FDA 510(k) Premarket Notification

K890791 SIEMENS SIRECUST 730 PATIENT MONITOR

CLEARED

May 2, 1989
74 days to decision

K890791 · Product code: **DPS** · Cardiovascular
Source: https://www.510kdatabase.net/k890791/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrocardiograph (DPS)
Date received	Feb 17, 1989
Decision date	May 2, 1989
Days to decision	74 days
Third-party review	No

APPLICANT

Company	**Siemens Medical Electronics**
Location	Danvers, MA, US
Contact	MURFITT, PHD
510(k) history	15 submissions · 15 cleared · 1988-1995