FDA 510(k) Premarket Notification

K900773 NUCLEAR MAX

CLEARED

May 23, 1990
96 days to decision

K900773 · 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