URGENT MEDICAL DEVICE RECALL

August 19, 2026

Dear Valued McKesson Customer:

BD has notified McKesson Medical-Surgical Inc. (MMS) of an Urgent Medical Device Recall regarding specific lot(s) of their Bard Silastic Foley Catheters. This notice has been issued to inform customers that there were errors reported in the affected product table used to identify affected product. BD wants to assure that all affected product within your facility has been properly identified and dispositioned. Affected product first shipped April 22, 2025.

McKesson Medical-Surgical Inc. has taken appropriate action per this notice.

For questions regarding this notification, please contact BD at **(844) 823-5433**.

A review of our records indicates that your company may have purchased items included in this notification. Carefully review the information in this letter and follow the instructions provided below.

Refer to the table below for a list of affected item(s) and lot number(s) distributed by McKesson Medical-Surgical

MMS #	MFG Catalog #	Description	Affected Lot(s)	Exp. Date
33069	33620 X	CATHETER, FOLEY SILAS 20FR 5CC	NGKS5060 NGKT3308 NGKV3689	1/31/2029 11/30/2029 11/30/2029
33066	33614 X	CATHETER, FOLEY SILAS 14FR 5CC	NGKN4037 NGKP1420 NGKQ0040 NGKQ3901 NGKS4937 NGKV3462 NGKW3897 NGKT3258	1/31/2029 9/30/2029 9/30/2029 12/31/2028 9/30/2029 6/30/2029 6/30/2029 9/30/2029
240490	33618 ✓	CATHETER, FOLEY SIL 18FR 5CC (10/CS)	NGKQ3370 NGKS3753 NGKT3258 NGKT3309 NGKU4129 NGKV1446 NGKV2345 NGKV3478 NGKV5029 NGKV0617	5/31/2029 9/30/2029 9/30/2029 9/30/2029 9/30/2029 7/31/2029 7/31/2029 7/31/2029 7/31/2029 9/30/2029
241211	33622 X	CATHETER, FOLEY SIL 22FR 5CC (10/CS)	NGKQ1016 NGKS0244 NGKU0967 NGKV5075 NGKX0441	1/31/2029 1/31/2029 9/30/2029 1/31/2030 1/31/2030
33060	33418 X	CATHETER, FOLEY SILAS 30CC	NGKW2813	1/31/2029
33061	33420 X	CATHETER, FOLEY SILAS 30CC	NGKQ1015	1/31/2029
241212	33616 X	CATHETER, FOLEY SIL 16FR 5CC (10/CS)	NGKQ0042 NGKQ3946 NGKR3420 NGKT3281 NGKU1015 NGKV0617 NGKV2275 NGKV2309 NGKW3771	9/30/2029 1/31/2029 9/30/2029 9/30/2029 11/30/2029 9/30/2029 9/30/2029 9/30/2029 9/30/2029

McKesson Customer Instructions:

1.) Immediately discontinue use of any product matching the affected item(s) and lot number(s) listed above. If you have no product matching the affected item(s) and lot number(s), no further action is needed.

2.) A copy of the Urgent Medical Device Recall from BD has been included for reference.

3.) If you have product affected by this notice, fill out the McKesson Reply Form and return it to our Corporate Customer Service Center via email at MMSRecalls@McKesson.com or fax at **(866) 871-0270**. To ensure timely credit to your account and support the completion of this notice, please respond within 30 days. **Please note**: Credit will only be issued for the affected Lot(s) of the product(s) listed above and entered on the reply form. **Please place a new order for replacement product if there is an immediate need**.

4.) Please dispose of any affected product in your possession in accordance with your institution's policies and procedures.

5.) If you have further distributed any of the item(s) referenced in this notification, provide your accounts with a copy of this notification.

We sincerely apologize for any inconvenience this notice may have caused you and your staff. If you have any questions about information provided in this communication, please contact our **McKesson Medical-Surgical Recall Message Center** at MMSRecalls@McKesson.com or call **(800) 688-8840**.

Thank you for your prompt attention,

McKesson Medical-Surgical Inc.

McKesson Medical-Surgical Inc.
Medical Device Recall Reply Form: RC-2026-087A
BD - Bard Silastic Foley Catheters

August 19, 2026

Complete this reply form and return all pages immediately via email to MMSRecalls@McKesson.com or fax at **(866) 871-0270** should you have affected product.

To ensure timely credit to your account and support the completion of this notice, please respond within 30 days.

Date: _____

Your Name: _____ Email Address: _____

Phone Number: _____ Fax Number: _____

Account: 20037573 District: 22610000
CARELINC MEDICAL EQUIP & SUPP
ATTN: RISK MANAGEMENT
89 54TH ST SW STE 1
GRAND RAPIDS, MI 49548-5503

☐ I acknowledge that I DO HAVE product affected by this notification and have disposed of this product in accordance with my institution's policies and procedures.

