

K800353 ADAPTER KITMay 28, 1980
103 days to decisionK800353 · Product code: **DTD** · Cardiovascular
Source: <https://www.510kdatabase.net/k800353/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pacemaker Lead Adaptor (DTD)
Date received	Feb 15, 1980
Decision date	May 28, 1980
Days to decision	103 days
Third-party review	No

APPLICANT

Company	Daig Corp.
Location	Mchenry, IL, US
510(k) history	63 submissions · 63 cleared · 1977-2000

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Device record: <https://www.510kdatabase.net/k800353/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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