FDA 510(k) Premarket Notification

**CLEARED**

# K111983 VISTA-S DEVICE MODEL 08/06-401-XXXXX, 08/06-402-XXXXXX

Nov 18, 2011
129 days to decision

K111983 · Product code: **ODP** · Orthopedic
Source: https://www.510kdatabase.net/k111983/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Intervertebral Fusion Device With Bone Graft, Cervical (ODP) |
| Date received | Jul 12, 2011 |
| Decision date | Nov 18, 2011 |
| Days to decision | 129 days |
| Third-party review | No |
| Summary / Statement | Summary |

## APPLICANT

| | |
|---|---|
| Company | **Zimmer Trabecular Metal Technology** |
| Location | Parsippany, NJ, US |
| Contact | KATHLEEN RUTHERFORD |
| 510(k) history | 11 submissions · 11 cleared · 2007-2014 |