

K810331 DUO-TELMar 20, 1981
39 days to decisionK810331 · Product code: **DRG** · Cardiovascular
Source: <https://www.510kdatabase.net/k810331/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transmitters And Receivers, Physiological Signal, Radiofrequency (DRG)
Date received	Feb 9, 1981
Decision date	Mar 20, 1981
Days to decision	39 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/k810331/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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