FDA 510(k) Premarket Notification

K903940 TYPE 9210 S/ECG

CLEARED

Dec 4, 1991
464 days to decision

K903940 · Product code: **DSI** · Cardiovascular
Source: https://www.510kdatabase.net/k903940/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Aug 27, 1990
Decision date	Dec 4, 1991
Days to decision	464 days
Third-party review	No

APPLICANT

Company	**S & W Medico Teknik**
Location	Walker, MI, US
Contact	METTE JORGENSEN
510(k) history	46 submissions · 46 cleared · 1984-1995