FDA 510(k) Premarket Notification

K820444 HEART-ALERT

CLEARED

Mar 12, 1982
22 days to decision

K820444 · Product code: **KHX** · Physical Medicine
Source: https://www.510kdatabase.net/k820444/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Platform, Force-measuring (KHX)
Date received	Feb 18, 1982
Decision date	Mar 12, 1982
Days to decision	22 days
Third-party review	No

APPLICANT

Company	**Medical Equipment Devices, Inc.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1982-1982