FDA 510(k) Premarket Notification

K880739 DYONICS DRILL GUIDE/ISOMETER SYSTEM

CLEARED

Mar 21, 1988
26 days to decision

K880739 · Product code: **LXI** · Orthopedic
Source: https://www.510kdatabase.net/k880739/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Guide, Drill, Ligament (LXI)
Date received	Feb 24, 1988
Decision date	Mar 21, 1988
Days to decision	26 days
Third-party review	No

APPLICANT

Company	**Dyonics, Inc.**
Location	Walker, MI, US
Contact	KENNETH E CARRIER
510(k) history	19 submissions · 19 cleared · 1978-1990