

Subject

URGENT- Field Safety Notice – Distributors Klinion Kliniray Gauze Compress X-Ray Klinion Kliniray Abdominal Gauze Compress X-Ray

Dear Sir/Madam.

Medeco BV initiated a product recall for <u>specific product codes and lot numbers</u> of the Klinion Kliniray Gauze Compress X-Ray and Klinion Kliniray Abdominal Gauze Compress X-Ray. Please note that it does <u>not</u> affect all product codes of the Klinion Kliniray Gauze Compress X-Ray and Klinion Kliniray Abdominal Gauze Compress X-Ray.

Description of the problem:

After assessment of these products, it became known that x-ray threads in gauze compresses can disintegrate. It has been confirmed that these products are not meeting the required criteria.

Theoretically, small pieces of thread could be left behind if the x-ray thread breaks or frays. This may, for example, lead to inflammation and/or granuloma formation when remaining in the body.

The identified product codes and lot numbers within this notice are potentially affected. For this reason and to address any potential risk of harm, all of the affected products should <u>not be used</u>.

Product identification procedure:

The only way to identify affected products is by checking product code and lot number to the recalled product list (see attachment 1).

Attachment 2 gives an example of a packaging label that highlights the location of the product code and lot number on the device labels. The label can be found on the primary packaging and the carton. The product code (reference number) is preceded by the word REF. The lot number is preceded by and is in 4500xxxxxxx format.

Contact

Medeco Customer Service

Email

info-medecobv@mediq.com

Date

2 October 2023

Our reference

FSN2023-01

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Advice on actions to be taken by distributor

Our records show that you have taken delivery of affected products (attachment 1). Please follow the steps below:

- 1. Immediately stop distributing and quarantine all affected products.
- Please forward the 'Field Safety Notice END USERS' to your customers and ask them to return the affected products to you together with a completed form 'Recall Response Form for END USERS'.
- When the completed forms have been returned to you, please complete
 the 'Recall Response Form for DISTRIBUTORS' and contact Medeco BV
 for further instructions on how to dispose affected products, to arrange
 credit and possibilities for replacement.
- 4. Please also return the completed 'Recall Response Form for DISTRIBUTORS' and all 'Recall Response Form for END USERS" via email to Medeco BV.

Please provide a completed response as soon as possible.

Transmission of this Field Safety Notice

This notice should be sent to anyone who has received the affected devices within your organisation and to any organisation where the affected devices have been transferred to.

We are committed to deliver products of high quality to our customers and we apologize for any inconvenience that this notice can cause.

If you have any questions related to this recall, please contact us at number provided above.

The relevant National Authorities have been informed about this Field Safety Corrective Action.





RECALL RESPONSE FORM for DISTRIBUTOR URGENT – FIELD SAFETY NOTICE

Please complete and return by email to info-medecobv@mediq.com

•		•				
Deliveries to yo	our facility:					
Invoice #	Product Code (REF number)	luct Medeco e Article Code (SAP		del	Quantity delivered (pieces)	
If yes If you	ou distributed : Have you no did not notify	I the product otified down to	further ? o your cus			
2. Do you	Yes / NO					
Record quantit Product Code REF number	Medec	each LOT to	·		Quantity (pieces)	
Please provide	e details of af	fected produ	cts that we	re distril	buted to your	
Customer/ Company nar		Medec Article Code (T No.	Quantity (pieces)	
. ,	(REF number	,	J. 1.			
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DISTRIBUTOR FORM Completed and returned from:

Name:	
Function:	
Company	
name:	
Address:	
Email address:	
Phone number:	
Signature:	
Date:	
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As requested, I have also provided a copy of all "Recall Response Forms for End Users" returned from customers who received affected product.



Appendix 1
Products including LOT numbers affected:

