# California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

Medtronic Navigation, Inc. Medtronic PIN, 100mm and 150mm Sterile Percutaneous

| Recall Date | Product Description | Recalling Firm | Recall Reason |
|---|---|---|---|
| 01/27/2025 | Medtronic PIN, 100mm STERILE PERCUTANEOUS REF: 9733235<br><br>Medtronic PIN, 150MM, STERILE PERCUTANEOUS REF: 97733236<br><br>The Sterile Percutaneous Reference Pin Set is intended to be used to place and remove a pin percutaneously into bony patient anatomy for rigid attachment of a patient reference during image-guided surgeries using a Medtronic computer-assisted surgery system. | Medtronic Navigation, Inc. | Due to an increase in complaint that the percutaneous reference pin would not fit into the patient reference frame or percutaneous pin adapter. |

| Recall Class | Product Identification | Distribution | Affected Dates |
|---|---|---|---|
| II | Sterile Percutaneous Pin, 100mm<br>Product Number: 9733235<br>UDI-DI code: 00613994247872<br>Serial/Lot Numbers:<br>2023071142 - 2024080530 | 67,370 units nationwide | December and prior |

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|---|---|---|---|
| | Sterile Percutaneous Pin, 150mm<br>Product Number: 9733236<br>UDI-DI code: 613994247865<br>Serial/Lot Numbers:<br>2023071143 - 2024080535 | | |

For additional information, please visit the FDA Website.