FDA 510(k) Premarket Notification

K020056 DUET SUTURE ANCHOR

CLEARED

Jun 26, 2002
169 days to decision

K020056 · Product code: **HWC** · Orthopedic
Source: https://www.510kdatabase.net/k020056/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Screw, Fixation, Bone (HWC)
Date received	Jan 8, 2002
Decision date	Jun 26, 2002
Days to decision	169 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Bionx Implants, Ltd.**
Location	Tampere, FI
Contact	TUIJA ANNALA
510(k) history	12 submissions · 12 cleared · 1999-2003