FDA 510(k) Premarket Notification

# K073531 ACCIN PEDICLE SCREW SYSTEM

**CLEARED**

Feb 4, 2008
49 days to decision

K073531 · Product code: **MNH** · Orthopedic
Source: https://www.510kdatabase.net/k073531/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Orthosis, Spondylolisthesis Spinal Fixation (MNH) |
| Date received | Dec 17, 2007 |
| Decision date | Feb 4, 2008 |
| Days to decision | 49 days |
| Third-party review | No |
| Summary / Statement | Summary |

## APPLICANT

| | |
|---|---|
| Company | **Accelerated Innovation, LLC** |
| Location | Clifton, NJ, US |
| Contact | MICHAEL KVITNITSKY |
| 510(k) history | 7 submissions · 7 cleared · 2008-2008 |