FDA 510(k) Premarket Notification

K052918 REPROCESSED EXTERNAL FIXATION DEVICES

CLEARED

Jan 27, 2006
102 days to decision

K052918 · Product code: **KTT** · Orthopedic
Source: https://www.510kdatabase.net/k052918/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Nail/blade/plate Combination, Multiple Component (KTT)
Date received	Oct 17, 2005
Decision date	Jan 27, 2006
Days to decision	102 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Alliance Medical Corp.**
Location	Phoenix, AZ, US
Contact	MOIRA BARTON
510(k) history	36 submissions · 36 cleared · 2001-2007