FDA 510(k) Premarket Notification

K062670 VERTIFLEX SPINAL SCREW SYSTEM

CLEARED

Jan 12, 2007
127 days to decision

K062670 · Product code: **MNH** · Orthopedic
Source: https://www.510kdatabase.net/k062670/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Orthosis, Spondylolisthesis Spinal Fixation (MNH)
Date received	Sep 7, 2006
Decision date	Jan 12, 2007
Days to decision	127 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Vertiflex (Tm), Incorporated**
Location	Carlsbad, CA, US
Contact	STEVE REITZLER
510(k) history	2 submissions · 2 cleared · 2007-2007