

K052918 REPROCESSED EXTERNAL FIXATION DEVICESJan 27, 2006
102 days to decisionK052918 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k052918/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 20, 2026