FDA 510(k) Premarket Notification

K061801 2.8 & 3.5MM LACTOSCREW SUTURE ANCHORS

CLEARED

Aug 7, 2006
42 days to decision

K061801 · Product code: **JDR** · Orthopedic
Source: https://www.510kdatabase.net/k061801/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Staple, Fixation, Bone (JDR)
Date received	Jun 26, 2006
Decision date	Aug 7, 2006
Days to decision	42 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Arthrotek, Inc.**
Location	West Hartford, CT, US
Contact	SUSAN ALEXANDER
510(k) history	17 submissions · 17 cleared · 1987-2006