FDA 510(k) Premarket Notification

K014090 ENDIUS SPINAL FIXATION SYSTEM

CLEARED

Mar 11, 2002
90 days to decision

K014090 · Product code: **KWP** · Orthopedic
Source: https://www.510kdatabase.net/k014090/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Interlaminal (KWP)
Date received	Dec 11, 2001
Decision date	Mar 11, 2002
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Endius, Inc.**
Location	Plainville, MA, US
Contact	GENE DIPOTO
510(k) history	33 submissions · 33 cleared · 1997-2008