FDA 510(k) Premarket Notification

K874535 MICROTEK MINI-MAX DRILL SYSTEM

CLEARED

Nov 23, 1987
20 days to decision

K874535 · Product code: **DZI** · Dental
Source: https://www.510kdatabase.net/k874535/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drill, Bone, Powered (DZI)
Date received	Nov 3, 1987
Decision date	Nov 23, 1987
Days to decision	20 days
Third-party review	No

APPLICANT

Company	**Microtek Medical, Inc.**
Location	Mchenry, IL, US
Contact	KEITH MCGEE
510(k) history	24 submissions · 24 cleared · 1984-2011

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k874535/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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