

Customer  
Hospital  
City  
Postal code  
Country  
*Attn.: XXX*

## **URGENT Field Safety Notice**

### ***safePICO A aspirating blood syringe***

Dear Customer

#### **Background**

Radiometer has become aware of an issue with the *safePICO A* aspirating blood syringe. The issue relates to the sterile barrier system for the product.

For this reason, Radiometer kindly requests you to stop using the affected product with immediate effect.

#### **Risk for the patient**

The described error may potentially result in bacterial bloodstream infection. The bloodstream infection may be asymptomatic but may also progress to sepsis or life-threatening septic shock. Immunosuppressed patients are at particular risk.

#### **Affected product**

All Lots of the *safePICO A* aspirating blood syringe, 957-204.

For EU Countries only the following is to be included in translated letter:

***EU Basic UDI-DI: 57006900067N9***

***(UDI = Unique Device Identifier – DI = Device Identifier)***

**Your actions**

Radiometer kindly request you to stop using the affected product with immediate effect.

To ensure patient safety in your facility, kindly follow the steps below:

- Check your inventory of the above *safePICO A* aspirating blood syringes.
- Check for the above *safePICO A* aspirating blood syringes distributed in your institution.
- Collect any of the above *safePICO A* aspirating blood syringes and put them in quarantine.
- Complete the Recall Response Form (last page of this letter) and return it to your Radiometer representative together with the quarantined *safePICO A* aspirating blood syringes within two weeks of receiving this letter.

To ensure you receive credit for the *safePICO A* aspirating blood syringes:

- Fill out a credit claim and send it to your Radiometer representative.

**Solution provided by Radiometer**

Currently, it is difficult to provide a specific resolution date for the *safePICO A* aspirating blood syringe.

Please contact your Radiometer representative for information about alternative samplers.

**Your help is appreciated**

If you are not the end-user of the affected product, please ensure that this letter is distributed to the final end-user.

If you have any questions, please contact your Radiometer representative.

Radiometer sincerely apologizes for the inconvenience this situation may cause you.

Best regards,

<State Radiometer distributor name>

# Recall Response Form

Concerning:

## **safePICO A aspirating blood syringe**

- I have received the customer advisory letter and reviewed both my current inventory of *safePICO A* aspirating blood syringes and those distributed in my institute. All affected samplers have been collected and I have returned the quantities of samplers stated below to my Radiometer representative.

Returned quantities:

<b>Item number</b>	<b>Description</b>	<b>Lot</b>	<b>Quantity</b> (boxes of 100 pcs.)
957-204	<i>safePICO A</i> aspirating blood syringe		

In case you have more than one Lot of any type of *safePICO A* in your institute, then please insert extra row(s) in the table and then state Lot numbers and quantities.

- I have no *safePICO A* aspirating blood syringes in my institute.

Hospital Name:	
Your Name:	
Date:	
Signature:	
Email Address:	