FDA 510(k) Premarket Notification

K032078 MITEK MICRO QUICKANCHOR

CLEARED

Aug 25, 2003
49 days to decision

K032078 · Product code: **NOV** · Orthopedic
Source: https://www.510kdatabase.net/k032078/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Anchor, Suture, Bone Fixation, Metallic (NOV)
Date received	Jul 7, 2003
Decision date	Aug 25, 2003
Days to decision	49 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Mitek Worldwide**
Location	Norwood, MA, US
Contact	SERGIO J GADALETA
510(k) history	10 submissions · 10 cleared · 2002-2004