FDA 510(k) Premarket Notification

# K133372 CROSSTREES PVA POD

**CLEARED**

Dec 6, 2013
35 days to decision

K133372 · Product code: **NDN** · Orthopedic
Source: https://www.510kdatabase.net/k133372/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Special |
| Device classification | Cement, Bone, Vertebroplasty (NDN) |
| Date received | Nov 1, 2013 |
| Decision date | Dec 6, 2013 |
| Days to decision | 35 days |
| Third-party review | No |
| Summary / Statement | Summary |

## APPLICANT

| | |
|---|---|
| Company | **Crosstrees Medical, Inc.** |
| Location | Philadelphia, PA, US |
| Contact | JANICE HOGAN |
| 510(k) history | 2 submissions · 2 cleared · 2013-2013 |