

**K032058 MODIFICATION TO REPROCESSED SYNTHESIS
EXTERNAL FIXATION DEVICES**Jul 30, 2003
28 days to decisionK032058 · Product code: KTT · Orthopedic
Source: <https://www.510kdatabase.net/k032058/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Appliance, Fixation, Nail/blade/plate Combination, Multiple Component (KTT)
Date received	Jul 2, 2003
Decision date	Jul 30, 2003
Days to decision	28 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/k032058/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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