FDA 510(k) Premarket Notification

K812354 KONTRON PMS 7000

CLEARED

Sep 25, 1981
37 days to decision

K812354 · Product code: **DQK** · Cardiovascular
Source: https://www.510kdatabase.net/k812354/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Computer, Diagnostic, Programmable (DQK)
Date received	Aug 19, 1981
Decision date	Sep 25, 1981
Days to decision	37 days
Third-party review	No

APPLICANT

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