

K931659 3 LEAD CONNECTORJan 7, 1994
277 days to decisionK931659 · Product code: **FPA** · General Hospital
Source: <https://www.510kdatabase.net/k931659/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Apr 5, 1993
Decision date	Jan 7, 1994
Days to decision	277 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Douglas Medical Products Corp.
Location	Mundelein, IL, US
Contact	DOUGLAS MEDICAL
510(k) history	19 submissions · 19 cleared · 1992-1996

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k931659/> 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 May 22, 2026