smiths medical
bringing technology to life

6000 Nathan Lane N.
Minneapolis, MN 55442
https://www.smiths-medical.com

URGENT: FIELD SAFETY NOTICE

Jelco® Optiva® IV Catheter 5063-AI

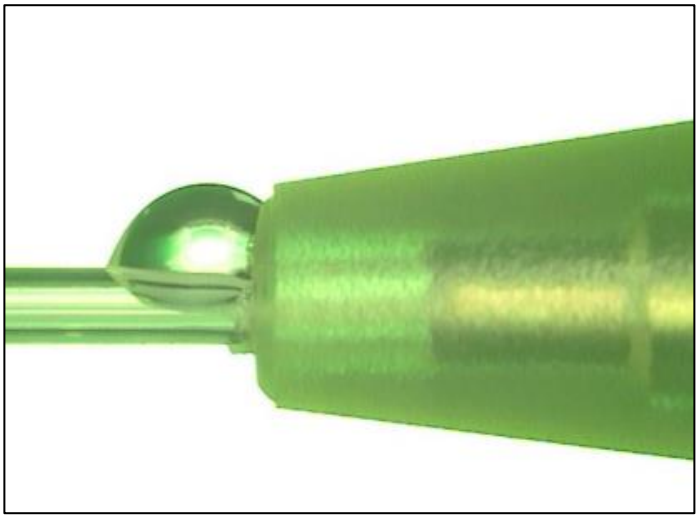
13th April 2023

Dear Valued Customers:

Smiths Medical, Inc. is issuing this Urgent Field Safety Notice to notify you of a potential defect with the Jelco® Optiva® IV Catheters. This letter details the issue and the required steps for you to perform.

Issue:
Smiths Medical has identified the potential for a manufacturing defect within specific lots of the 24-gauge Jelco® Optiva® IV catheters which may result in leakage at the insertion site.

Potential Risk:
The potential leakage is the result of catheter damage at proximal end of the catheter and near the tip of the female luer connector. A potential concern would be any fluid loss that would disrupt routine fluid delivery, drug delivery or administration of blood. To date, Smiths Medical has not received any reports of serious injury or death associated with this issue.



Affected Product:
Our records indicate that you may have received some of the affected products, which were distributed in Ireland in November 2022. The affected item and lot numbers are provided in Table 1, below:

Table 1: Affected Product and Lot Numbers

List Number	Product Description	Lot Number
5063-AI	Jelco® Optiva® IV Catheter - 24G X 19mm	4331853

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Required Actions for Users:

1) Please discontinue the use and distribution of the affected product immediately. Check your inventory and quarantine all affected product at your facility.

2) Inform potential users of the product in your organization of this notification and complete the attached response form, even if you do not have affected product. Please indicate on the form whether you intend to return this product to Smiths Medical or destroy it locally and return the completed response form to EMEA-Quality@icumed.com .

3) If you have distributed the product further, immediately notify your accounts that received the product identified in the Affected Product / Table 1 sections of this notification and ask your customers to complete a response form and return to you for overall completion.

4) Please contact Customer Service using the information provided below for assistance reordering replacement product.

Follow up Actions by Smiths Medical:
Upon receipt of the affected product, or upon receipt of a Certificate of Destruction, Smiths Medical will credit you for any product returned/destroyed. You will only receive credit for product that you return or that you certify has been destroyed locally. NOTE: Credits for product purchased through a distributor will be credited by the distributor.

For further inquiries, please contact Smiths Medical using the information provided below.

Smiths Medical Contact	Contact Information	Areas of Support
Global Complaint Management	globalcomplaints@icumed.com	To report adverse events or product complaints
Customer Support	ukcs@icumed.com	Additional information or assistance

HPRA has been notified of this action.

Smiths Medical is committed to patient safety and is focused on providing exceptional product reliability and the highest level of customer satisfaction. Thank you for your prompt support on this important matter. We appreciate your cooperation.

Sincerely,

Joe Canavan
Vice President, Quality, Consumables

Enclosures:
- Customer Response Form (pages 3 of this notice)
- Certificate of Destruction (separate file)

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URGENT: FIELD SAFETY NOTICE – RESPONSE FORM

Jelco® Optiva® IV Catheter 5063-AI

13th April 2023

Check your inventory and complete the information below, even if you do not have the affected product. *Failure to complete all sections of this page may result in improper, delayed or denied credit.*

Please return the completed form to EMEA-Quality@icumed.com, Smiths Medical Customer Service and your local sales representative.

Name of Hospital / Facility	
Hospital / Facility Address	
Telephone Number	
Name and Title of Person Completing this Form	
Signature of Person Completing this Form	
Date	
If Purchased through a distributor, please list distributor name/location here for traceability purposes	

Please select one:

☐ I have **NO** affected products (complete and return this form to the e-mail address above)

☐ **YES**, I have affected products

If you have affected product on hand, please complete table below:

TABLE 1

List Number	Lot Number	Quantity in inventory	PO, debit memo or invoice

If you have distributed the product further, please complete table below with collated information received from your customers and respond to ICU Medical with the overall information.

TABLE 2

List Number	Lot Number	Quantity destroyed locally by customer	Quantity returned to distributor

☐ I have followed the instructions provided to me and I will **destroy** affected products on site (complete and return provided Certificate of Destruction to the email addresses on the certificate).

☐ I have followed the instructions provided to me and I will contact my Smiths Medical CS Representative to make arrangements to **return** the affected products.

☐ I have followed the instructions provided to me. If affected product is not being returned, please explain below:

Adverse events and complaints associated with the use of this product should be reported and emailed to the Competent Authority or to Smiths Medical's Global Complaint Management Department at globalcomplaints@icumed.com.