ACL11SPS NERVE BLOCK - regional anesthesia kit Clint Pharmaceuticals, 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



REF ACL11SPS

To Reorder Call: 1-800-677-5022

PROCEDURAL COMPONENTS:

- PREP COMPONENTS:

NERVE BLOCK

- Needle-Pro and the color orange applied to the needle protection device are trademarks of the Smiths Medical family of co The symbol ® indicates the trademark is registered in the U.S. Patent and Trademark Office and certain other countries.

Manufactured for: Clint Pharmaceuticals 629 Shute Lane Old Hickory, TN 37138



Packaged in USA with US and imported parts PLTACL11SPS REV. 003 01/11

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Caution - Do Not Reuse - Latex Free - Do not use if package is damaged - Sterflized using ethylene oxide - Caution: Federal (U.S.A.) law restricts this device to sale by or on the order of a physician

Packaged in USA with US and imported parts

PLTACL11SPS REV: 003 01/11

ACL11SPS NERVE BLOCK

regional anesthesia kit kit

Product Information

MEDICAL DEVICE NHRIC:55553-468 Product Type Item Code (Source)

Packaging

	0 0			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NHRIC:55553-468-02	30 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



Product Information		
Item Code (Source)	NDC:52380-0001	
Route of Administration	TOPICAL	

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
PO VIDO NE-IO DINE (UNII: 85H0 HZU99 M) (IO DINE - UNII:9679 TC07X4)	PO VIDO NE-IO DINE	0.10 mg in 1 mL		

Inactive Ingredients				
Ingredient Name	Strength			
CITRIC ACID MO NO HYDRATE (UNII: 2968 PHW8 QP)				
SO DIUM PHO SPHATE, DIBASIC (UNII: GR686LBA74)				
SODIUM HYDROXIDE (UNII: 55X04QC32I)				
NONOXYNOL-9 (UNII: 48Q180SH9T)				
WATER (UNII: 059QF0KO0R)				

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	01/19/2009			

Labeler - Clint Pharmaceuticals, Inc. (609197785)

Registrant - Smiths Medical ASD, Inc. (137835299)

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

Esta	blishment			
	Name	Address	ID/FEI	Business Operations

Aplicare, Inc. 107255002 manufacture

Revised: 6/2012 Clint Pharmaceuticals, Inc.