FDA 510(k) Premarket Notification

K061657 HITCH LACTOSORB SUTURE ANCHOR

CLEARED

Jul 25, 2006
42 days to decision

K061657 · Product code: **JDR** · Orthopedic
Source: https://www.510kdatabase.net/k061657/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Staple, Fixation, Bone (JDR)
Date received	Jun 13, 2006
Decision date	Jul 25, 2006
Days to decision	42 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k061657/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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