Qty	Unit of Measure	Affected Lot #(s)	MMS #	MFG Catalog #	Description
			33069	33620	CATHETER, FOLEY SILAS 20FR 5CC
			33066	33614	CATHETER, FOLEY SILAS 14FR 5CC
			240490	33418	CATHETER, FOLEY SIL 18FR 5CC (10/CS)
			241211	33622	CATHETER, FOLEY SIL 22FR 5CC (10/CS)
			33060	33418	CATHETER, FOLEY SILAS 30CC
			33061	33420	CATHETER, FOLEY SILAS 30CC
			241212	33616	CATHETER, FOLEY SIL 16FR 5CC (10/CS)

*Discard Affected lot numbers only

*** The affected lot number(s) are listed on the McKesson customer letter. Please dispose of affected product in accordance with your institution's policies and procedures.**

Credit will only be issued for product(s) listed above with affected lots.

If you have any questions about information provided in this communication, please contact the McKesson Recall Message Center at MMSRecalls@McKesson.com or call (800) 688-8840.

See instructions on the reverse side of this form to access
McKesson Medical-Surgical's online product ordering system, "SupplyManager", for a fillable form.

Instructions for the McKesson Medical-Surgical online product ordering system – "SupplyManager", to access and download a "fillable" PDF reply form.

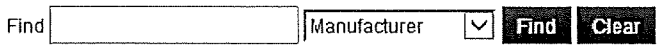
1) It is important to download the correct reply form for the specific recall you are responding to.
    a. Reply forms have a specific designation, example: RC-202X-XXX.
        - "202X" is the recall year, and "XXX" is the 3-digit unique numeric identifier for the recall.

2) Go to https://mms.mckesson.com/ and log in to "SupplyManager", with your username and password.

3) On the home page, under 'Essential Tasks' click 'Your Account.' Under 'Resources' a support link titled "Product Recalls" can be found on the right side.

4) Click on the hyperlink "Product Recalls" (this will open a listing of recalls for the last 3 months).

5) On the recalls list page, locate the "Find" box.
    a. From the drop-down options, select one of the following: "Keyword", "McKesson Item #" or "Manufacturer".
    b. Enter a Keyword, McKesson Item #, or Manufacturer name in the Find box and click "Find".



    c. A list of issued recalls will be made visible for you to select from.

6) Click on the blue hyperlink, found under the heading "Recall Notice," for the notice details you want to access.

7) The PDF customer documents associated with the notice will be displayed, *this includes the reply form to download and complete for your response.*

8) Click on the hyperlink(s) to open the Customer Document(s). Save/Download this document to your computer. Once the documents are saved, close out the document window.

9) Submit completed reply form to MMSRecalls@McKesson.com.

10) If you wish to view additional recalls, return to the home recall page by clicking the blue "View All Recalls" button at the upper right corner of the page above the blue alert banner.

BD (C.R. Bard) Urology and Critical Care
8195 Industrial Blvd
Covington, GA 30014
www.bd.com

BD
Advancing the world of health

URGENT Medical Device Recall

Bard® Silastic® Foley Catheters

August 13, 2026

For the Attention of: Recall Coordinator, Director of Nursing, Director of Purchasing, Director of Risk Management

This is not a new or expanded recall. This notice is to inform you that there were errors reported in the affected product table used to identify affected product (Attachment 2). This letter provides a corrected affected product table with the items that have been updated highlighted. This is being provided to assure that all affected product within your facility has been properly identified and dispositioned.

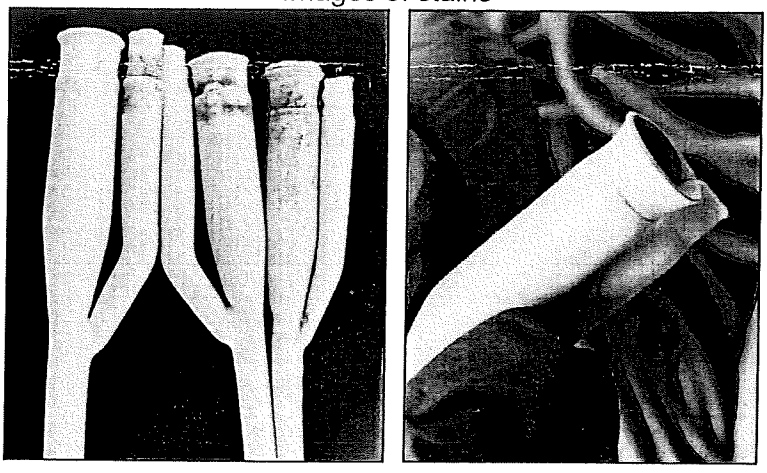
The prior letter sent 23 April 2026 was to inform you that C.R. Bard® Urology and Critical Care, a wholly owned subsidiary of Becton, Dickinson and Company (BD), is voluntarily initiating a correction for the devices listed below. All product and associated lots listed in Attachment 2 (Affected Product) are impacted by this correction. The following products are included:

