FDA 510(k) Premarket Notification

K930893 MITEK SUPER QUICKANCHOR, MODIFICATION

CLEARED

Dec 8, 1993
313 days to decision

K930893 · Product code: **JDR** · Orthopedic
Source: https://www.510kdatabase.net/k930893/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Staple, Fixation, Bone (JDR)
Date received	Jan 29, 1993
Decision date	Dec 8, 1993
Days to decision	313 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Mitek Surgical Products, Inc.**
Location	Dedham, MA, US
Contact	ROBERT P ZOLETTI
510(k) history	31 submissions · 26 cleared · 1989-1996