FDA 510(k) Premarket Notification

K051697 SELECTIVE-AXIS POSTERIOR NONCERVICAL PLATING SYSTEM

CLEARED

Sep 21, 2005
90 days to decision

K051697 · Product code: **MNH** · Orthopedic
Source: https://www.510kdatabase.net/k051697/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Orthosis, Spondylolisthesis Spinal Fixation (MNH)
Date received	Jun 23, 2005
Decision date	Sep 21, 2005
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Medical Device Advisory Development Group**
Location	Irvine, CA, US
Contact	C. STEPHEN LAWRENCE
510(k) history	1 submissions · 1 cleared · 2005-2005