FDA 510(k) Premarket Notification

K850870 MEDIFETTE ERGOMETER

CLEARED

Jun 12, 1985
103 days to decision

K850870 · Physical Medicine
Source: https://www.510kdatabase.net/k850870/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Date received	Mar 1, 1985
Decision date	Jun 12, 1985
Days to decision	103 days
Third-party review	No

APPLICANT

Company	**Ketronic, Inc.**
Location	Netherlands, US
Contact	VANDILL
510(k) history	10 submissions · 9 cleared · 1984-1985