FDA 510(k) Premarket Notification

K942379 DAIG DIAGNOSTIC ELECTROPHYSIOLOGY CATHETER

CLEARED

Dec 27, 1994
232 days to decision

K942379 · Product code: **DRF** · Cardiovascular
Source: https://www.510kdatabase.net/k942379/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF)
Date received	May 9, 1994
Decision date	Dec 27, 1994
Days to decision	232 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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