- Bard® Silastic® Foley Catheter (French sizes 16, 18, 20, 22, 24) 30cc balloons
- Bard® Silastic® Foley Catheter (French sizes 14, 16, 18, 20, 22, 24) 5cc balloons

Description of the problem:

BD has received complaints, and confirmed through internal inspection that stains are present on the surface of the affected silastic catheters. The cause of the stains was determined to be an increased amount of a raw material used in the manufacturing process. This raw material may be harmful if present in increased amounts on the surface of the catheter. Please see images below.

Images of stains



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Complaint & Adverse Event Statement

To date, there has been 1 (one) complaint and no adverse events reported worldwide related to this issue.

Clinical Impact:

The current issue may result in the use of urinary catheters that exhibit staining on the catheter surface. Use of an affected device could potentially lead to local irritation, erythema at or near the area of contact, pain, dysuria, hematuria, or a burning sensation. These events are not considered life-threatening; however, they represent patient injuries that may require evaluation or medical intervention.

Advice for Clinical Users:

For any patients requiring Foley catheterization, an alternative urinary catheter should be used. If an impacted catheter is currently in place and continued catheterization is required, the device should be replaced with an alternative Foley catheter as soon as clinically appropriate.

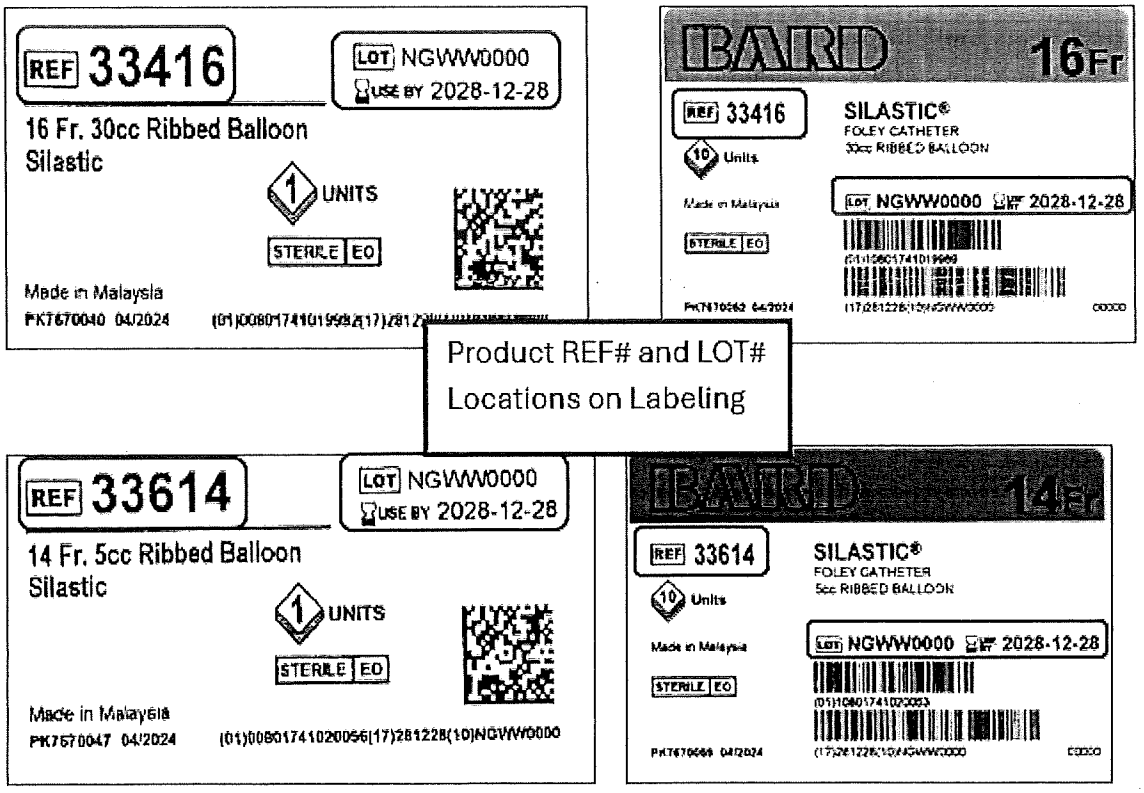
If devices have been safely used without issue, no additional treatment or clinical follow up is required.

