FDA 510(k) Premarket Notification

K023324 MODIFICATION TO EBI XFIX DFS SYSTEM

CLEARED

Oct 18, 2002
14 days to decision

K023324 · Product code: **KTT** · Orthopedic
Source: https://www.510kdatabase.net/k023324/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Appliance, Fixation, Nail/blade/plate Combination, Multiple Component (KTT)
Date received	Oct 4, 2002
Decision date	Oct 18, 2002
Days to decision	14 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Ebi, L.P.**
Location	Parsippany, NJ, US
Contact	FREDERIC TESTA
510(k) history	95 submissions · 94 cleared · 1997-2010