FDA 510(k) Premarket Notification

K800353 ADAPTER KIT

CLEARED

May 28, 1980
103 days to decision

K800353 · 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