FDA 510(k) Premarket Notification

K821711 CARDIO TEHCHANALYZER

CLEARED

Oct 4, 1982
117 days to decision

K821711 · Product code: **DSI** · Cardiovascular
Source: https://www.510kdatabase.net/k821711/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Jun 9, 1982
Decision date	Oct 4, 1982
Days to decision	117 days
Third-party review	No

APPLICANT

Company	**Cardio Technology , Ltd.**
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1982-1983