FDA 510(k) Premarket Notification

K883109 MINIMON 7136B PATIENT MONITOR

CLEARED

Jan 17, 1989
176 days to decision

K883109 · Product code: **DRT** · Cardiovascular
Source: https://www.510kdatabase.net/k883109/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Jul 25, 1988
Decision date	Jan 17, 1989
Days to decision	176 days
Third-party review	No

APPLICANT

Company	**Kontron, Inc.**
Location	Everett, MA, US
Contact	DAVID CROMWICK
510(k) history	1 submissions · 1 cleared · 1989-1989