ACL4KC SINGLE SHOT EPIDURAL - 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

To Reorder Call: 1-800-677-5022





SINGLE SHOT EPIDURAL **20G HUSTEAD**

PREP COMPONENTS

- Towel Povidone-lodine Solution, ¾ oz.

3 Sponge Applicators PROCEDURAL COMPONENTS:

1 Hustead Epidural Needle Calibrated

- with Wings (20G x 3 ½ in.) Pulsator® L.O.R. Syringe
- (7ml, Luer Slip) 18G x 1 ½ in. Needle
- 25G x 2 in. Needle
- 30G x 1 in. Needle
- Plastic Syringe (10ml, Luer Lock) Plastic Syringes (3ml, Luer Lock) Needle Stick Pad

- Gauze Sponges Fenestrated Drape

- WARNING(5):
 A needle stick with a contaminated needle may cause infectious disease.
 The use of excessive force while placing needles into the stick pad may cause the needle to protrude through the bottom of the tray which may result in a contaminated needle stick.

 PRECAUTIONS:
 Use Aseptic technique.
 To help prevent needle-stick injuries, needles should not be recapped or purposely bent. If excessive resistance is met during needle insertion, do not force the needle as damage may occur. To help avoid needle breakage, do not attempt to straighten a bent needle; discard it and complete the procedure with a replacement needle.
- needle. After use, place sharps in a suitable sharps container. Dispose of contaminated product in a safe manner according to Centers for Disease Control and Prevention, USA and Federal/State/Local regulations (EPA, OSHA) and health care facility guidelines or local equivalent:
- regulations (EPA, OSHA) and health care facility guidelines of local equivalent. Do NOT Resterilize. To be used only by individuals familiar with single shot epidural procedures, For specific techniques and procedures, refer to standard textbooks.

Pulsator is a trademark of Smiths Medical. The symbol @ indicates the trademark is registered in the U.S. Patent and Trademark Office and certain other countries.





SINGLE SHOT EPIDURAL 20G HUSTEAD \[\times_{\text{out}} \] \[\text{yvyy-mm} \]

RE ACL4KC







PTACL4KC REV. 002 09/11

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



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

FOLD

ACL4KC SINGLE SHOT EPIDURAL

regional anesthesia kit kit

Product Information

Product Type MEDICAL DEVICE Item Code (Source) NHRIC:55553-444

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NHRIC:55553-444-02	24 in 1 CASE		
1		1 in 1 PACKAGE, COMBINATION		

Quantity of Parts

	5			
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)				
SODIUM PHO SPHATE, DIBASIC (UNII: GR686LBA74)				
SODIUM HYDROXIDE (UNII: 55X04QC32I)				
NONOXYNOL-9 (UNII: 48Q180SH9T)				
WATER (UNII: 059QF0KO0R)				

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

Marketing InformationMarketing CategoryApplication Number or Monograph CitationMarketing Start DateMarketing End DateOTC monograph not finalpart333A03/01/1984

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

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

Revised: 6/2012 Clint Pharmaceuticals, Inc.