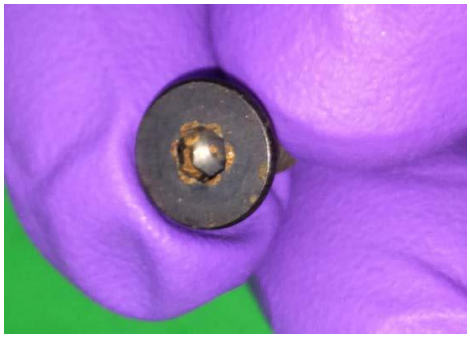
April 30, 2019

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

**Subject:** **URGENT MEDICAL DEVICE RECALL**

**Affected Product:** Oxford Unicompartmental Knee Drill Guides (Phase 3/Domed Lateral)

| Item Number | Lot Number | Item Description | Item Number | Lot Number | Item Description |
|---|---|---|---|---|---|
| 32-420321 | ZB160201 | Phase 3 Femoral Drill Guide Small | 32-420323 | ZB170301 | Phase 3 Femoral Drill Guide Large |
| 32-420321 | ZB160401 | | 32-420323 | ZB170401 | |
| 32-420321 | ZB161101 | | 32-420323 | ZB170901 | |
| 32-420321 | ZB170401 | | 32-420324 | ZB110301 | Phase 3 Femoral Drill Guide Extra Large |
| 32-420321 | ZB170601 | | 32-421060 | ZB160601 | Phase 3 Femoral Drill Guide Extra Small |
| 32-420321 | ZB170801 | | 32-421060 | ZB170201 | |
| 32-420321 | ZB170901 | | 32-421060 | ZB170601 | |
| 32-420322 | ZB130801 | Phase 3 Femoral Drill Guide Medium | 32-421930 | ZB160101 | System Domed Lateral Femoral Drill Guide Small |
| 32-420323 | ZB161102 | Phase 3 Femoral Drill Guide Large | 32-421930 | ZB160801 | |
| 32-420323 | ZB170201 | | 32-421931 | ZB150901 | System Domed Lateral Femoral Drill Guide Medium |



Zimmer Biomet is conducting a lot specific medical device recall for the Oxford Unicompartmental Knee Drill Guides (Phase 3/Domed Lateral) due to the incorrect raw material used by the supplier in the manufacturing of the screw component, which material could potentially lead to corrosion.

| 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 – alternative device obtained and used.* | *Device used and corrosion identified during surgery – extension to surgery > 30 minutes to obtain new device.* |
| 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 – alternative device obtained and used.* | *Adverse tissue reaction/inflammatory response due to foreign debris. Potential revision surgery required as a result.* |

Our records indicate that you may have received one or more of the affected products. The affected units were distributed between March 2016 and August 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.

**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 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,

[redacted]

Kevin W. Escapule
Post Market Surveillance and Regulatory Compliance Director

ZIMMER BIOMET

# ATTACHMENT 1
## Certificate of Acknowledgement

**IMMEDIATE RESPONSE REQUIRED – TIME SENSITIVE ACTION NEEDED**

**Affected Product:** Oxford Unicompartmental Knee Drill Guides (Phase 3/Domed Lateral)

**Field Action Reference:** ZFA 2018-00529

**Do you have affected product in your facility?**
(Hospital Facility Only: Please mark the appropriate response.)

☐ **Yes**, we currently have one or more affected items in our facility.

☐ **No**, we currently have no affected items in our facility.

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.