

K780569 L-12, L-21, AND L-22 ELECTRODESJun 14, 1978
64 days to decisionK780569 · Product code: **DTB** · CardiovascularSource: <https://www.510kdatabase.net/k780569/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Permanent Pacemaker Electrode (DTB)
Date received	Apr 11, 1978
Decision date	Jun 14, 1978
Days to decision	64 days
Third-party review	No

APPLICANT

Company	Coratomic, Inc.
Location	Mchenry, IL, US
510(k) history	15 submissions · 14 cleared · 1976-1986

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Device record: <https://www.510kdatabase.net/k780569/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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