

### B. Braun Singapore Pte Ltd

Company Reg No.: 198703127H GST Reg No.: M2-0078143-X

TO WHOM IT MAY CONCERN

CC Chairman Medical Board and relevant Head of Departments

Date: 11 March 2019

# **Urgent FIELD SAFETY NOTICE - Certofix® Quattro Recall**

To whom it may concern,

we, the B. Braun Melsungen AG have decided to recall the CERTOFIX® QUATTRO PRODUCT RANGE in the context of a FIELD SAFETY CORRECTIVE ACTION from the market.

The Certofix® Quattro catheters are distributed in various sterile procedural trays. As the defect cannot be restricted, the whole Certofix® Quattro Product range has to be recalled. Depending on the shelf-life of the products and the distribution on the market, the below mentioned batches have to be considered as initial batches. From then onwards all consecutive batches are affected.

Article No.	Article Name	Initial Batch	Expiry Date
4167767	CERTOFIX QUATTRO V 815	16AXXXXXXX or following	January 2021 or later
4167775	CERTOFIX QUATTRO V 820		
4167783	CERTOFIX QUATTRO V 830		
4167767-07	CERTOFIX QUATTRO V 815-EU/SA		
4167775-07	CERTOFIX QUATTRO V 820-EU/SA		
4167783-07	CERTOFIX QUATTRO V 830-EU/SA		
4167775S	CERTOFIX SAFETY QUATTRO S 820		
4167775S-07	CERTOFIX SAFETY QUATTRO S 820-EU/SA		

71077731 07 02111011111111111111111111111111111	2019 or later
4167767P-S1         CERTOFIX PROTECT QUATTRO V815 - SET1           4167775P         CERTOFIX PROTECT QUATTRO V 820         4CXXXXXX /           4167775P-07         CERTOFIX PROTECT QUATTRO V 820-EU/SA         14CXXXXXXX           March 2	?019 or later
4167775P CERTOFIX PROTECT QUATTRO V 820 4167775P-07 CERTOFIX PROTECT QUATTRO V 820-EU/SA 4CXXXXXX / 14CXXXXXXX March 2	2019 or later
4167775P-07 CERTOFIX PROTECT QUATTRO V 820-EU/SA 14CXXXXXXX March 2	2019 or later
	2019 or later
44.0777FD C4 CERTOLIV PROTECT QUATTRO VISO SET1 or following	
4167775P-S1   CERTOFIX PROTECT QUATTRO V820 - SET1   or following	
4167783P CERTOFIX PROTECT QUATTRO V 830	
4167783P-07 CERTOFIX PROTECT QUATTRO V 830-EU/SA	
4167783P-S1 CERTOFIX PROTECT QUATTRO V830 - SET1	
4161094 PROSET CERTOFIX PROTECT QUATTRO V820	March 19 or later
4161098 PROSET CERTOFIX PROTECT QUATTRO V815	
4161886 PROSET CERTOFIX QUATTRO V830	
4163492 PROSET CERTOFIX QUATTRO S820	
4168178 PROSET CERTOFIX QUATTRO S820	
4168894 PROSET CERTOFIX QUATTRO S830 17CXXXXXXX March	
4167230SM PROSET CERTOFIX SAFETY QUATTRO S820 or following	
4167280SM PROSET CERTOFIX SAFETY QUATTRO S830	
4167312SM PROSET CERTOFIX SAFETY QUATTRO S830	
4167784S PROSET CERTOFIX SAFETY QUATTRO S820	
4168030S PROSET CERTOFIX SAFETY QUATTRO S820	
4161886P PROSET CERTOFIX PROTECT QUATTRO V830	

# Reason for the Recall

Certofix® Quattro is a four-lumen catheter for catheterization of the superior vena cava using the Seldinger technique. Central venous catheters are well known and routinely used for central vein puncture and establishing access to the blood system of a patient to deliver infusion solutions, blood or pharmaceuticals in carrier solutions or monitoring systemic pressure.

In order to avoid accumulation of blood or fluid in the dead space between side hole and catheter tip, the lumen is closed with a plug.

In the course of our internal quality and post market surveillance activities we have found that this plug may not stay in its intended position. This deviation is caused by supplier related component quality issues.

While no serious injuries to patients, users, or third parties have been reported to date, there is a remaining risk of an undersupply, infusion of the plug into the patient and thrombus formation up to embolism.

All other Certofix® catheter variants (Mono, Duo, Trio, Quinto) are <u>not</u> affected as the lumen closure has a different design and is completely produced in-house.



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#### Actions to be taken by the customer:

Our records have shown that your institution has received the affected Certofix® Quattro Catheter as specified above.

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 products in your organization and other concerned persons are informed about this Field Safety Corrective Action. If you are a distributor, please forward this correction notification to your customers.
- Identify, quarantine and return affected goods.
- Do not use affected devices anymore.
- Already placed catheters can stay in situ until therapy is finished depending on individual risk benefit assessment. Check channels for patency.
- Confirm receipt of this information.

# If more information is needed, please contact

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Kindly accept our apologies for any inconveniences.

Yours sincerely,

Teng Hui Min

Senior Regulatory Affairs Executive

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