

K900773 NUCLEAR MAXMay 23, 1990
96 days to decisionK900773 · Product code: **IYX** · Radiology
Source: <https://www.510kdatabase.net/k900773/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Camera, Scintillation (gamma) (IYX)
Date received	Feb 16, 1990
Decision date	May 23, 1990
Days to decision	96 days
Third-party review	No

APPLICANT

Company	Scientific Imaging, Inc.
Location	Fairfield, CT, US
Contact	LEAR, M.D.
510(k) history	3 submissions · 3 cleared · 1988-2005

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k900773/>, 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 14, 2026