	
chloroxylenol (UNII: 0F32U78V2Q)	
ethyl nitrate (UNII: E1ZT886LR5)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52641-700-02	75 in 1 BOX	06/23/2011	
1		1.2 mL in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:52641-700-01	50 in 1 BOX	06/23/2011	
2		1.2 mL in 1 PACKET; 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	06/23/2011	

Labeler - ReliaMed (049386139)

Registrant - Safetec of America, Inc. (874965262)

Establishment

Name	Address	ID/FEI	Business Operations
Safetec of America, Inc.		874965262	MANUFACTURE(52641-700)

Revised: 6/2013 ReliaMed