

K821711 CARDIO TEHCHANALYZEROct 4, 1982
117 days to decisionK821711 · 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

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Device record: <https://www.510kdatabase.net/k821711/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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