



Urgent Safety Notice

Recall
concerning

Priflex®, Spiral-reinforced PRIFLEX cannula, with low pressure cuff, size 8

13.05.2026

Sender:

Primed Halberstadt Medizintechnik GmbH
Straße des 20. Juli 1
38820 Halberstadt

Target group:

Users, Distributors, Hospital Logistics, Medical Device Safety Officers

Identification of the affected medical devices:

Item-Nr.	Description
200815	Priflex®, Spiral-reinforced PRIFLEX cannula, with low pressure cuff, size 8

Potentially affected batches: LOT **02.09.2025**

Description of the problem, including the identified root cause:

We have received information regarding an incident involving one of our tracheal cannulas. In this instance, the inflation channel of the cuff system inverted inward into the inner lumen. This results in a narrowing of the inner lumen of the tracheal cannula. This may lead to respiratory distress for the patient. Depending on the patient's medical condition, medical intervention may be required.

The described defect may be observed during the cuff integrity check performed prior to inserting the tracheal cannula.

The defect may also occur during the use of the tracheal cannula.

For patients currently being treated with the affected product, we recommend the immediate replacement of the tracheal cannula.

Should immediate replacement not be possible due to the current treatment situation, the tracheal cannula must be closely monitored for signs of this defect and replaced immediately if the defect occurs.



What actions should the recipient take?

Please perform the following actions:

1. Identify the products in your facility/company
2. Completely remove the product inventory from any use
3. Separate the products and make them available for collection
4. Confirmation to Primed Halberstadt Medizintechnik GmbH that the goods are ready for collection

We ask you to include the safety notice regarding the potentially affected tracheal cannulas in your documentation. We also ask you to forward the safety notice to other hospitals and departments that have received the affected tracheal cannulas.

Please complete the enclosed acknowledgement form in full and return it. Keep yourself informed of this notification and the associated measures.

Sincerely,

Contact:

Steffen Schaefer

(verantwortliche Person nach Artikel 15 MDR 2017/745 /
responsible person according to Article 15 MDR 2017/745)

Tel.: +49 3941- 668 6

vigilanz@primed-halberstadt.de | www.primed-halberstadt.de

Signature



Subject: Safety Notice Receipt Confirmation

Dear Sir or Madam,

Thank you in advance for your cooperation. Please complete this document and return it to your local Primed Halberstadt Medizintechnik GmbH representative or via one of the following methods:

Fax: +49 (0) 3941 - 245 65

E-Mail: vigilanz@primed-halberstadt.de

Mail:

Primed Halberstadt Medizintechnik GmbH
Straße des 20. Juli 1
38820 Halberstadt
Germany

Confirmation of receipt:

I confirm receipt of the safety information issued by Primed Halberstadt Medizintechnik GmbH.

I understand the information and have passed it on to all responsible employees, departments, and facilities affected by this action.

After implementing the above-mentioned measures, we have reached the following conclusion:

Please check/complete the relevant box.

- We have the full quantity in stock. No copies have been transferred to third parties.
- We have the following number of affected products in stock: _____ pcs
- The specified product has already been distributed to third parties in quantities of _____ pcs. We will notify the customer base that received the above-mentioned product(s) of the recall.
- We do not have any of the affected products in stock and will notify our customers who have received the above-mentioned product(s) of the recall.

Name/Name, Vorname/Surname

Unternehmen/Company

Abteilung (Department)/Funktion (Position)

Unterschrift/Signature

Datum/Date