FDA 510(k) Premarket Notification

K924719 ANCOR

CLEARED

Dec 30, 1992
100 days to decision

K924719 · Product code: **IZI** · Radiology
Source: https://www.510kdatabase.net/k924719/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Angiographic (IZI)
Date received	Sep 21, 1992
Decision date	Dec 30, 1992
Days to decision	100 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Siemens Medical Solutions USA, Inc.**
Location	Hoffman Estates, IL, US
Contact	ANNE PINTO
510(k) history	778 submissions · 778 cleared · 1980-2026