Cue X6R5004 Product Performance Report

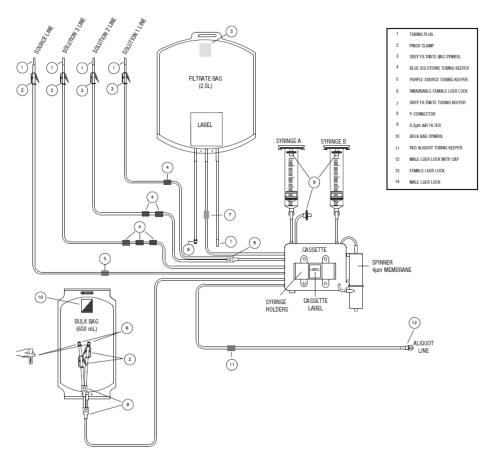


Important: If reaction or injury has occurred call Fresenius Kabi Post-Market Quality Assurance at 1-800-933-6925.

ncident Date: Instrument S/N.: Software Version: UDI No.: Lot No.: Spinner Video Jet No.:		
Problem Detected with: □ Primary Set □ Sampling Set(s)* □ Both *Note: Each Primary Set includes three, individually packaged Sampling Sets. Sampling Sets are not available as a separate product code.		
Primary Set (Complete this section if problem was detected with Primary Set)		
When Was the Problem Detected? ☐ Before Use ☐ Set Installation ☐ Set Checks ☐ Set Prime ☐ During Procedure ☐ After Procedure		
Problem Type (Mark all applicable) Packaging: □ Packaging Open □ Mispacked □ Discolored □ Missing or Illegible Label □ Missing or Separated Component (e.g., keeper) □ Tubing: □ Flattened □ Kinked □ Hole □ Cut/Sliced □ Blocked/Occluded □ Discolored □ Incorrect Length □ Separated (e.g. from Cassette) □ Spinner (Separation Device): □ Cracked □ Leaking □ Noise Cassette: □ Cracked □ Hole □ Leaking □ Poor Fit in Cassette Enclosure Syringe Assemblies: □ Cracked □ Leaking □ Missing or Separated Component □ Poor Fit on Syringe Guides Air Filter: □ Cracked □ Leaking □ Discolored □ Blocked/Occluded Swabbable Female Luer Lock: □ Cracked □ Leaking □ Discolored □ Blocked/Occluded Container: □ Hole □ Improper Seal around Container Port □ Leaking □ Discolored □ Missing Label		
Other: Associated Alert Name/Code (if applicable):		

Please circle specific components on the diagram where incident occurred

Cue X6R5004 Primary Set



Sampling Set (Complete this section if problem was detected with S	Sampling Set)	
When Was the Problem Detected? ☐ Before Use ☐ During Use ☐ After Use		
Component (e.g., pinch clamp) Tubing: Flattened Kinked Hole Cut/Slice Pouch: Hole Cut/Sliced Leaking Discolored B Male/Female Luer Locks: Cracked Component (e.g., pinch clamp) Kinked Hole Cut/Sliced Leaking Discolored B Male/Female Luer Locks: Cracked Leaking Discolored D	locked/Occluded ☐ Poor Fit	
Other:		
Please circle specific components on the diagram where incide	ent occurred	
Cue Sampling Set		
9 (3) (1) (2) POUCH (~6 mL) (3)		
Additional Problem Description / Explanation		
Picture available for evaluation? Yes □ No □ If a picture is available, please e-mail a clear picture along with this report to mdpmqa.usa@fresenius-kabi.com		
Please answer the following questions: 1. Was there any adverse event or injury? Yes No 2. Was the procedure successfully completed? Yes No N/A 3. If no, was the procedure stopped due to a soft goods incident? Yes No N/A 4. Was product lost? Yes No N/A 5. Did the procedure involve clinical or patient material? Yes No N/A Check box if you do NOT wish to receive response letters. E-mail address for letter recipient (if applicable)		
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Kit Return to Fresenius Kabi 1. Sample available for evaluation? Yes \(\scale \) No \(\scale \) 2. Return label needed? Yes \(\scale \) No \(\scale \) 3. Sample return box needed? Yes \(\scale \) No \(\scale \) Center Authorized Signature/Date:	Customer Information (please print) The following information is required to receive a credit Facility Name: Contact Name: Account Number (if known): Operator Name: Street Address:	
Fax this report to 1-888-858-2983 or E-mail this report to mdpmqa.usa@fresenius-kabi.com and include a copy of this form when returning a kit.	Street Address:	