

K942503 UNICATH FAMILYJul 20, 1994
55 days to decisionK942503 · Product code: IZI · Radiology
Source: <https://www.510kdatabase.net/k942503/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Angiographic (IZI)
Date received	May 26, 1994
Decision date	Jul 20, 1994
Days to decision	55 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Angiographic Devices Corp.
Location	Littleton, MA, US
Contact	WAYNE V LOOMER
510(k) history	1 submissions · 1 cleared · 1994-1994

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k942503/> 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 20, 2026