

K812914 MICRO-PREPROCESSOR UNITDec 29, 1981
71 days to decisionK812914 · Product code: **DSI** · Cardiovascular
Source: <https://www.510kdatabase.net/k812914/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Oct 19, 1981
Decision date	Dec 29, 1981
Days to decision	71 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/k812914/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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