

FSN & FSCA Ref: 2019FA0009

Date: 22 Oct 2019

## <u>Urgent Field Safety Notice</u> Product Removal - Torcon NB<sup>®</sup> Advantage Catheter

For Attention of: Chief Executive / Risk Management / Purchasing

#### Contact details of local representative (name, e-mail, telephone, address etc.)

Cook Medical Europe Ltd. O'Halloran Road National Technology Park Limerick, Ireland

E-mail: European.FieldAction@CookMedical.com

Phone: Please refer to the attached Country Contacts List

For any further information or support concerning the information within this FSN, please contact your local Cook Medical Sales Representative or Cook Medical Europe Ltd.



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# <u>Urgent Field Safety Notice</u> Product Removal - Torcon NB® Advantage Catheter Risk Addressed by FSN

	Information on Affected Devices				
	1. Device Type(s)				
1.	The Torcon NB® Advantage Catheters are sterile, single-use products. The catheters are available in a variety of French sizes, endhole sizes, lengths, materials and designs (e.g., polyethylene or nylon, non-braided or braided with 1:1 torque).				
	2. Commercial name(s)				
1.	Torcon NB® Advantage Catheter				
	3. Primary clinical purpose of devi	ce(s)			
1.	Per the Instructions for Use T_NBADV_REV0: Intended for use in the peripheral and coronary vascular system, including the carotid arteries, in angiographic procedures by physicians trained and experienced in angiographic techniques.  Per the Instructions for Use T_CE_ANGIO88_REV5: Intended for use in angiographic procedures by physicians trained and experienced in angiographic techniques.				
	4. Device model / Catalogue / Part number(s)	5. Affected serial or lot number range			
	HNB5.0-38-100-P-NS-DAV (G08895)	9708139			
	HNB5.0-38-100-P-NS-H1 (G06005)	9714540			
	HNB5.0-38-100-P-NS-JB1 (G05997)	9726996			
	HNB5.0-38-100-P-NS-JL4 (G05923)	NS9701376			
	HNB5.0-38-100-P-NS-SIM1 (G04922)	9700385, 9713001			
	HNB5.0-38-100-P-NS-SIM2 (G05976)	9713002			
1.	HNB5.0-38-100-P-NS-VERT (G06074)	9710402			
	HNB5.0-38-110-P-12S-PIG (G10741)	9709328			
	HNB5.0-35-65-P-NS-TIPS (G19706)	9890919			
	HNB5.0-38-65-P-2S-C1 (G06827)	NS9906241			
	HNB5.0-38-65-P-NS-SHK1.0 (G05975)	9900974			
	HNB5.0-35-100-P-8S-NEFF (G11026)	NS9720276			
	HNB6.0-38-65-P-NS-BMC (G32465)	9908469, 9911359, 9936124			
	HNB7.0-38-100-P-NS-MPA (G08591)	9713048			



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	Reason for Field Safety Corrective Action (FSCA)			
	Description of the product problem			
2.	Pouches may be undersealed, potentially compromising the sterility of the product. This issue is related specifically to the chevron seal.			
	2. Hazard giving rise to the FSCA			
2.	Use of an affected product could pose significant risk to the patient, as local or systemic infection may occur.			

	Type of Action to Mitigate the Risk						
	1. Action To Be Taken by the User						
3.	☑ Identify Devices ☑ Quarantine Devices ☑ Return Devices						
		Please complete the enclosed Reply Form. Where product is indicated as being returned, our Customer Services department will contact you to organize the return and issue you with the relevant Returns Authorization number. Please include contact details on the Reply Form.					
		Returned Product should be addressed to: Cook Medical EUDC Robert-Koch-Straße, 2 52499 Baesweiler GERMANY					
		Credit will be provided for the returned affected products where applicable.					
3.	2. Is Customer Reply Required? Form is attached specifying deadline for return.  Yes		Yes				
3.	3.	Action Being Taken by the Manufacturer					

General Information			
4.	1. FSN Type	New	
4.	Further advice or information already expected in follow-up FSN?	No	



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General Information Continued			
	Manufacturer information     For contact details of local representative refer to page 1 of this FSN		
4.	a. Company Name	Cook Incorporated	
	b. Address	750 Daniels Way Bloomington, IN 47402, United States	
4.	The Competent (Regulatory) Authority of your country has been informed about this communication to customers.		
4.	5. Name/Signature	Cook Incorporated	

### **Transmission of this Field Safety Notice**

This notice needs to be passed on all those who need to be aware within your organisation or to any organisation where the potentially affected devices have been transferred.

Please transfer this notice to other organisations on which this action has an impact.

Please maintain awareness on this notice and resulting action for an appropriate period to ensure effectiveness of the corrective action.

Please report all device-related incidents to the manufacturer, distributor or local representative, and the national Competent Authority if appropriate, as this provides important feedback.



### **Field Action Customer Reply Form**

1. Field Safety Notice (FSN) information		
FSN Reference number	2019FA0009	
FSN Date	22 Oct 2019	
Product/Device name	Torcon NB® Advantage Catheter	
Product Part Number(s)	HNB5.0-38-100-P-NS-DAV (G08895) HNB5.0-38-100-P-NS-H1 (G06005) HNB5.0-38-100-P-NS-JB1 (G05997) HNB5.0-38-100-P-NS-JL4 (G05923) HNB5.0-38-100-P-NS-SIM1 (G04922) HNB5.0-38-100-P-NS-SIM2 (G05976) HNB5.0-38-100-P-NS-VERT (G06074) HNB5.0-38-110-P-12S-PIG (G10741) HNB5.0-35-65-P-NS-TIPS (G19706) HNB5.0-38-65-P-S-C1 (G06827) HNB5.0-38-65-P-NS-SHK1.0 (G05975) HNB5.0-35-100-P-8S-NEFF (G11026) HNB6.0-38-65-P-NS-BMC (G32465) HNB7.0-38-100-P-NS-MPA (G08591)	
Batch/Serial Number(s)	9700385, 9708139, 9709328, 9710402, 9713001, 9713002, 9713048, 9714540, 9726996, 9890919, 9900974, 9908469, 9911359, 9936124, NS9701376, NS9720276, NS9906241	

2. Customer Details		
Account Number		
Healthcare Organisation Name		
Organisation Address		
Contact Name		
Title or Function		
Telephone number		
Email		



3. Customer action undertaken on behalf of Healthcare Organisation Please mark boxes below to indicate actions have been completed. If action is not applicable, please write N/A in the column on the right.					
	I confirm receipt of the Field Safety Notice and that I read and understood its content.			ood	
	The information and required actions have been brought to the attention of all relevant users and executed.			n of	
	I have affected devices to return - enter Lot number and quantities in table below.			ıble	
	No affected devices remain in our organisation's inventory				
Print	Print Name				
Signature					
Date	Date				
4. F	Return acl	knowledgement to ser	nder		
Email			European.FieldAction@CookMedical.com		
Fax			+ 353 61 239294		
Deadline for returning the customer reply form		turning the customer	Please return this form within 5 business days of receipt, even if you do not have any of the affected product(s).		
Cust	omer Help	line	Please refer to the attached Country Contacts List		
If you are returning any affected product, please indicate the part number, lot number and quantity					
Product Part Number		Number	Product Lot Number	Quantity	

It is important that your organisation takes the actions detailed in the FSN and confirms that you have received the FSN.

Your organisation's reply is the evidence we need to monitor the progress of the corrective actions.