

K802103 ECG RECALL & EDIT PRODUCT VERSION-A3601Sep 26, 1980
23 days to decisionK802103 · Product code: **DSI** · Cardiovascular
Source: <https://www.510kdatabase.net/k802103/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Sep 3, 1980
Decision date	Sep 26, 1980
Days to decision	23 days
Third-party review	No

APPLICANT

Company	General Electric Co.
Location	Mchenry, IL, US
510(k) history	254 submissions · 254 cleared · 1976-2011

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Device record: <https://www.510kdatabase.net/k802103/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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