

K913940 RESPONSE(TM) STEERABLE CATHETERJan 26, 1993
511 days to decisionK913940 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k913940/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 3, 2026