FDA 510(k) Premarket Notification

K896386 MINIMON 7133B PATIENT MONITOR

CLEARED

Dec 13, 1989
37 days to decision

K896386 · Product code: **DPS** · Cardiovascular
Source: https://www.510kdatabase.net/k896386/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrocardiograph (DPS)
Date received	Nov 6, 1989
Decision date	Dec 13, 1989
Days to decision	37 days
Third-party review	No

APPLICANT

Company	**Kontron Instruments, Inc.**
Location	Mchenry, IL, US
Contact	DAVID CROMWICK
510(k) history	57 submissions · 57 cleared · 1981-1993