ACL7 SINGLE SHOT EPIDURAL - regional anesthesia kit Clint Pharmaceuticals, Inc.

Drug Facts

APLICARE THREE POVIDONE-IODINE SWABSTICKS [Aplicare, Inc.]

Three 4-inch saturated swabsticks

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



REF ACL7

To Reorder Call: 1-800-677-5022

PREP COMPONENTS:

PROCEDURAL COMPONENTS:

SINGLE SHOT EPIDURAL

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

Caution - Do Not Reuse - Does not contain Natural Rubber Lates - Do not use if package is damaged - Sterifized using ethylene oxide - Caution. Federal (U.S.A.) law restricts this device soale by or on the order of a physical



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ACL7 SINGLE SHOT EPIDURAL

regional anesthesia kit kit

Product Information

MEDICAL DEVICE NHRIC:55553-454 Product Type Item Code (Source)

Packaging Item Code Package Description **Marketing Start Date Marketing End Date** 1 NHRIC:55553-454-02 30 in 1 CASE 1 1 in 1 PACKAGE, COMBINATION

Quantity of Parts Part# **Package Quantity Total Product Quantity** Part 1 1 PACKET 6.5 mL

Part 1 of 1

APLICARE POVIDONE-IODINE TRIPLES

povidone-iodine solution



Product Information		
Item Code (Source)	NDC:52380-3101	
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)	POVIDONE-IODINE	9.8 g in 96.4 mL	

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

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52380-3101-5	6.5 mL in 1 PACKET		

Marketing Information				
ı	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
l	OTC monograph not final	part333A	03/01/1998	

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

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.