

K894330 DAIG VESSEL DILATORSep 12, 1989
60 days to decisionK894330 · Product code: **DRE** · CardiovascularSource: <https://www.510kdatabase.net/k894330/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dilator, Vessel, For Percutaneous Catheterization (DRE)
Date received	Jul 14, 1989
Decision date	Sep 12, 1989
Days to decision	60 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k894330/>, 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