FDA 510(k) Premarket Notification

K913940 RESPONSE(TM) STEERABLE CATHETER

CLEARED

Jan 26, 1993
511 days to decision

K913940 · Product code: **DRA** · Cardiovascular
Source: https://www.510kdatabase.net/k913940/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Steerable (DRA)
Date received	Sep 3, 1991
Decision date	Jan 26, 1993
Days to decision	511 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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