

K941453 CHECK VALVE SOLUTION SETAug 16, 1994
144 days to decisionK941453 · Product code: **FPA** · General HospitalSource: <https://www.510kdatabase.net/k941453/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Mar 25, 1994
Decision date	Aug 16, 1994
Days to decision	144 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k941453/> 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