FDA 510(k) Premarket Notification

K031290 MODIFICATION TO EASYS TRANSVERSE CONNECTION

CLEARED

May 13, 2003
20 days to decision

K031290 · Product code: **MNH** · Orthopedic
Source: https://www.510kdatabase.net/k031290/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Orthosis, Spondylolisthesis Spinal Fixation (MNH)
Date received	Apr 23, 2003
Decision date	May 13, 2003
Days to decision	20 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Scient'X USA, Inc.**
Location	Round Rock, TX, US
Contact	J.D. WEBB
510(k) history	4 submissions · 4 cleared · 2003-2009