Medeco	Product	LOT number	Name and description
Article	REF		
Number	number		
3016720	111180	4500131696	KLINION GAASK 10X10 12L XRAY 50ST
3016720	111180	4500142292	KLINION GAASK 10X10 12L XRAY 50ST
3016720	111180	4500142298	KLINION GAASK 10X10 12L XRAY 50ST
3016720	111180	4500159100	KLINION GAASK 10X10 12L XRAY 50ST
3016722	111209	4500131696	KLINION GAASK 10X20 12L XRAY 50ST
3016722	111209	4500134619	KLINION GAASK 10X20 12L XRAY 50ST
3016722	111209	4500139418	KLINION GAASK 10X20 12L XRAY 50ST
3016722	111209	4500142298	KLINION GAASK 10X20 12L XRAY 50ST
3016722	111209	4500150886	KLINION GAASK 10X20 12L XRAY 50ST
3016722	111209	4500154489	KLINION GAASK 10X20 12L XRAY 50ST
3016722	111209	4500159105	KLINION GAASK 10X20 12L XRAY 50ST
3016736	115001	4500142292A	KLINION GAASK 5X 5 12L XRAY *S* PIP10
3016736	115001	4500142292B	KLINION GAASK 5X 5 12L XRAY *S* PIP10
3016737	115005	4500131696	KLINION GAASK 10X10 12L XRAY *S* PIP5
3016737	115005	4500136318	KLINION GAASK 10X10 12L XRAY *S* PIP5
3016737	115005	4500139418	KLINION GAASK 10X10 12L XRAY *S* PIP5
3016737	115005	4500142292A	KLINION GAASK 10X10 12L XRAY *S* PIP5
3016737	115005	4500142292B	KLINION GAASK 10X10 12L XRAY *S* PIP5
3016739	115007	4500131696	KLINION GAASK 10X10 12L XRAY *S* PIP10
3016739	115007	4500134619	KLINION GAASK 10X10 12L XRAY *S* PIP10
3016739	115007	4500139418A	KLINION GAASK 10X10 12L XRAY *S* PIP10
3016739	115007	4500139418B	KLINION GAASK 10X10 12L XRAY *S* PIP10
3016739	115007	4500142301	KLINION GAASK 10X10 12L XRAY *S* PIP10
3016739	115007	4500142293	KLINION GAASK 10X10 12L XRAY *S* PIP10
3016739	115007	4500142299	KLINION GAASK 10X10 12L XRAY *S* PIP10
3016739	115007	4500150886	KLINION GAASK 10X10 12L XRAY *S* PIP10
3016739	115007	4500159105	KLINION GAASK 10X10 12L XRAY *S* PIP10
3016739	115007	4500163488	KLINION GAASK 10X10 12L XRAY *S* PIP10
3016740	115020	4500131696	KLINION GAASK 10X20 12L XRAY *S* PIP5
3016740	115020	4500148507B	KLINION GAASK 10X20 12L XRAY *S* PIP5
3016740	115020	4500154489	KLINION GAASK 10X20 12L XRAY *S* PIP5
3016741	115021	4500123945	KLINION GAASK 10X20 12L XRAY *S* PIP10
3016741	115021	4500131696	KLINION GAASK 10X20 12L XRAY *S* PIP10
3016741	115021	4500134619A	KLINION GAASK 10X20 12L XRAY *S* PIP10
3016741	115021	4500134619B	KLINION GAASK 10X20 12L XRAY *S* PIP10
3016741	115021	4500139418	KLINION GAASK 10X20 12L XRAY *S* PIP10
3016741	115021	4500142293	KLINION GAASK 10X20 12L XRAY *S* PIP10
3016741	115021	4500142298	KLINION GAASK 10X20 12L XRAY *S* PIP10
3016741	115021	4500148507	KLINION GAASK 10X20 12L XRAY *S* PIP10
3016741	115021	4500150886	KLINION GAASK 10X20 12L XRAY *S* PIP10
3016741	115021	4500154490	KLINION GAASK 10X20 12L XRAY *S* PIP10
3016741	115021	4500157250	KLINION GAASK 10X20 12L XRAY *S* PIP10
3016741	115021	4500157230	KLINION GAASK 10X20 12L XRAY *S* PIP10
3016741	115021	4500159105	KLINION GAASK 10X20 12L XRAY *S* PIP10
3010741	110021	7300133103	INCIDION OFFICIAL TOTAL ANAL S FIFTU



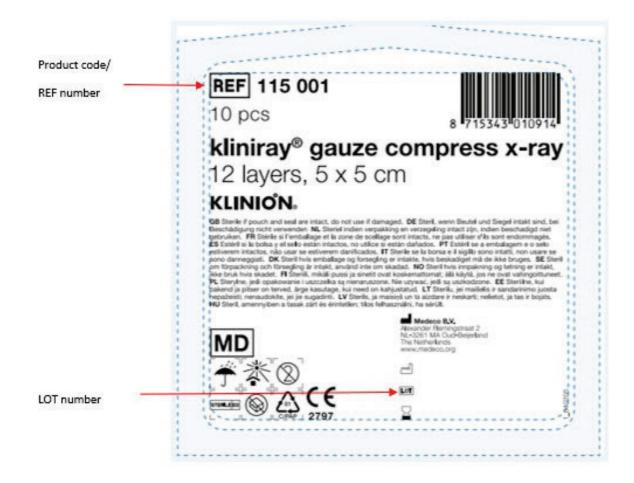
A MEDIQ COMPANY

Medeco	Product	LOT number	Name and description
Article	REF		
Number	number		
3016742	115023	4500139418A	KLINION GAASK 10X20 24L XRAY *S* PIP5
3016742	115023	4500139418B	KLINION GAASK 10X20 24L XRAY *S* PIP5
3016742	115023	4500142299	KLINION GAASK 10X20 24L XRAY *S* PIP5
3016742	115023	4500150887	KLINION GAASK 10X20 24L XRAY *S* PIP5
3016742	115023	4500157250	KLINION GAASK 10X20 24L XRAY *S* PIP5
3016744	115046	4500134619	KLINION GAASK 20X25 16L XRAY *S* PIP5
3016744	115046	4500142299	KLINION GAASK 20X25 16L XRAY *S* PIP5
3016746	115048	4500163297	KLINION GAASK 20X25 32L XRAY *S* PIP2
3016749	115062	4500139419A	KLINION BUIKGAAS 45X45 4L XRAY *S*



Appendix 2 Example of Packaging Labelling

Primary packaging





Outer carton label

