

June 4, 2019

Dear Healthcare Professional,
 cc: Chairman Medical Board and relevant Head of Departments

Subject: **URGENT MEDICAL DEVICE RECALL**

Affected Product: Ultra-Drive® Irrigation Tubing Assembly

Item Number	Description	Lot Numbers					
423834	Ultra-Drive® Irrigation Tubing Assembly	217761	215048	212966	210002	211237	211325
		216489	215524	212622	210867	210848	209586
		219913	212928	215445	215329	210566	214051
		220935	214669	218667	215623	210972	210488
		219661	211172	217767	217556	214499	214995
		217370	212076	214005	214546	214255	214822
		216611	212488	214118	219305		



Zimmer Biomet is conducting a medical device recall for the Ultra-Drive® Irrigation Tubing Assembly due to insufficient data to support the labeled shelf life of 10 years.

Risks		
Describe immediate health consequences (injuries or illness) that may result from use of or exposure to the product issue.	Most Probable	Highest Severity
	None	<i>Extension of Surgery < 30 minutes while another tubing set is obtained or alternative method is used</i>
Describe long range health consequences (injuries or illness) that may result from use of or exposure to the product issue.	Most Probable	Highest Severity
	None	<i>Infection, surgical interventions (Biologic Response)</i>

Our records indicate that you may have received one or more of the affected products. The affected units were distributed between September 2009 and March 2018.

Risk Manager Responsibilities:

1. Review this notification and ensure that affected personnel are aware of the contents.
2. If you have affected product at your facility, assist your Zimmer Biomet sales representative and quarantine all affected product. Your Zimmer Biomet sales representative will remove the affected product from your facility.
3. Complete **Attachment 1 – Certificate of Acknowledgement** and send to CorporateQuality.PostMarket@zimmerbiomet.com. This form must be returned even if you do not have affected products at your facility.
4. Retain a copy of the acknowledgement form with your recall records in the event of a compliance audit of your facility’s documentation.
5. If you have further questions or concerns after reviewing this notice, please call customer service at 574-371-3071 between 8:00 am and 5:00pm EST, Monday through Friday. Calls received outside of call center operating hours will receive a voicemail prompt or be transferred to an on-call representative in the event of an emergency. Alternatively, your questions may be emailed to CorporateQuality.PostMarket@zimmerbiomet.com.

Surgeon Responsibilities:

1. Review this notification for awareness of the contents.
2. There are no specific patient monitoring instructions related to this recall that are recommended beyond your existing follow-up schedule.
3. Complete **Attachment 1 – Certificate of Acknowledgement** and send to CorporateQuality.PostMarket@zimmerbiomet.com.
4. Retain a copy of the acknowledgement form with your recall records in the event of a compliance audit of your facility’s documentation.
5. If you have further questions or concerns after reviewing this notice, please call customer service at 574-371-3071 between 8:00 am and 5:00pm EST, Monday through Friday. Calls received outside of call center operating hours will receive a voicemail prompt or be transferred to an on-call representative in the event of an emergency. Alternatively, your questions may be emailed to CorporateQuality.PostMarket@zimmerbiomet.com.



Other Information

This medical device recall was reported to the U.S. Food and Drug Administration and will be reported to other Competent Authorities, Notified Bodies, and Regulatory Authorities as required.

- Med Watch Reporting: Adverse reactions or quality problems experienced with the use of this product may be reported to the FDA's Med Watch Adverse Event Reporting program either online, by mail, or by fax.
- Online: www.fda.gov/medwatch/report.htm
- Mail: Use postage paid, pre-addressed form FDA 3500, available at: www.fda.gov/MedWatch/getforms.htm
- Fax: 1-800-FDA-0178

Under 21 CFR 803, manufacturers are also required to report any serious injuries where a product has contributed or may have contributed to the event. Please keep Zimmer Biomet informed of any adverse events associated with this product or any other Zimmer Biomet product by emailing product.experience@zimmerbiomet.com.

Please be aware that the names of user facilities notified are routinely provided to the Competent Authorities for audit purposes.

The undersigned confirms that this notice has been delivered to the appropriate Regulatory Agencies.

Thank you for your assistance. We regret any inconvenience caused by this recall.

Sincerely,



Kevin W. Escapule
Post Market Surveillance & Regulatory Compliance Director



ATTACHMENT 1
Certificate of Acknowledgement

IMMEDIATE RESPONSE REQUIRED – TIME SENSITIVE ACTION NEEDED

Affected Product: Ultra-Drive® Irrigation Tubing Assembly

Field Action Reference: ZFA 2018-00634

<p>Please check one as applicable:</p> <p><input type="checkbox"/> Hospital Facility <input type="checkbox"/> Surgeon</p> <p>Do you have affected product in your facility? (Hospital Facility Only: Please mark the appropriate response.)</p> <p><input type="checkbox"/> Yes, we currently have one or more affected items in our facility.</p> <p><input type="checkbox"/> No, we currently have no affected items in our facility.</p>
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By signing below, I acknowledge that the required actions have been taken in accordance with this recall notice.

Printed Name: _____ **Signature:** _____

Title: _____ **Telephone:** () _____ - _____ **Date:** ____/____/____

Facility Name: _____

Facility Address: _____

City: _____ **State:** _____ **ZIP:** _____

Note: This form must be returned to Zimmer Biomet before this action is closed for your account. It is important that you complete this form and email a copy to CorporateQuality.PostMarket@zimmerbiomet.com or fax to 574-373-3589.