

K800696 ELECTRONIC PACEMAKER WAVEFORM ANALYZERAug 14, 1980
139 days to decisionK800696 · Product code: **KRE** · CardiovascularSource: <https://www.510kdatabase.net/k800696/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Analyzer, Pacemaker Generator Function, Indirect (KRE)
Date received	Mar 28, 1980
Decision date	Aug 14, 1980
Days to decision	139 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/k800696/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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