



**Urgent Medical Device Recall Notice**  
**R-2018-37**

December 13, 2018

<Insert Address>

This letter is to inform you that Smith & Nephew Inc., have voluntarily initiated a recall to remove a single lot of JOURNEY DCF AP Femoral Cutting Blocks Size 3, due to a manufacturing error. The knob on the affected devices was laser etched on the wrong side.

Please see product details below:

Product Number	Description	Batch Number
74012413	JOURNEY DCF AP Femoral Cutting Block Size 3	18BM19943

Shipment Date: 5/1/2018 – 6/8/2018

**Potential Risk of Use of the Product**

In the event the affected device is presented for use, the surgeon would notice that the device is not in the neutral position. The surgeon would make the required adjustments to ensure the appropriate cuts are made. Therefore, the use of or exposure to the affected device presents a low risk of adverse health consequences.

**Required Actions:**

- Please follow the instructions on the attached Response Form.

Enclosure: Response Form

