

Medline Industries, LP. Three Lakes Drive Northfield, IL 60093-2753 Toll Free: 866.359.1704
Email: Recalls@Medline.com

medline.com

Medline Industries, LP IMMEDIATE ACTION REQUIRED RECALL Medline-Labeled Syringes

Fax: 866.767.1290

April 22, 2024

Dear Valued Medline Customer:

This letter is to inform you that Medline Industries, LP issued a recall for specific lots of item number SYR101010 and SYR101020. It has been identified that testing documentation does not support that the syringes can successfully deliver accurate volumes across the full range of their claimed graduated capacity and may pose a risk to patient health. Syringes not being able to deliver the intended dose may cause inaccurate dosing leading to the need for additional treatment or other medication. Affected product shall be destroyed per credit per instructions below. Please destroy the affected product in accordance with your state and local regulations. No product is to be returned to Medline. Adverse reactions or quality problems experienced with the use of this product may be reported to the FDA's MedWatch Adverse Event Reporting program either online, by regular mail or by fax.

REQUIRED ACTION:

- 1. Immediately check your stock for **item number and lot number (All Lot Numbers Starting with 897),** listed on the destruction form, quarantine all affected product. Destroy all product belonging to the affected lot numbers listed on the destruction form.
- 2. Please return the completed enclosed destruction form listing the quantity of affected product destroyed. Even if you do not have any affected product please complete and return the form, as indicated on the destruction form.
 - When we receive your completed destruction form your account will be issued a credit, if applicable.
- 3. If you have transferred possession of this product to another individual, to another department or another location in your organization you are required by FDA regulations to notify them of this recall communication.

If you have any questions, please contact 866-359-1704 or email recalls@medline.com

As the distributor of this product, we sincerely regret the inconvenience. We, like you, place the health and safety of your patients first and foremost.

Sincerely,

Karin Johnson

Karin Johnson
Product Specialist Sr.

Recall Ref#: R-24-075-X23



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URGENT REMEDIAL ACTION DESTRUCTION FORM

April 22, 2024

Recall Ref# R-24-075-X23 1081285 CARELINC MEDICAL EQUIPMENT 89 54TH ST SW GRAND RAPIDS, MI 49548-5503

Medline Industries, LP

Medline-Labeled Syringes

Please follow your lo	cal and state	Ordinances when disposing/destroying	of this or any product.	
Medline Item (used to purch		Lot Number Affected (All lot numbers Starting with 897)	Individual Quantity Destroyed*	
SYR1010	110		EA	
SYR1010	20		EA	
		*Indicate zero "0" if you do not have	ve any lots and/or items to destroy.	
If you have transferred in your o	d possession of th organization you a	nis product to another individual, to another depa are required by FDA regulations to notify them of	ertment or another location this recall communication.	
By signing b	elow you are verifying	that you have received, read, understood and performed any	actions specified in the recall letter.	
Print Name			Title	
Signature	Date	Emai	Email Address	
Telephone Number		Fax	(Number	

Even if you have no affected stock:
Please return the completed form via email to recalls@medline.com or fax 866-767-1290

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