

K770865 MONITOR, PACEMAKERJun 14, 1977
34 days to decisionK770865 · Product code: **DTC** · CardiovascularSource: <https://www.510kdatabase.net/k770865/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Analyzer, Pacemaker Generator Function (DTC)
Date received	May 11, 1977
Decision date	Jun 14, 1977
Days to decision	34 days
Third-party review	No

APPLICANT

Company	Adcor Electronics, Inc.
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1977-1977

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Device record: <https://www.510kdatabase.net/k770865/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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