FDA 510(k) Premarket Notification

K830282 ELECTRONIC EXERCISE #313-PULSE RATE

CLEARED

Apr 14, 1983
78 days to decision

K830282 · Product code: **DRT** · Cardiovascular
Source: https://www.510kdatabase.net/k830282/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Jan 26, 1983
Decision date	Apr 14, 1983
Days to decision	78 days
Third-party review	No

APPLICANT

Company	**Walton Manufacturer Co.**
Location	Walker, MI, US
510(k) history	1 submissions · 1 cleared · 1983-1983