FDA 510(k) Premarket Notification

K802182 WISCONSIN HEART RATE MONITOR

CLEARED

Oct 31, 1980
52 days to decision

K802182 · Product code: **DRT** · Cardiovascular
Source: https://www.510kdatabase.net/k802182/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Sep 9, 1980
Decision date	Oct 31, 1980
Days to decision	52 days
Third-party review	No

APPLICANT

Company	**Lunar Radiation Corp.**
Location	WI, US
510(k) history	6 submissions · 6 cleared · 1980-1988