

Urgent Medical Device Correction

Kangaroo Enteral Feed Pumping Sets

March 1, 2021

of the

problem:

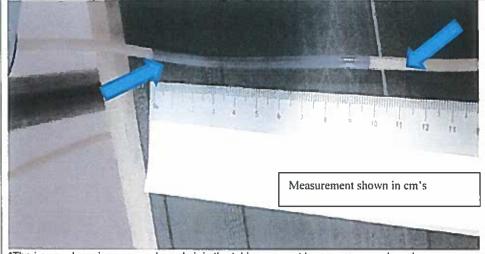
Dear Valued Customer,

Cardinal Health is initiating a Medical Device Product Correction to advise you of the potential for air appearing in the enteral feed pumping set tubing during set-up. The purpose of this communication is to advise you of this situation and to provide steps to help address the issue prior to the availability of the corrected product.

Product Correction	Cardinal Health has initiated a field safety corrective action for the following Kangaroo enteral feed pump sets.		
Overview	Product Code	Product Description	Affected Lots
	612054	Kangaroo™ ePump INT. 500ML Feed Only ST	
	613650	Kangaroo™ ePump INT. 1000ML Feed Only ST	
	614650	Kangaroo™ ePump INT. Proximal Spike ST	
	672055	Kangaroo™ ePump INT. 500ML Feed Only NS	
	673656	Kangaroo™ ePump INT. 1000ML Feed Only NS	
	673662	Kangaroo™ ePump INT. 1000ML Feed/Flush NS	
	674655	Kangaroo™ ePump INT. Proximal Spike NS	
	674668	Kangaroo™ ePump Joey 500ML Feed/Flush DEHP Free	
	674669	Kangaroo™ ePump INT. Proximal Spike and Flush NS	
	716154	Kangaroo™ ePump 100ML Burette Sterile	
	762055	Kangaroo™ Joey 500ML Pump Set	All product manufactured before Sep 3, 2020
	763656	Kangaroo™ Joey 1000ML Pump Set	(lot 202580052 and below)
	763662	Kangaroo™ Joey 1000ML Feed/Flush Set	
	765100	Kangaroo™ Joey ENPlus Spike Feed and Flush 1000ML	
	765559	Kangaroo™ Joey ENPlus Spike Set	
	772055	Kangaroo™ ePump DEHP Free 500ML Feed Only	
	773656	Kangaroo™ ePump DEHP Free 1000ML Feed Only	
	773662	Kangaroo™ ePump DEHP Free 1000ML Feed and Flush	
	775100	Kangaroo™ ePump ENPlus Spike Feed and Flush Set 1000ML	
	775659	Kangaroo™ ePump ENPlus Spike Pump Set	
	776150°	Kangaroo™ ePump Burette Re-certification Burette 100ML*	
	*Although this product code is not used for feeding, the air in tubing issue may affect the recertification/calibration process and the same steps in Attachment A should be followed.		
	<u>Usage</u> Kangaroo Enteral Feed Pump sets are used for enteral feeding either as feed/flush pump sets or feed only pump sets.		
Why you are being contacted:		eiving this letter because our records indicate to nteral Feed Pump sets in the past.	hat you have purchased Cardinal Health
Description	What is the	issue?	

Directo

The issue relates to air entry into the feed set tubing, which appears as air space in the tubing.



*The image above is an example and air in the tubing may not be as pronounced as shown.

Why are we sending this Product Correction?

Air entry into the feeding set tubing could result in the following health consequences: vomiting, dehydration, hypoglycemia, abdominal pain, and abdominal distention in patients.

What other actions is Cardinal Health taking?

Cardinal Health is implementing the appropriate corrective actions to ensure that the issue is corrected.

Actions requested on your part:

- FOLLOW the steps described in Attachment A (Air in Tubing Check Process) to check the feed set.
- COMMUNICATE with all personnel that utilize these feeding sets of the potential risk of air entering feeding set tubing.
- 3) NOTIFY any customers to whom you may have distributed/forwarded affected product to or will send the product on to about this product correction notice and share a copy of this notice and Attachment A.
- 4) POST a copy of this notification in your storeroom where the product is stored

Available Assistance:

Please contact the Customer Service group for any questions related to this action or to return defective product:

- Hospital—800-964-5227
- Federal Government—800-444-1166
- Distributor—800-635-6021
- All other customers—888-444-5440

For questions related to the notification and/or acknowledgement form that are not adequately addressed in this letter, please contact the market action team at: GMB-FieldCorrectiveAction@cardinalhealth.com or call 800-292-9332.

Additional Information:

Adverse Events Reporting Process

Cardinal Health has notified the U.S. Food & Drug Administration that we are taking this action. In the event you have experienced quality problems or adverse events related to the products listed above, send an email to: GMB-PRComplaints@cardinalhealth.com.

The FDA can be contacted to report any adverse events experienced with these products:

 Online @ http://www.fda.gov/Safety/MedWatch/HowToReport/default.htm (form available to fax or email) or call FDA 1-800-332-1088.

Regulatory Notification

The FDA and other applicable regulatory bodies have been notified and are aware Cardinal Health is voluntarily taking this action.

We apologize for any inconvenience this communication may cause. We know that you place high value in our products, and we appreciate your cooperation in this matter. Cardinal Health is committed to maintaining your confidence in the safety and quality of the products that we supply.

Respectfully yours,

Moazzam Khan Director, QRA Management

Event ID: Event-2020-03389



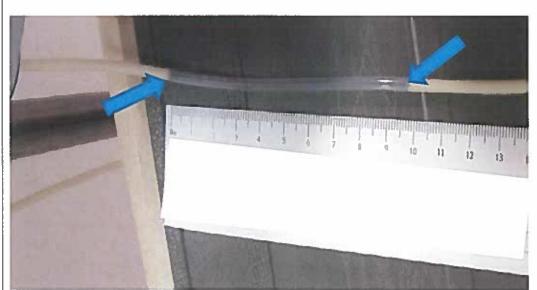
Attachment A (Air in Kangaroo Enteral Feed Pump Sets Tubing Check Process)

To check the feed set for entry of air in the tubing (see the example below) follow the steps described below.

- Press "Run" on the pump after installing and priming the feeding set.
- After 30 seconds, observe the tubing between the pump and the patient to check for air being drawn into the line.
- It is not uncommon, and not a problem, to see a small amount of air in the feed tubing (approximately 1.3 cm [1/2 inch]) after the pump checks for any blockages.
- Do not use the feeding set if you see a single segment of air measuring greater than 2.5 cm (1 inch).
- Do not use the feeding set if you see several air bubbles that cumulatively measure greater than 2.5 cm (1 inch)

NOTE:

The air in tubing event is found to be intermittent and is not observed in all lots. Thus, if a defective feeding set is identified, it does not necessarily mean that the entire lot is defective.



*This is an example and the air in the tubing may not be as pronounced as shown in the image.