

K780175 RATE PROGRAMMER MODEL 520Apr 21, 1978
77 days to decisionK780175 · Product code: **KRG** · CardiovascularSource: <https://www.510kdatabase.net/k780175/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Programmer, Pacemaker (KRG)
Date received	Feb 3, 1978
Decision date	Apr 21, 1978
Days to decision	77 days
Third-party review	No

APPLICANT

Company	Intermedics, Inc.
Location	Mchenry, IL, US
510(k) history	211 submissions · 201 cleared · 1977-1996

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Device record: <https://www.510kdatabase.net/k780175/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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