

AMICUS Separator Kit Performance Report

Important: If reaction or injury has occurred call Fresenius Kabi Post-Market Quality Assurance at 1-800-933-6925.

Incident Date: Instrument S/N.: Software Version: UDI No.: Product Code: Lot No.:

When Was the Problem Detected?

Set Up Prime % Exchange Depletion Depletion/Exchange Reinfusion After Procedure/QC

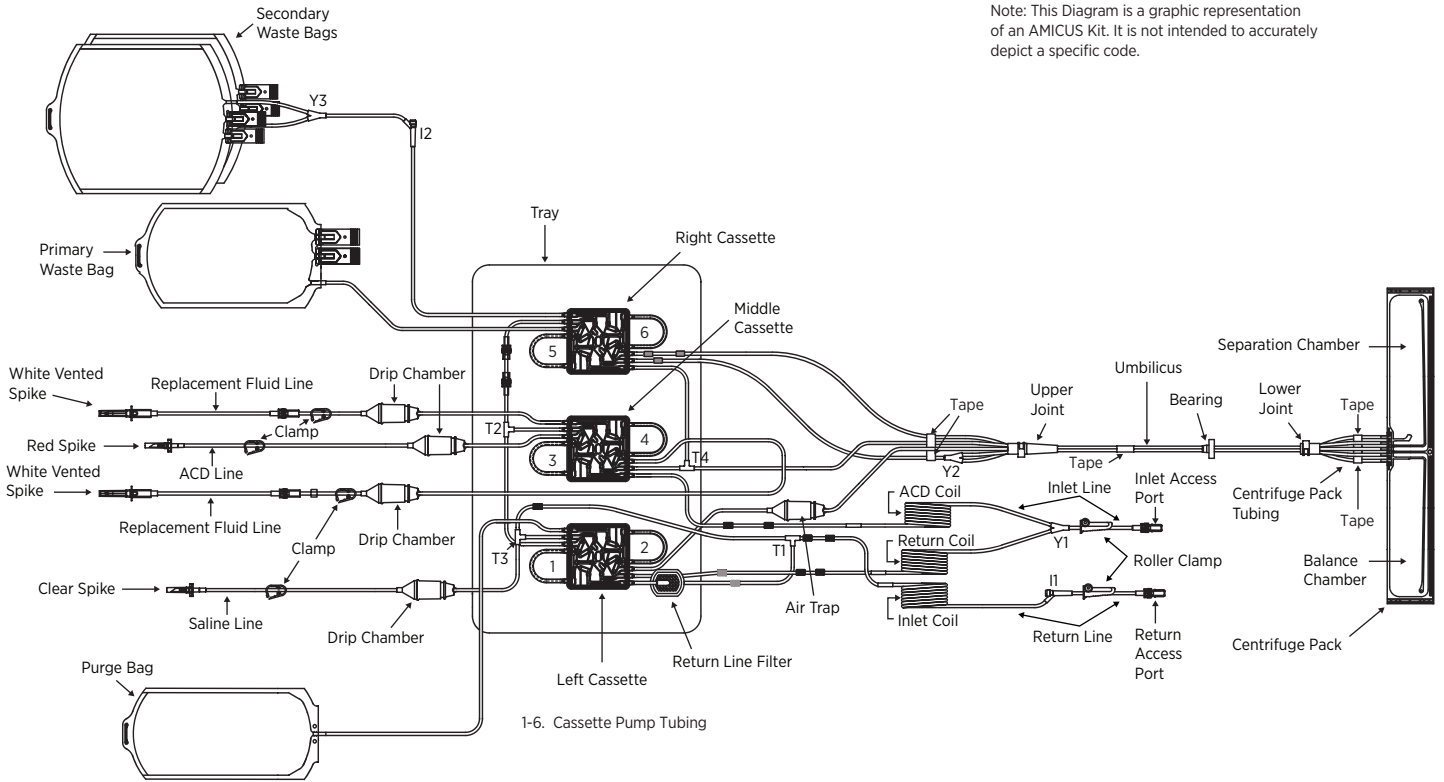
Problem Type (Mark all applicable) Cut/Slice Kink Particulate Matter Restriction/No Flow Alarms (Specify) Leak Separated Blood Leak Cracked Excess Air Missing RBC Contamination Other (Specify Below)

Please answer the following questions:

- 1. Was there any adverse event or injury? Yes No
2. If applicable, list name of any drug administered:
3. Was the procedure successfully completed? Yes No N/A
4. If no, was the procedure stopped due to a soft goods incident? Yes No N/A
5. Was product lost? Yes No N/A
Check box if you do NOT wish to receive response letters.

E-mail address for letter recipient (if applicable)

Please circle specific components on the diagram where incident occurred



Note: This Diagram is a graphic representation of an AMICUS Kit. It is not intended to accurately depict a specific code.

Additional Problem Description / Explanation

Kit Return To Fresenius Kabi

- 1. Sample available for evaluation? Yes No
2. Sample return box needed? Yes No Return label only
3. Picture available for evaluation? Yes No
Please e-mail a clear picture along with this report to MDComplaintSupport@fresenius-kabi.com

Center Authorized Signature/Date:

Customer Information (please print)

The following information is required to receive a credit

Facility Name: Contact Person: Account Number (if known): Operator Name: Street Address: City/State/Zip: Phone Number: Contact Person's E-mail:

Fax this report to 1-888-858-2983 or E-mail to MDComplaintSupport@fresenius-kabi.com and include a copy of this form when returning a kit.