

Urgent Field Safety Notice

Drill Sleeve Guide
FSCA-identifier 22-08-2019-00001
Type of action: Field Safety Notice (FSN)

Date: September 10, 2019

JAMIE ATKINS SEVERN HEALTHCARE 42 KINGFISHER COURT HAMBRIDGE ROAD NEWBURY, BERKSHIRE, RG145SJ, UK

Attention Valued Customer:

Details on affected devices:

	Produ	ıct and Distribu	tion Table		
Product Name	Manufacturer's Catalog Number	Lot Number	Distribution Date	Expiration Date	Quantity
Drill Sleeve Guide	DSG-90-2.3N	417250	7/21/2014	2016-05	2
Drill Sleeve Guide	DSG-90-2.3N	417280	10/31/2014	2016-08	1
Drill Sleeve Guide	DSG-90-2.3N	511130	10/8/2015	2017-03	1
Drill Sleeve Guide	DSG-90-2.3N	511130	11/9/2015	2017-03	1
Drill Sleeve Guide	DSG-6.3-090-2.4N	515211	2/26/2016	2017-11	1
Drill Sleeve Guide	DSG-6.3-090-2.4N	618010	8/4/2016	2018-01	2
Drill Sleeve Guide	DSG-6.3-090-2.4N	812050	5/18/2018	2020-05-01	4

Description of the problem:

On August 22, 2019, Ad-Tech Medical Instrument Corporation decided to voluntarily recall all drill sleeve guides which are intended to be used only with the 2.4mm diameter Cranial Drill Bit. This recall has been initiated due to an investigation being performed that has identified potential issues with both of our raw material suppliers. The inner diameter of the drill sleeve guide raw material was found to be under tolerance, potentially resulting in the drill bit seizing in the guide during surgery. Although these devices do not make patient contact and are not expected to result in death, this issue could result in a delay in procedure.

The worst-case severity has been determined to be Moderate. This deficiency is not expected to result in death, but there is a possibility an additional surgery could be required if a drill bit seizes within a guide and replacements are not available on-hand. A delayed or additional surgery constitutes additional medical intervention. There is no further impact to the patient, as once the drill bit becomes stuck within the guide, neither the drill sleeve guide nor the drill bit can further interact with the patient.

Actions to be taken by the Customer:

- Immediately examine your inventory and quarantine any product subject to recall. Since the devices are reusable, if you have any of this product in stock with an unknown lot number, please return those as well. In addition, if you may have further distributed this product, please identify your customers and notify them at once of this product recall. Your notification to your customers may be enhanced by including a copy of this recall notification letter. Please return product subject to recall to:
 - Ad-Tech Medical Instrument Corporation 400 West Oakview Parkway
 Oak Creek, WI 53154
- Contact an Ad-Tech Customer Support Specialist for a Return Material Authorization (RMA) number
- Replacements will not be provided
- Credit will be issued for all non-expired returned product
- Please send acknowledgement of this letter as soon as possible to Ad-Tech's Regulatory Department:

FAX: 262-634-5668Telephone: 262-634-1555

o Email: Regulatory@adtechmedical.com

EU contact information:

o E C Rep Ltd

Telephone: (44) 1704 544 944 FAX: (44) 1704 544 050

Email: Janet.Borgerson@ecrep.ie

The undersigned confirms that this notice has been notified to the appropriate Regulatory Agency

Authorized by:

Kathleen Barlow

Signature:

Little:

Regulatory Team Representative and CAPA/Complaints Manager



MEDICAL DEVICE RECALL RETURN RESPONSE Acknowledgement and Receipt Form

Response is Required

JAMIE ATKINS SEVERN HEALTHCARE 42 KINGFISHER COURT HAMBRIDGE ROAD NEWBURY, BERKSHIRE, RG145SJ, UK

DRILL SLEEVE GUIDES

I have read and understand the recall instructions provided in the September 10, 2019 letter. Yes No
Any adverse events associated with recalled product? Yes No
If yes, please explain:
x x
Affected Product Information:

Affected Product Information Table				
Product Name	Catalog Number	Lot Number	Quantity In Inventory	Quantity Returned
Drill Sleeve Guide	DSG-90-2.3N	417250		
Drill Sleeve Guide	DSG-90-2.3N	417280		
Drill Sleeve Guide	DSG-90-2.3N	511130		
Drill Sleeve Guide	DSG-90-2.3N	511130		

Drill Sleeve Guide	DSG-6.3-090-2.4N	515211	
Drill Sleeve Guide	DSG-6.3-090-2.4N	618010	
Drill Sleeve Guide	DSG-6.3-090-2.4N	812050	

Return	Respo	onse	Box:
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Return Response Box:			
Please provide any ad	dditional information, if applicable.		
Distributors:			
I have checked my	y stock and have quarantined inventory consisting of units.		
by (specify date a	nd notified my customers that were shipped or may have been shipped this product nd method of notification); <or> of customers who received/may have received this product. Please notify my</or>		
Questions: (when	applicable)		
Please have so	meone from the Regulatory Department contact me		
Signature of Recei	ipt		
Name/Title			
Telephone			
Email Address			

PLEASE FAX COMPLETED RESPONSE FORM TO: 262-634-5668, ATTENTION: REGULATORY DEPARTMENT OR MAIL TO: AD-TECH MEDICAL INSTRUMENT CORPORATION, 400 WEST OAKVIEW PARKWAY, OAK CREEK, WI 53154

OR EMAIL TO: Regulatory@adtechmedical.com