

K803286 CARDIOMEMOMar 17, 1981
77 days to decisionK803286 · Product code: **KRE** · Cardiovascular
Source: <https://www.510kdatabase.net/k803286/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Analyzer, Pacemaker Generator Function, Indirect (KRE)
Date received	Dec 30, 1980
Decision date	Mar 17, 1981
Days to decision	77 days
Third-party review	No

APPLICANT

Company	Instromedix, Inc.
Location	Mchenry, IL, US
510(k) history	32 submissions · 32 cleared · 1977-1997

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Device record: <https://www.510kdatabase.net/k803286/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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