FDA 510(k) Premarket Notification

K802312 CC101 ECG MONITORING SYSTEM

CLEARED

Jan 7, 1981
106 days to decision

K802312 · Product code: **DSI** · Cardiovascular
Source: https://www.510kdatabase.net/k802312/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Sep 23, 1980
Decision date	Jan 7, 1981
Days to decision	106 days
Third-party review	No

APPLICANT

Company	**Anthro-Metrics Corp.**
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1981-1981