

K894431 MODIFIED LABELING TO PERMANENT LEAD INTRODUCERSep 6, 1989
51 days to decisionK894431 · Product code: **DYB** · Cardiovascular
Source: <https://www.510kdatabase.net/k894431/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Introducer, Catheter (DYB)
Date received	Jul 17, 1989
Decision date	Sep 6, 1989
Days to decision	51 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k894431/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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