FDA 510(k) Premarket Notification

K894500 DAIG ELECTROPHYSIOLOGY CATHETER

CLEARED

Sep 18, 1989
61 days to decision

K894500 · Product code: **DRF** · Cardiovascular
Source: https://www.510kdatabase.net/k894500/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF)
Date received	Jul 19, 1989
Decision date	Sep 18, 1989
Days to decision	61 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