FDA 510(k) Premarket Notification

K800500 PULSEOMETER

CLEARED

Apr 8, 1980
35 days to decision

K800500 · Product code: **DRT** · Cardiovascular
Source: https://www.510kdatabase.net/k800500/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Mar 4, 1980
Decision date	Apr 8, 1980
Days to decision	35 days
Third-party review	No

APPLICANT

Company	**Kirklands Sporting Goods Co.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1980-1980