A1052-22 AMNIOCENTESIS 22G QUINCKE - regional anesthesia kit Smiths Medical ASD, Inc.

APLICARE POVIDONE-IODINE SOLUTION (povidone-iodine solution) solution [Aplicare, Inc.]

3/4 Ounce Povidone Iodine Packet

Povidone-iodine 10%

Antiseptic

Warnings

Do not use

- if allergic to iodine
- in the eyes

For external use only

Ask a doctor before use if injuries are

- deep or puncture wounds
- serious burns

Stop use and ask a doctor if

- redness, irritation, swelling or pain persists or increases
- infection occurs

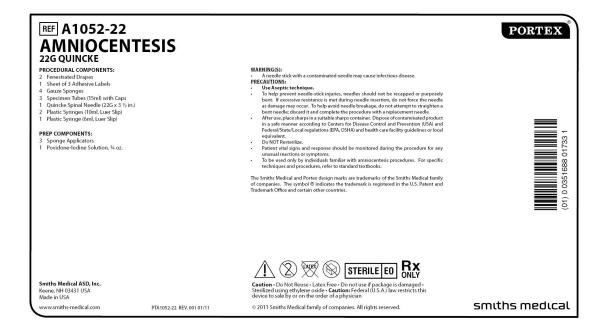
Avoid pooling beneath patient

Keep out of reach of children. In case of accidental ingestion, seek professional assistance or consult a poison control center immediately.

Package Label Display Panel



Package Label Display Panel



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AMNIOCENTESIS 22G QUINCKE

A1052-22 AMNIOCENTESIS 22G QUINCKE

regional anesthesia kit kit

Product Information

Product TypeMEDICAL DEVICEItem Code (Source)NHRIC:51688-1236

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NHRIC:51688-1236-2	10 in 1 CASE			
1		1 in 1 PACKAGE, COMBINATION			

Quantity of Parts				
Part #	Package Quantity	Total Product Quantity		
Part 1	1 PACKET	22.5 mL		

Part 1 of 1

APLICARE POVIDONE-IODINE

povidone-iodine solution

l	Product Information			
	Item Code (Source)	NDC:52380-0001		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PO VIDO NE-IO DINE (IO DINE)	PO VIDONE-IO DINE	0.10 mg in 1 mL

Inactive Ingredients			
Ingredient Name	Strength		
CITRIC ACID MO NO HYDRATE			
SO DIUM PHO SPHATE, DIBASIC			
SO DIUM HYDRO XIDE			
NONOXYNOL-9			
WATER			

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:52380-0001-3	22.5 mL in 1 PACKET				

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	03/01/1984		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
premarket notification	K965017	12/14/2005		

Labeler - Smiths Medical ASD, Inc. (137835299)

Establishment Name Address ID/FEI Business Operations Smiths Medical ASD, Inc. 137835299 relabel, manufacture

Establishment				
Name	Address	ID/FEI	Business Operations	
Aplicare, Inc.		107255002	manufacture	

Revised: 11/2012 Smiths Medical ASD, Inc.