FDA 510(k) Premarket Notification

K894460 SIEMENS SIRECUST 730, PATIENT MONITOR W/ARRHYTHMIA

CLEARED

Oct 2, 1989
76 days to decision

K894460 · Product code: **DSI** · Cardiovascular
Source: https://www.510kdatabase.net/k894460/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Jul 18, 1989
Decision date	Oct 2, 1989
Days to decision	76 days
Third-party review	No

APPLICANT

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