

B. Braun Melsungen AG
Division Hospital Care
Safety Officer Medical Devices

34209 Melsungen

Our Reference FSCA-2023-01-06
Contact: Dr. Stephan Krause
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Date: Jan 06, 2023

URGENT Field Safety Corrective Action – Infusomat Space Transfusion Line - leakage

Dear Sir or Madam,

The B. Braun Melsungen AG has decided to proactively recall defined article batch combinations of Infusomat Space Transfusion Lines in the course of a Field Safety Corrective Action from the market:

Article Number	Article Name	Batch
8270066SP-01	INF.SP.LINE,TRANS,PVC,LL,250CM-EU	22F19E8ST5
8270066SP-01	INF.SP.LINE,TRANS,PVC,LL,250CM-EU	22G01E8ST5
8270066SP-01	INF.SP.LINE,TRANS,PVC,LL,250CM-EU	22G03E8ST5
8270066SP-01	INF.SP.LINE,TRANS,PVC,LL,250CM-EU	22H25E8ST5

Reason for the Recall

In the course of our Post Market Surveillance activities, we identified the risk for leakages on the above mentioned article batch combinations.

Chairwoman of the Supervisory Board:
Dr. Annette Belier

Executive Board:
Markus Strotmann
(Chairman)
Priv.-Doz. Dr. Stefan Ruppert
Jürgen Stihl

Corporate Office: Melsungen
Register Court:
Local Court Fritzlar
HRB 11 000
WEEE-Reg.-No. DE 42690900

Address:
B. Braun Melsungen AG
Carl-Braun-Straße 1
34212 Melsungen
Germany

The potential leakage is located between the tube and the patients' Luer connector as indicated on the picture below:



Whilst no serious injuries to patients, users, or third parties have been reported to date, the deviation might harbour the risk for the patient of microbial contamination, under supply, open patient access, or air infusion.

In view of the identified risks, we decided to recall the above listed devices from the market.

Based on internal controls and available post market data, the effect can be limited to the above mentioned article batches combinations.

Actions to be taken

Our records have shown that your institution has received one or more of the affected article batch combinations.

We kindly ask you to initiate the following activities immediately and with priority:

- Review this Field Safety Notice in its entirety and ensure that all users of the above mentioned product in your organization and other concerned persons are informed about this Field Safety Notice.
- If you are a distributor, please forward this correction notification to your customers.
- Identify, quarantine and return affected articles.
- Do not use affected devices anymore.
- It is recommended to exchange devices from the above mentioned batches, which are currently in use.
- Confirm receipt of this information by completing the attached confirmation slip and return this to B. Braun using the contact details provided.

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If more information is needed or you require devices for replacement, please contact

Local contact 1

Name

Title

Email

telephone

Local contact 2

We believe in improving people's health through everything we do. Patient and user safety is our highest priority. Kindly accept our apologies for any inconveniences caused and thank you in advance for your cooperation to resolve this matter quickly.

Yours sincerely,