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WEEE-Reg.-Nr. DE 36963167 VAT-ID: DE 120679179



Field Safety Notice

Name of the affected product: 3M™ Ranger™ Blood/Fluid Warming High Flow Sets, catalog

numbers 24355 and 24370

FSCA-identifier: 2022-11 FSCA Ranger **Type of action:** Disposal of affected products

Date: November 7th, 2022

Attention: 3M Health Care Business Customers

Dear Customer,

3M is notifying all customers in impacted countries of the above-mentioned 3M™ Ranger™ Blood/Fluid Warming High Flow Sets, catalog numbers 24355 and 24370 of a field safety corrective action.

Description of the problem and potential hazard and risk for the patient/user:

This corrective action has been initiated due to the identification of a manufacturing issue with the auto-venting bubble trap. There is the risk of a blood or fluid leak while priming the sets and/or during fluid administration. The products could be unable to deliver the therapy as per intended use and could expose the user to blood, blood products, and IV fluids during a leak.

Details on affected devices:

The following lots of 3M™ Ranger™ Blood/Fluid Warming High Flow Sets are subject to this field safety corrective action and were supplied after March 22nd, 2022:

Catalogue number	Lot number
24355	HX9137, HX9158, HX9162, HX9167, HX9169, HX9179, HX9181, HX9183, HX9184, HX9189, HX9190, HX9192, HX9198, HX9200, HX9202, HX9204, HX9214
24370	HM9186, HM9196, HM9213

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Action to be taken by all customer:

- 1. Ensure all your internal and external customers are informed about this corrective action.
- 2. Please inform us if you supplied the affected items to customers outside your home country.
- 3. Please identify the affected product listed above and immediately cease use of affected models and lots of 3M[™] Ranger[™] Blood/Fluid Warming High Flow Sets.
- 4. Please discard all remaining affected product listed above per facility procedures.
- 5. Complete and return by e-mail to meddev.de@mmm.com the enclosed Acknowledgement Form, indicating that the corrective action was understood and executed. Please also indicate the number of devices you have disposed of.

Transmission of this Field Safety Notice:

Please pass on this notice immediately to all departments who might use the concerned products. In addition, ensure that the information is provided to any organisation where the concerned products potentially have been distributed.

Thank you for your immediate attention and cooperation. We apologise for any inconvenience this matter may cause.

Contact reference person:

If you have questions, please contact your local 3M representative.

The undersigned confirms that this notice has been notified to the appropriate Regulatory Agency.

Dr. Marie Isabel Cobbers Safety Officer 3M Deutschland GmbH, Health Care Business Carl-Schurz-Strasse 1, 41453 Neuss, Germany Mail: meddev.de@mmm.com 3M Deutschland GmbH page 3

Acknowledgement Form – FSN 2022-11 FSCA Ranger

Email completed form to: meddev.de@mmm.com

Please examine your inventory to determine if you have any of the 3M™ Range	r™ Blood/Fluid
Warming High Flow Sets listed. Note, this corrective action only applies to the affe	ected catalogue
numbers and lots listed and does NOT apply to any other 3M™ Ranger™ produc	ts.

		Quantity of sets in
	ve examined our inventory and identified and dispos ^M Ranger™ Blood/Fluid Warming High Flow Sets:	ed of the following number
	wledge that you have read and understood this lette s requested.	r and will complete the

Catalogue number	Lot number	Quantity of sets in inventory that have been disposed
24355	HX9137, HX9158, HX9162, HX9167, HX9169, HX9179, HX9181, HX9183, HX9184, HX9189, HX9190, HX9192, HX9198, HX9200, HX9202, HX9204, HX9214	
24370	HM9186, HM9196, HM9213	

We have examined our inventory and do not have the affected 3M™ Ranger™
Blood/Fluid Warming High Flow Sets in stock.

Email completed form to: meddev.de@mmm.com

Person completing this form:

Name	Company /Hospital Name	
Signature	City, Country	
Date	Phone	
	E-mail	