# ANTICOAGULANT SODIUM CITRATE- sodium citrate solution Medsep Corporation

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Anticoagulant Sodium Citrate Solution, USP Code 798-60

# ANTICOAGULANT SODIUM CITRATE SOLUTION, USP

Sterile, non-pyrogenic fluid path. Sterilized by steam.

# Rx only.

250 ml volume bag.

Store at room temperature.

# INDICATIONS AND USAGE

For use with automated plasmapheresis equipment only. Use according to equipment manufacturer's instructions. This unit should be used for plasmapheresis only.

#### **WARNINGS**

Not for direct intravenous infusion.

DO NOT VENT.

Avoid excessive heat. Protect from freezing.

#### **GENERAL PRECAUTIONS**

Do not use unless solution is clear and no leaks are detected.

Discard unused portion.

## **HOW SUPPLIED**

Single Use Container.

# INFORMATION FOR PATIENTS

Medsep Corporation, A Subsidiary of Pall Corp., Covina, CA 91722, USA

Visit us at www.pall.com/medical

For Pall customer service, call: 1.800.645.6578

#### CONTAINER TEXT DISPLAYED



# ANTICOAGULANT SODIUM CITRATE SOLUTION, USP

250 mL

Code 798-60

For use with automated plasmapheresis equipment only. Each 250 mL of anticoagulant contains 10.0 g sodium citrate (dihydrate), USP (pH adjusted with citric acid, USP). CAUTION: Not for direct intravenous infusion. Do not use unless anticoagulant is clear. This unit should be used for plasmapheresis only. Sterile, nonpyrogenic fluid path. Sterilized by steam. Rx Only.

Medsep Corporation
A Subsidiary of Pall Corporation
Covina, CA 91722, USA

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# Anticoagulant Sodium Citrate Bag Text

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Issued August 2010, 147798603

## ANTICOAGULANT SODIUM CITRATE

sodium citrate solution

Product Information				
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:62646-798	
Route of Administration	INTRAVENOUS			

Active Ingredient/Active Moiety				
Ingredient Name	<b>Basis of Strength</b>	Strength		
SODIUM CITRATE (UNII: 1Q73Q2JULR) (CITRIC ACID ANHYDROUS - UNII:XF417D3PSL)	SODIUM CITRATE	10 g in 250 mL		

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62646-798-60	250 mL in 1 BAG		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA760305	04/15/2011	

# **Labeler** - Medsep Corporation (928224765)

# **Registrant** - Medsep Corporation (928224765)

Establishment				
Name	Address	ID/FEI	Business Operations	
Medsep Corporation		928224765	manufacture, relabel, repack	

Revised: 5/2011 Medsep Corporation