SODIUM CITRATE- anticoagulant sodium citrate solution solution
Haemonetics Manufacturing Inc

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Anticoagulant Sodium Citrate Solution, USP

DESCRIPTION
Re-Order Product Code: 798-60
250 mL volume. Single use container.
Sterile, nonpyrogenic fluid path. Sterilized by steam.
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.

CONTRAINDICATIONS
Not for direct intravenous infusion.

WARNINGS
Avoid excessive heat. Protect from freezing. Do NOT vent.

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

General
Rx only. Discard unused portion.
Recommended Storage: Store at room temperature (25°C / 77°F).

HOW SUPPLIED
250 mL volume bag.
Each 250 ml contains:
10.0 g sodium citrate (dihydrate), USP

PACKAGE/LABEL PRINCIPAL DISPLAY PANEL
Bag Label
HAEMONETICS®
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.
DO NOT VENT.

References
HAEMONETICS and THE Blood Management Company are trademarks of Haemonetics Corporation.
Manufactured for:
Haemonetics Corporation
400 Wood Road
Braintree, MA 02184, USA
143798604

SODIUM CITRATE
anticoagulant sodium citrate solution solution
Product Information

Product Type: HUMAN PRESCRIPTION DRUG  
Item Code (Source): NDC:53157-798  
Route of Administration: INTRAVENOUS  
DEA Schedule: CII

Active Ingredient/Active Moiety

<table>
<thead>
<tr>
<th>Ingredient Name</th>
<th>Basis of Strength</th>
<th>Strength</th>
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</thead>
<tbody>
<tr>
<td>SODIUM CITRATE (UNII: 1Q73Q2JULR) (ANHYDROUS CITRIC ACID - UNII:XF417D3PSL)</td>
<td>SODIUM CITRATE</td>
<td>10.0 g in 250 mL</td>
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</tbody>
</table>

Packaging

<table>
<thead>
<tr>
<th>#</th>
<th>Item Code</th>
<th>Package Description</th>
<th>Marketing Start Date</th>
<th>Marketing End Date</th>
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</thead>
<tbody>
<tr>
<td>1</td>
<td>NDC:53157-798-60</td>
<td>36 in 1 CARTON</td>
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<tr>
<td>1</td>
<td></td>
<td>250 mL in 1 BAG; Type 0: Not a Combination Product</td>
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Marketing Information

Marketing Category: NDA  
Application Number or Monograph Citation: NDA760305  
Marketing Start Date: 06/30/1978  
Marketing End Date: |

Labeler - Haemonetics Manufacturing Inc (078598396)

Registrant - Haemonetics Manufacturing Inc (078598396)

Establishment

<table>
<thead>
<tr>
<th>Name</th>
<th>Address</th>
<th>ID/FEI</th>
<th>Business Operations</th>
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<tbody>
<tr>
<td>Haemonetics Manufacturing Inc</td>
<td>078598396</td>
<td>MANUFACTURE(53157-798) , STERILIZE(53157-798) , LABEL(53157-798)</td>
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Revised: 2/2015