FDA 510(k) Premarket Notification

K894431 MODIFIED LABELING TO PERMANENT LEAD INTRODUCER

CLEARED

Sep 6, 1989
51 days to decision

K894431 · 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