FDA 510(k) Premarket Notification

K844731 KONTRON SUPERMON 7210 PATIENT MONITOR

CLEARED

May 16, 1985
163 days to decision

K844731 · Product code: **DRT** · Cardiovascular
Source: https://www.510kdatabase.net/k844731/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Dec 4, 1984
Decision date	May 16, 1985
Days to decision	163 days
Third-party review	No

APPLICANT

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