Distribution Timeframe and Label Photo:

The affected product was distributed from 22 Apr 2025 – 28 Jan 2026

Product Label

Case Label



BD (C.R. Bard) Urology and Critical Care
8195 Industrial Blvd
Covington, GA 30014
www.bd.com

Please Take the Following Actions:

1. **Immediately Discontinue Use**
2. Please check all inventory locations within your institution for the Bard® Silastic® Foley Catheters listed in Attachment 2.
3. Immediately quarantine and destroy all devices within your facility's control per your facility's procedures.
4. Share and post this Advisory letter within your facility network and forward to any customers you may have distributed the product to ensure awareness. Ensure the contents of this Product Advisory are read and understood by those within your organization.
5. ~~Complete the attached Distributor Customer Response Form and~~ **return to your distributor, whether or not you have any of the impacted product**, to document your receipt of this notification and receive credit for the affected product from your distributor. ~~(Attachment 1 – Distributor Response Form).~~
6. Report any adverse health consequences experienced with the use of this product to BD. Events may also be reported to the FDA's MedWatch Adverse Event Reporting program via:

   Web: MedWatch website at www.fda.gov/medwatch
   Phone: 1-800-FDA-1088 (1-800-332-1088)
   Mail: MedWatch, HF-2, FDA, 5600 Fisher's Lane, Rockville, MD 20852-9787

7. Report any complaints experienced with the use of this product to BD via the North American Regional Complaint Center:
   Phone: 1-844-8BD-LIFE (1-844-823-5433) say "Product Complaints" when prompted
   Mon–Fri 8:00am and 5:00pm CT
   Email: productcomplaints@bd.com

Affected Product:

Please see Attachment 2 for affected product information.

Actions Taken by BD:

BD is investigating the cause of the issue and will implement corrective action based on the investigation results to prevent re-occurrence.

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Contact Information: If you require further assistance please contact:

BD Contact	Contact Information	Areas of Support
BD Technical Support	1-844-8BD-LIFE (1-844-823-5433) Say "Product Complaint" when prompted M-F 8am - 5pm CT or Email: productcomplaints@bd.com	Product Complaints, Technical Support

BD is committed to advancing the world of health. Our primary objectives are patient and user safety and providing you with quality products. We apologize for any inconvenience this issue may have caused and thank you in advance for helping us to resolve this matter as quickly and effectively as possible.

Sincerely,

Haytham Gareer

Haytham Gareer
Senior Vice President, Medical Affairs
BDI Business Units

Elizabeth Gaipa

Elizabeth Gaipa
Vice President, Quality Management
BD UCC

Enclosures:
Attachment 1 – Distributor Customer Response Form
Attachment 2 – Affected Product

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Attachment 2- Affected Product

Product Name	Catalog #	Lot#	Exp Date	Pri UDI (DI+PI) (Product)	Sec UDI (DI+PI) (Case)
Bard® Silastic® Foley Catheter	33416	NGKX0444	1/31/2029	00801741019982	10801741019989
	33418	NGKW2813	1/31/2029	00801741019999	10801741019996
	33420	NGKQ1015	1/31/2029	00801741020001	10801741020008
	33422	NGKW3707	1/31/2029	00801741020018	10801741020015
	33424	NGKW2669	1/31/2029	00801741020025	10801741020022
Bard® Silastic® Foley Catheter	33614	NGKN4037	1/31/2029	00801741020056	10801741020053
		NGKP1420	9/30/2029		
		NGKQ0040	9/30/2029		
		NGKQ3901	12/31/2028		
		NGKS4937	9/30/2029		
		NGKT3258	9/30/2029		
		NGKV3462	6/30/2029		
		NGKW3897	6/30/2029		
	33616	NGKQ0042	9/30/2029	00801741020063	10801741020060
		NGKQ3946	1/31/2029		
		NGKR3420	9/30/2029		
		NGKT3281	9/30/2029		
		NGKU1015	11/30/2029		
		NGKV2275	9/30/2029		
		NGKV2309	9/30/2029		
		NGKW3771	9/30/2029		
	33618	NGKQ3370	5/31/2029	00801741020070	10801741020077
		NGKS3753	9/30/2029		
		NGKT3309	9/30/2029		
		NGKU4129	9/30/2029		
		NGKV0617	9/30/2029		
		NGKV1446	7/31/2029		
		NGKV2345	7/31/2029		
		NGKV3478	7/31/2029		
		NGKV5029	7/31/2029		
	33620	NGKS5060	1/31/2029	00801741020087	10801741020084
		NGKT3308	11/30/2029		
		NGKV3689	11/30/2029		
	33622	NGKQ1016	1/31/2029	00801741020094	10801741020091
		NGKS0244	1/31/2029		
		NGKU0967	9/30/2029		
		NGKV5075	1/31/2030		
		NGKX0441	1/31/2030		
	33624	NGKQ1722	9/30/2029	00801741020100	10801741020107
		NGKQ3862	1/31/2029		
		NGKT3259	9/30/2029		
		NGKX0456	1/31/2030		