FDA 510(k) Premarket Notification

K103033 TRABECULAR METAL FUSION DEVICE MODEL 06-101-XXXXX, 06-102-XXXXX AND 08-100-XXXXX

CLEARED

Jan 10, 2011
89 days to decision

K103033 · Product code: **ODP** · Orthopedic
Source: https://www.510kdatabase.net/k103033/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Cervical (ODP)
Date received	Oct 13, 2010
Decision date	Jan 10, 2011
Days to decision	89 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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