LOVO X6R4902 Product 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.: Lot No.:	Software Version: UDI No.:
When Was the Problem Detected? □ Before Use □ Kit Installation □ Disposable Kit Check □ I	Disposable Kit Prime □ During Procedure □ After Procedure
Problem Type (Mark all applicable) Packaging: □ Packaging Open □ Mispacked □ Illegible Labe Tubing: □ Flattened □ Kinked □ Hole □ Cut/Sliced □ Container: □ Leaking Fluid □ Improper Seal around Containe Other:	er Port Discolored
Additional Problem Description/Explanation	
	
Please circle specific components on the diagram where incident occurred	
ANCILLARY BAG	PROCESSING KIT CONNECTION POINT
Picture available for evaluation? Yes \square No \square If a picture is available, please e-mail a clear picture along with the	is 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 □ 3. If no, was the procedure stopped due to a soft goods incident? 4. Was product lost? Yes □ No □ N/A □ 5. Did the procedure involve clinical or patient material? Yes □ No □ Check box if you do NOT wish to receive response letters. □	
	-mail address for letter recipient (if applicable)
Kit Return to Fresenius Kabi 1. Sample available for evaluation? Yes □ No □ 2. Return label needed? Yes □ No □ 3. Sample return box needed? Yes □ No □	Customer Information (please print) The following information is required to receive a credit Facility Name: Contact Name:
Center Authorized Signature/Date:	Account Number (if known): Operator Name: Street Address: City/State/Zip:
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.	Phone Number: Contact Person's E-mail:

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