FDA 510(k) Premarket Notification

K895425 MINIMON 7137B PATIENT MONITOR

CLEARED

Sep 26, 1989
19 days to decision

K895425 · Product code: **DQA** · Anesthesiology
Source: https://www.510kdatabase.net/k895425/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Sep 7, 1989
Decision date	Sep 26, 1989
Days to decision	19 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