FDA 510(k) Premarket Notification

K924665 QUICKANCHOR

CLEARED

Oct 27, 1992
47 days to decision

K924665 · Product code: **JDR** · Orthopedic
Source: https://www.510kdatabase.net/k924665/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Staple, Fixation, Bone (JDR)
Date received	Sep 10, 1992
Decision date	Oct 27, 1992
Days to decision	47 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